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# Cation deficiency enables reversal of dopant segregation at perovskite oxide surfaces under anodic potential†

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Surface instability of perovskite oxides caused by aliovalent dopant segregation and resultant formation of insulating surface layers and precipitates, such as SrO<sub>x</sub>, limits the performance and durability of these materials in energy conversion, including in solid oxide fuel and electrolysis cells. Previous studies showed that reversal (re-incorporation) of such dopant segregation layers can occur under both cathodic and anodic potential in Sr-doped lanthanum manganite (LSM) as a model perovskite system. Sr segregation under anodic and reversal of segregation under cathodic potential could be explained by a defect equilibrium reaction involving A-site cation vacancies and Mn oxidation state. Sr segregation under cathodic and reversal of segregation under anodic potential is also known to take place, but it has been poorly understood and its governing mechanism is not known. In this study, we show that defect reactions involving cation vacancies and oxygen holes at high oxygen pressures can account for re-incorporation of the segregated dopant from the surface into the bulk of lanthanum manganite under anodic conditions. We use Ca-doped lanthanum manganite (LCM) as a model perovskite oxide. We have demonstrated with X-ray photoelectron and absorption spectroscopy that the precipitation and re-incorporation of the surface CaO<sub>x</sub> under cathodic and anodic potential, respectively, are coupled with the reduction and oxidation of the perovskite surface. Under oxidizing, anodic conditions, cation-deficient perovskite unit cells form on/near the surface, allowing for the dissolution of the excess Ca at the surface back into the perovskite lattice. Notably, applying anodic potential could remove half of the surface Ca precipitates within a few minutes, thus improving the oxygen exchange kinetics of the LCM surface significantly. These results establish a highly efficient oxidative route for activating perovskite oxide surfaces and advance the quantitative understanding of dopant segregation and re-incorporation reactions.

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## Introduction

Perovskite oxides (ABO<sub>3</sub>) are widely used in clean energy conversion applications, including solid oxide fuel and electrolysis cells (SOFC/ECs),<sup>1–6</sup> solar thermochemical fuel production,<sup>7–9</sup> and gas separation membranes.<sup>10</sup> Many state-of-the-art perovskites, such as La<sub>1–x</sub>Sr<sub>x</sub>MnO<sub>3</sub>,<sup>11–16</sup> La<sub>1–x</sub>Sr<sub>x</sub>CoO<sub>3</sub>,<sup>17–22</sup> La<sub>1–x</sub>Sr<sub>x</sub>Co<sub>1–y</sub>Fe<sub>y</sub>O<sub>3</sub>,<sup>23–27</sup> and SrTi<sub>x</sub>Fe<sub>1–x</sub>O<sub>3</sub> (ref. 28–31) suffer from degradation of their surface chemistry and oxygen exchange kinetics at elevated temperatures. This process limits

their long-term stability in the aforementioned applications. This degradation is primarily because of dopant segregation to the surface and precipitation in the form of insulating phases, such as SrO<sub>x</sub>. These insulating surface layers block the electron transfer and oxygen exchange pathways, impeding the oxygen exchange kinetics<sup>12,24,32</sup> and reducing the attainable power densities.<sup>2</sup>

In our previous studies, we have demonstrated that electrostatic ( $E_{\text{elec}}$ ) and elastic ( $E_{\text{ela}}$ ) energies of dopant are important driving forces for dopant segregation on perovskite oxides.<sup>33,34</sup> Predominantly under cathodic overpotentials (or low  $pO_{2,\text{eff}}$ ), negatively charged dopant (Sr'<sub>La</sub>) can segregate to the surface due to the electrostatic attraction ( $E_{\text{elec}}$ ) to the positively charged oxygen vacancies ( $V_{\text{O}}^{\bullet}$ ) enriched at the surface. Under anodic overpotentials (or high  $pO_{2,\text{eff}}$ ), too, the dopant can segregate and precipitate in the form of a binary oxide at the surface, e.g. SrO<sub>x</sub>. This is because precipitation of the larger dopant cation ( $R_{\text{La}^{3+}} < R_{\text{Sr}^{2+}}$ ) to the surface relaxes the elastic energy ( $E_{\text{ela}}$ ) in the perovskite phase which contracts when oxidized. Due to their different overpotential dependence,  $E_{\text{elec}}$  and  $E_{\text{ela}}$  dominate at

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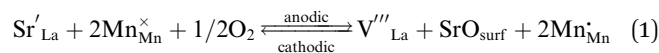
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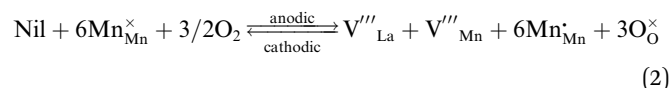
the low and high  $pO_{2,eff}$  regimes, respectively, giving rise to a convex total segregation behavior with respect to  $pO_{2,eff}$  (red and blue solid lines in Fig. 1a).

Importantly, the fact that dopant segregation is a thermodynamically driven process implies that it can be reversed if the reverse reaction is kinetically unrestricted. Several previous studies reported that dopant segregation induced by anodic potential (or under oxidizing environment) could be reversed with cathodic potential.<sup>13,35,36</sup> Jiang<sup>13</sup> and Janek<sup>36</sup> groups have reported that applying cathodic potential to  $(La_{0.8}Sr_{0.2})_{0.9}MnO_3$  and  $La_{1-x}Sr_xCoO_3$  incorporated the surface  $SrO_x$  back into the perovskite phase. This led to the improvement of the surface oxygen exchange rate by several orders of magnitude.<sup>13</sup> Based on the role of  $E_{ela}$  described in Fig. 1a, we can reason that the reduction of  $E_{ela}$  under reducing conditions (or cathodic potential) is one reason for this reversal of segregation. However, increasing contribution from  $E_{elec}$  under reducing conditions can actually drive segregation as well.<sup>19,34,37</sup> Therefore, it is difficult to reconcile this reversal of segregation under cathodic conditions only by considering  $E_{ela}$  and  $E_{elec}$  as shown in Fig. 1a. In addition to the elastic and electrostatic energy contributions, we believe that we must also consider the defect equilibria in the perovskite oxide in the segregation in the form of dopant oxide precipitates, and reversal of such segregation on perovskite oxides, as we have recently studied theoretically.<sup>38</sup> Eqn (1) demonstrates the known defect reaction that can explain the dopant segregation (forward) under anodic conditions and dopant re-incorporation (backward) under cathodic conditions.<sup>36,39</sup>



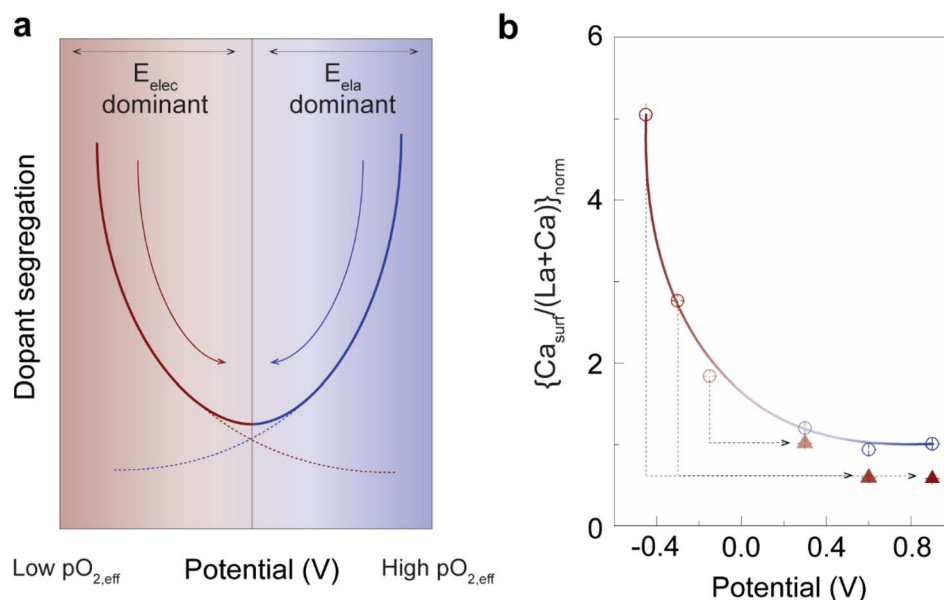
( $Sr'_{La}$  = Sr on the La lattice site,  $Mn^{\times}_{Mn}$  = Mn on the Mn lattice site,  $V'''_{La}$  = La vacancy,  $Mn^{\times}_{Mn}$  = Mn with single positive charge on the Mn lattice site)

In contrast, the opposite segregation behavior, dopant segregation under cathodic and its reversal under anodic conditions, is also observed in similar perovskite oxides such as  $La_{1-x}Sr_xCo_{1-y}Fe_yO_3$  (ref. 20, 23 and 24) and  $La_{1-x}Sr_xMnO_3$ .<sup>40-42</sup> However, the defect reaction in eqn (1) cannot explain this segregation and reversal phenomenon. Also, as described in Fig. 1a, going toward anodic conditions suppresses the  $E_{elec}$  contribution but it also increases the elastic energy contribution to Sr segregation. Hence, the  $E_{ela}$  and  $E_{elec}$  argument fails to explain the dopant re-incorporation under anodic conditions after segregation at cathodic conditions as well. Therefore, in addition to eqn (1), we need to consider another defect equilibrium to fully explain this contradicting segregation and reversal phenomenon. To this end, a defect reaction that we may have to consider is:



(nil =  $V'''_{La} + V'''_{Mn} + 3V''_O$  (empty unit cells),  $V''_O$  = O vacancy,  $V'''_{Mn}$  = Mn vacancy,  $O^{\times}_O$  = O on the O lattice site)

This Schottky reaction under oxidizing conditions is known from the bulk point defect equilibrium in doped lanthanum manganites where cation vacancies are formed as the perovskite



**Fig. 1** Electrochemically driven dopant segregation and re-incorporation. (a) Schematic rendering of the dependence of dopant segregation on electrochemical potential for an acceptor-doped lanthanum manganite, described by the sum of the two driving forces; red solid and dashed line = electrostatic energy ( $E_{elec}$ ), blue solid and dashed line = elastic energy ( $E_{ela}$ ). (b) Segregation vs. potential graph of  $La_{0.8}Ca_{0.2}MnO_3$  (LCM). The thin-film LCM samples were annealed under different potentials at 770 °C for 1 h in pure oxygen by using the lateral polarization method (open circles).<sup>34</sup> Another set of LCM samples were annealed with the two-step polarization approach to investigate the reversal of Ca segregation. In this approach, the samples were initially annealed for 1 h under cathodic potential, followed by anodic potential for another 1 h; black dashed arrows indicate the change of the applied local potential during the two-step polarization experiment. The resulting  $\{Ca_{surf}/(La+Ca)\}_{norm}$  after this two-step polarization experiment are plotted as triangles. All the error bars are smaller than the size of the data points.



oxides incorporate oxygens into the lattice as they are oxidized.<sup>43,44</sup> Importantly, this reaction also describes the formation of new empty perovskite unit cells (labeled as 'nil') with cation vacancies. Hinted by this, we reasoned that these cation-deficient unit cells at the surface could take in the segregated Sr, leading to the dopant re-incorporation under anodic conditions. The defect reaction in eqn (2) can be coupled with the dopant re-incorporation reaction occurring at the surface under anodic conditions.

In this study, we propose a new defect reaction that successfully explains the dopant segregation under cathodic and re-incorporation under anodic potential. We used thin-film  $\text{La}_{0.8}\text{Ca}_{0.2}\text{MnO}_3$  (LCM) as a primary model system because dopant segregation in LCM increases monotonically as potential goes from anodic to cathodic,<sup>34</sup> which corresponds exactly to the dopant segregation and re-incorporation reactions that we aim to investigate. With X-ray photoelectron spectroscopy (XPS) and X-ray absorption spectroscopy (XAS) measurements, we quantitatively analyzed the dopant segregation and re-incorporation reactions on LCM induced by electrochemical potential and found that dopant re-incorporation reaction on LCM is coupled to its surface oxidation process. We have found that such an oxidation process at the 'surface' is mediated predominantly by oxygen, O, rather than Mn which becomes redox-active only in the bulk. Also, the fact that lanthanum manganites equilibrate with cation vacancies under oxidizing conditions (eqn (2)) is key to formulating the defect reaction explaining the dopant re-incorporation reaction. Electrochemical impedance spectroscopy (EIS) showed that applying anodic potential to LCM improved its oxygen exchange kinetics, *i.e.*, reduced impedance, in a short time (<5 min) by removing the insulating  $\text{CaO}_x$  at the surface. This process was reversible over several cycles. Also, we show that the results from LCM can be transferred to  $\text{La}_{0.8}\text{Sr}_{0.2}\text{MnO}_3$  (LSM) which is one of the state-of-the-art air electrodes in solid oxide fuel and electrolysis cells. These findings advance our understanding of the governing mechanisms of surface instability and provide insights for operating perovskite oxides to maintain their surface chemical stability in a range of energy conversion applications.

## Experimental methods

### Sample preparation and characterizations

$\text{La}_{0.8}\text{Ca}_{0.2}\text{MnO}_3$ ,  $\text{La}_{0.8}\text{Sr}_{0.2}\text{MnO}_3$ , and  $\text{La}_{0.8}\text{Ba}_{0.2}\text{MnO}_3$  thin films were deposited onto (100)-oriented single crystals of YSZ (yttria-stabilized zirconia, 8 mol%  $\text{Y}_2\text{O}_3$ , MTI Corp.) *via* pulsed laser deposition (PLD). During the deposition, the substrate temperature was maintained at 700 °C and the oxygen pressure of the chamber was kept at 0.5 mTorr. The oxygen pressure was increased to 500 mTorr  $\text{O}_2$  while cooling to room temperature. In order to remove pre-existing surface dopant oxide layers, which form at the surface during PLD deposition, the thin-film samples were immersed in high-purity deionized water for 1 min. The effect of this pre-treatment process was demonstrated in earlier work.<sup>45</sup> The thickness of the thin films was measured by Bruker DXT Stylus profilometer and was measured 80–100 nm. To examine the dopant segregation and re-

incorporation on  $\text{La}_{0.8}\text{Ca}_{0.2}\text{MnO}_3$ ,  $\text{La}_{0.8}\text{Ca}_{0.2}\text{MnO}_3$  thin-film samples from the same PLD batch were used for each set of potential experiments to avoid any experimental artefact. Out-of-plane X-ray diffraction (XRD, Rigaku Smartlab) measurement showed that all the films were in the perovskite phase and had preferential texture along (110) and (100). The surfaces of the films were imaged by high-resolution scanning electron microscopy (HRSEM) using a Zeiss Merlin HRSEM with an in-lens secondary electron detector.

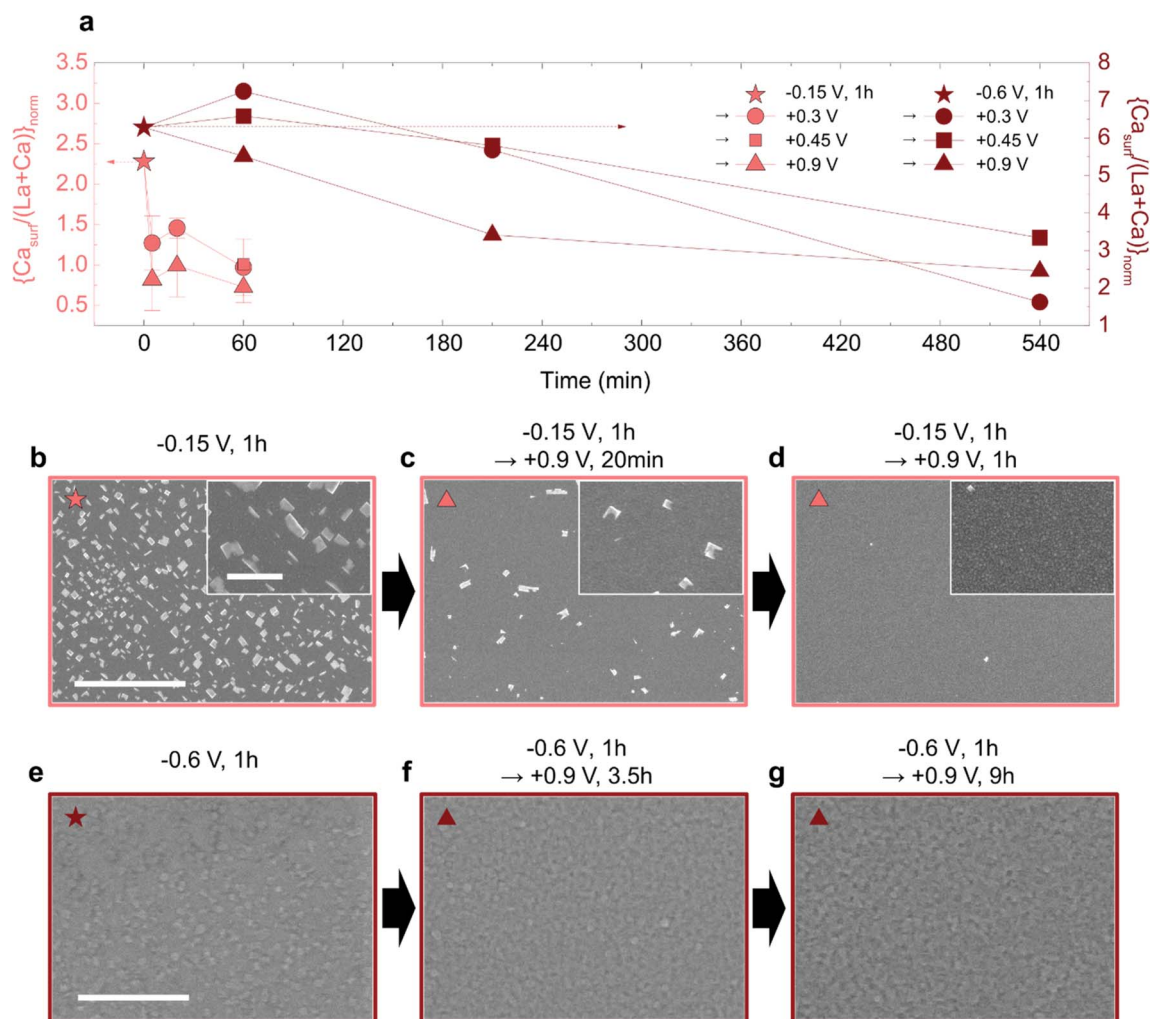
### X-ray photoelectron spectroscopy (XPS) measurement

The chemical composition of the films' near-surface region was characterized by X-ray photoelectron spectroscopy (XPS) using a Physical Electronics Versaprobe II X-ray photoelectron spectrometer equipped with a monochromated Al  $K\alpha$  X-ray source. The angle between the analyzer and the surface plane was set to 45°. Please note that all the XPS experiments were done *ex situ*. For quantitative analysis, core-level emission lines with similar inelastic mean free paths ( $\lambda$ ) were chosen; La 4d ( $\lambda = 1.66$ – $1.69$  nm), Mn 3p ( $\lambda = 1.72$ – $1.74$  nm), Ba 4d ( $\lambda = 1.70$  nm), Sr 3d ( $\lambda = 1.65$  nm), and Ca 2p ( $\lambda = 1.43$  nm).<sup>46</sup> The resulting spectra were analyzed to quantify the amount of each component by normalizing their peak area by the sensitivity factor of each core-level orbital.<sup>47</sup> The quantification of the  $\text{Ca}_{\text{surf}}$  on LCM samples was done in this way. In Fig. 2, 4, and 6, the amount of the  $\text{Ca}_{\text{surf}}$  was normalized with the total A-site cations ( $\text{Ca}_{\text{surf}}/(\text{La} + \text{Ca})$ ), and further normalized with the  $\text{Ca}_{\text{surf}}/(\text{La} + \text{Ca})$  of a separate reference sample that had the least segregated state. This reference sample was annealed solely under +0.3 V for 1 h. In Fig. 1 and 7, the normalization was done in the same way but the reference sample was the as-prepared LCM and LSM. We used the reference sample because LCM films from different PLD batches often showed a big difference in the absolute amount of  $\text{Ca}_{\text{surf}}$  even under the same annealing conditions. Due to the limit of the number of thin films we could deposit in a PLD batch at once, we used the amount of  $\text{Ca}_{\text{surf}}$  on one of the LCM thin films annealed under +0.3 V for 1 h as a reference segregation state of each PLD batch.

### Lateral polarization method

Lateral polarization method enabled us to investigate a wide range of potentials with a small number of thin-film samples, thus minimizing experimental artifacts. Possible experimental artifacts include using samples from different PLD batches or different experimental conditions such as variations in sputtering conditions or annealing temperatures. This method is based on the original design by Huber *et al.*<sup>48</sup> and the utility of this method for segregation study was previously demonstrated by our group<sup>34</sup> (Fig. S1†). For the preparation of the cells, Pt paste (SPI Corp.) was applied at the back side and one edge of YSZ crystals before the thin-film growth process to form a porous Pt counter electrode (CE). The deposited Pt electrode was heated at 700 °C in the air for 30 min to evaporate the solvent and sinter the electrode. After depositing the thin films, electrical contact points were created by sputter deposition of a Ti adhesion layer (10 nm) followed by a Pt contact layer (100





**Fig. 2** Re-incorporation of surface segregates under anodic potential and the resulting changes in the surface morphology of the LCM film. (a) Amount of  $Ca_{surf}$  vs. time duration of anodic potential. Two different cathodic potentials,  $-0.15$  V (pink star, y-axis on the left) and  $-0.6$  V (brown star, y-axis on the right), were first applied to the LCM thin films to induce dopant segregation. Different anodic potentials,  $+0.3$  V (circles),  $+0.45$  V (squares), and  $+0.9$  V (triangles), were then applied for different time durations to induce dopant re-incorporation. The error bars in (a) are standard deviation of the results of the three different sample sets deposited from different PLD batches. SEM images of the sample annealed under (b)  $-0.15$  V, 1 h, (c)  $-0.15$  V, 1 h  $\rightarrow$   $+0.9$  V, 20 min, (d)  $-0.15$  V, 1 h  $\rightarrow$   $+0.9$  V, 1 h, (e)  $-0.6$  V, 1 h, (f)  $-0.6$  V, 1 h  $\rightarrow$   $+0.9$  V, 3.5 h, and (g)  $-0.6$  V, 1 h  $\rightarrow$   $+0.9$  V, 9 h. The colored symbols in the top-left corner of the images indicate the corresponding data points in (a). Scale bars in (b) and (e) indicate  $4 \mu\text{m}$  and the one in the inset of (b) indicates  $800 \text{ nm}$ .

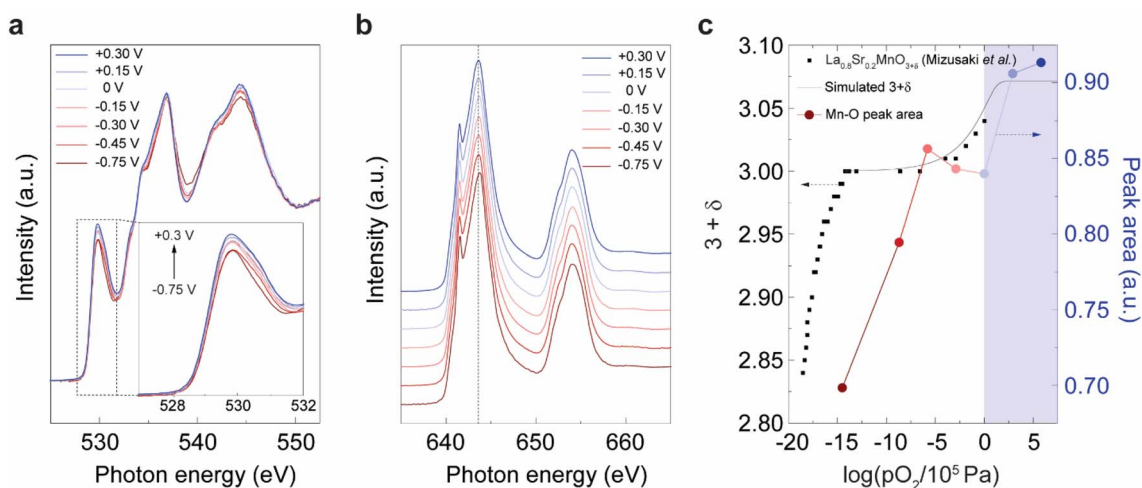
nm). The regions for sputtering were defined by a Kapton mask which covered the film surface, leaving a  $2 \text{ mm}$  gap at the side for depositing the upper side of the CE. The WE was deposited through a  $3 \times 7 \text{ mm}^2$  rectangular opening of the mask at a distance of  $3 \text{ mm}$  from the CE. With such a configuration, we could create the electrical connection between the WE and CE along the perovskite thin-film electrode. Upon application of electrical potential to the WE, the thin film serves as an ohmic resistor for the electronic current, resulting in a linear voltage drop across the distance between the WE and CE. The local potential applied at a local measurement point can be calculated with the applied potential on the WE and the distance between the measurement point and the WE (Fig. S1b†). This was proven by an *in situ* XPS measurement performed in our previous study.<sup>34</sup> After potential experiments, the exposed

surface of the thin film in between the WE and CE was analyzed with a variety of characterization tools, such as XPS and SEM, by scanning along the sample surface. Further details can be found in our previous report.<sup>34</sup> In the two-step polarization approach, cathodic and anodic potentials were sequentially applied to LCM for 1 h each, and the potential change was done gradually with a rate of  $1 \text{ mV s}^{-1}$  to prevent any electrochemical shock to the films.

#### X-ray absorption spectroscopy (XAS) measurement

XAS reveals the oxidation state of a sample in an element-specific way, and thus allows us to find the relationship between the oxidation of the samples and dopant re-incorporation reaction, and also which chemical species are involved in the oxidation process. In general, the redox-active





**Fig. 3** Redox-active species at the surface of LCM. (a) O K-edge and (b) Mn L-edge spectra in total-electron-yield (TEY) mode measured from the LCM thin-film samples annealed under different potentials. The inset of (a) shows a magnified view of the Mn 3d–O 2p hybridization peak at  $\sim 530$  eV. (c) Comparison between the integrated area of the Mn 3d–O 2p peaks under different potentials (colored circles, y-axis on the right) and the oxygen non-stoichiometry curve of LSM obtained under similar experimental conditions<sup>43</sup> (black dots, y-axis on the left). The blue-colored region in (c) indicates the range of the effective  $pO_2$  ( $\log(pO_2) > 0$ ) induced by applying anodic potential.

species of perovskite oxides are transition metal cations or oxygen anions as they constitute the valence band.<sup>49–51</sup> Hence, we investigated the O K-edge and Mn L-edge spectra of the  $La_{0.8}Ca_{0.2}MnO_3$  (LCM) samples used in our experiments. Please also note that all the XAS experiments were done *ex situ*. The oxidation of the LCM thin-film samples was probed by using XAS at beamline 23-ID-2 of the National Synchrotron Light Source II in Brookhaven National Laboratory (BNL). O K-edge and Mn L-edge X-ray absorption spectra were collected from each sample. The spectra were collected both in the total-electron-yield (TEY) and partial-fluorescence-yield (PFY) modes, corresponding to a probing depth of less than 10 nm and 100 nm, respectively. In the O K-edge spectra of  $La_{0.8}Ca_{0.2}MnO_3$  thin-film samples, the integrated area of the peak at  $\sim 530$  eV ( $I_{val}$ ) serves as an indicator of the concentration of the oxygen holes in the valence band, thus the oxidation of the sample. Thus, we used  $I_{val}$  to quantify the oxidation extent of LCM. Possible contributions from the formation of binary oxides or hydroxides, such as CaO, Ca(OH)<sub>2</sub>, CaCO<sub>3</sub>, La<sub>2</sub>O<sub>3</sub>, and MnO<sub>x</sub>, to the  $I_{val}$  are negligible.<sup>52–56</sup> The concentration of the lattice Ca ( $Ca_{latt}$ ) which could also influence the  $I_{val}$  does not show any trend during the oxidation of LCM (Fig. S22†), thus confirming that the increase in  $I_{val}$  is attributed to the oxidation of LCM.

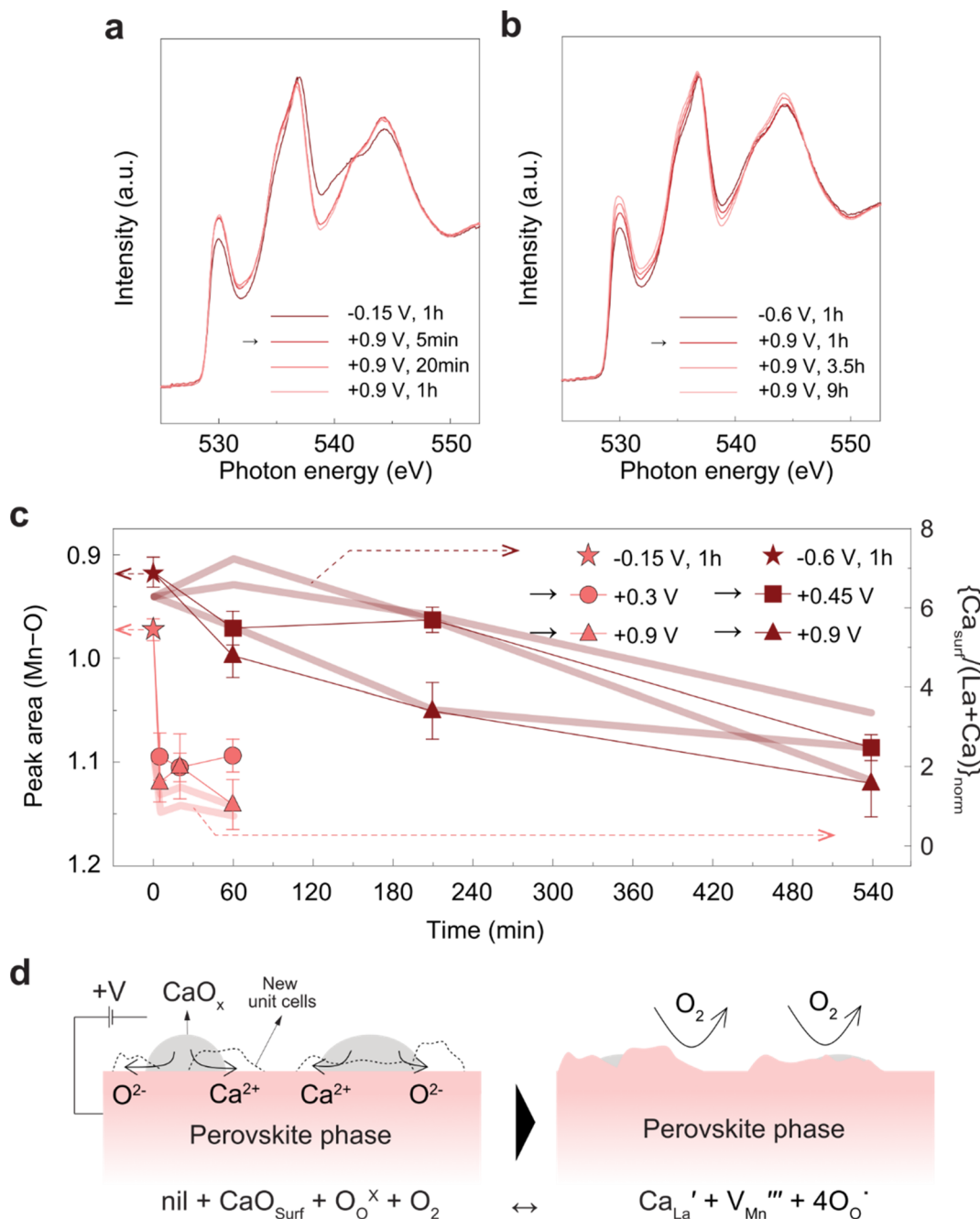
## Results and discussion

### Dopant segregation and re-incorporation on Ca-doped LaMnO<sub>3</sub> under electrochemical potential

First, we investigated the segregation behavior of LCM under different potentials using the *lateral polarization* approach (Fig. S1†). For this, two LCM thin-film samples were kept at 770 °C in 1 atm O<sub>2</sub> for 1 h under a wide range of potentials,  $-0.6$  and  $+1.2$  V at the WE, respectively. For the following *ex situ*

analysis, we chose three local points between the CE and WE on the two LCM samples. The local potentials applied during the lateral polarization experiment on the sample annealed under  $-0.6$  V (at its WE) were  $-0.45$ ,  $-0.3$ , and  $-0.15$  V, and those on the sample annealed under  $+1.2$  V (at its WE) were  $+0.3$ ,  $+0.6$ , and  $+0.9$  V. The extent of the resulting segregation was analyzed with X-ray photoelectron spectroscopy (XPS). Ca 2p core-level spectra were deconvoluted into the surface ( $Ca_{surf}$ , red envelopes in Fig. S2a†) and lattice components ( $Ca_{latt}$ , blue envelopes in Fig. S2a†). This allowed the quantification of the surface CaO<sub>x</sub>-like precipitates ( $Ca_{surf}$ ) which were formed as a result of segregation. The amount of  $Ca_{surf}$  was first divided by the total A-site cations and the resulting  $Ca_{surf}/(La + Ca)$  was then normalized by the  $Ca_{surf}/(La + Ca)$  of the as-prepared sample. The resulting  $\{Ca_{surf}/(La + Ca)\}_{norm}$  are plotted with respect to the applied local potential as open circles in the segregation graph of LCM in Fig. 1b. The monotonic relationship between the amount of  $Ca_{surf}$  and potential is consistent with our previous results.<sup>34</sup> In our previous work, we confirmed that such a monotonic increase in the amount of  $Ca_{surf}$  observed by XPS was in line with the increase in the surface roughness of the LCM films calculated by AFM.<sup>34</sup> Next, in order to examine the reversibility of Ca segregation, another set of LCM samples were annealed with a *two-step polarization* approach. In this approach, cathodic and anodic potentials were sequentially applied to the LCM samples for 1 h each. The same local cathodic potentials,  $-0.45$ ,  $-0.3$ , and  $-0.15$  V, as in the previous sample were applied for the first 1 h, followed by local anodic potential of  $+0.3$ ,  $+0.6$ , and  $+0.9$  V, respectively, for another hour. The resulting amounts of the  $Ca_{surf}$  after this two-step polarization experiment were again quantified by XPS, normalized by the same method as before ( $\{Ca_{surf}/(La + Ca)\}_{norm}$ ), and are plotted as triangles in Fig. 1b. It can be seen that the amounts of the  $Ca_{surf}$  on these samples were close to



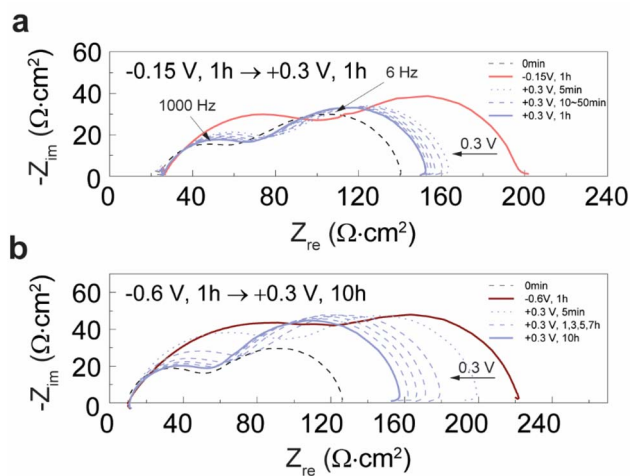


**Fig. 4** Dopant re-incorporation under anodic potential correlates with the oxidation of the LCM surface. (a and b) Total-electron-yield O K-edge XAS spectra measured from the LCM thin films annealed under different potentials. (c) Integrated area of the Mn 3d–O 2p peak in O K-edge XAS with respect to the time duration of anodic potential (pink and brown data points, y-axis on the left); samples polarized under  $-0.15$  V, 1 h (pink star),  $-0.15$  V  $\rightarrow$   $+0.3$  V (pink circles),  $-0.15$  V  $\rightarrow$   $+0.9$  V (pink triangles),  $-0.6$  V, 1 h (brown star),  $-0.6$  V  $\rightarrow$   $+0.45$  V (brown squares), and  $-0.6$  V  $\rightarrow$   $+0.9$  V (brown triangles). The temporal changes of the amount of  $Ca_{surf}$  (Fig. 2a) are also plotted as thick solid lines (y-axis on the right). (d) Proposed dopant re-incorporation reaction on the LCM surface under anodic potential.

those on the sample annealed only under anodic potential that we got before (circles at the same potentials). This indicates that Ca segregation resulting from cathodic potential can be reversed by applying anodic potential afterwards. This result was further supported by additional experiments and SEM measurement which will be explained in the next section. Recalling the underlying energetics of dopant segregation

described in Fig. 1a, this desegregation phenomenon could be related to the suppression of  $E_{elec}$  component (red dashed arrow at high  $pO_2$ ) under anodic conditions. However, as mentioned earlier, such energetics alone cannot explain the reaction that promotes the removal of dopant precipitates and their re-incorporation into the perovskite phase under anodic potential. In order to better understand this reversal reaction, we





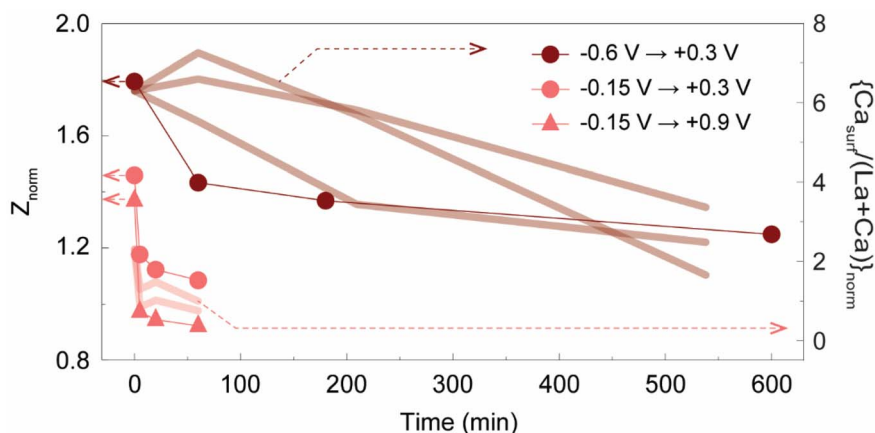
**Fig. 5** Activation of the LCM surface under different potentials. (a) After annealing an LCM film under  $-0.15$  V for 1 h at  $770$  °C (pink solid line), electrochemical impedance spectroscopy (EIS) was taken at every 5 or 10 min while applying  $+0.3$  V at the same temperature (blue dashed lines). Please note that all the EIS were consistently taken under  $0$  V. (b)  $-0.6$  V was first applied to LCM film for 1 h (brown solid line) and the potential was switched to  $+0.3$  V. During the annealing under  $+0.3$  V, EIS spectra were taken periodically under  $0$  V (blue dashed lines).

investigated the dopant re-incorporation process under various combinations of cathodic and anodic potentials.

### Dopant re-incorporation under different cathodic and anodic potentials

We further investigated dopant re-incorporation with the conventional potential method (Fig. S1a†) under different magnitudes of the initial cathodic potential and the subsequent anodic potential. Two different cathodic potentials,  $-0.15$  V or

$-0.6$  V, were applied for 1 h to first induce Ca segregation on LCM and three different anodic potentials,  $+0.3$ ,  $+0.45$ , or  $+0.9$  V, were thereafter applied for different time durations to reverse the dopant segregation. We found in our previous study<sup>34</sup> that applying high negative potential ( $\leq -0.6$  V) to the LCM surface resulted in significantly different surface morphology and amount of Ca segregation compared to those treated under low negative potential ( $\sim -0.15$  V). In this study, the two cathodic potentials,  $-0.15$  V and  $-0.6$  V, were chosen to investigate how such different initial states of LCM induced under these potentials affect the following dopant re-incorporation reaction under anodic potentials. Also, we avoided applying too high negative potential not to destroy the perovskite structure of LCM. After the polarization experiment, the amount of the resulting  $\text{Ca}_{\text{surf}}$  was quantified by XPS and plotted with respect to the time duration of anodic potential as shown in Fig. 2a. First, when dopant segregation is induced under  $-0.15$  V, more than twice the amount of  $\text{Ca}_{\text{surf}}/(\text{La} + \text{Ca})$  is formed at the surface of LCM (a pink star, y-axis on the left) compared to a reference sample annealed under  $+0.3$  V for 1 h (relative value of 1 in the plot). As can be seen in Fig. 2a (pink data points), when anodic potentials are sequentially applied to the sample, more than a half of the  $\text{Ca}_{\text{surf}}$  is removed in 5 min and the amount of  $\text{Ca}_{\text{surf}}$  gets closer to that on the reference sample over time ( $\{\text{Ca}_{\text{surf}}/(\text{La} + \text{Ca})\}_{\text{norm}}$  approaching 1). The corresponding XPS spectra are available in Fig. S3a.† Also, the amount of the  $\text{Ca}_{\text{surf}}$  removed under anodic potential was almost independent of the magnitude of the anodic potential (between  $+0.3$  (circles),  $+0.45$  (a square), and  $+0.9$  V (triangles)). These XPS results were further supported by scanning electron microscopy (SEM) measurement. Fig. 2b–d show that the number of the  $\text{CaO}_x$  precipitates on the surface decreases with increasing time spent at a potential of  $+0.9$  V and the surface of LCM starts to exhibit a grainy structure (inset of Fig. 2d). This surface morphology matches well with that of the as-prepared sample (Fig. S4†). The samples polarized at



**Fig. 6** Normalized impedance of the LCM film ( $Z_{\text{norm}}$ ) calculated from the electrochemical impedance spectra shown in Fig. 5.  $Z_{\text{norm}}$  is plotted with respect to the time duration of anodic potential;  $-0.15$  V (1 h)  $\rightarrow$   $+0.3$  V (pink circles),  $-0.15$  V (1 h)  $\rightarrow$   $+0.9$  V (pink triangles), and  $-0.6$  V (1 h)  $\rightarrow$   $+0.3$  V (brown circles).  $Z_{\text{norm}} = Z/Z_0$ ;  $Z$  = polarization resistance of LCM measured at  $0$  V after applying cathodic potential for 1 h and sequentially applying anodic potential for different time durations;  $Z_0$  = polarization resistance of LCM measured at  $0$  V before applying any potentials (right after the temperature reached to  $770$  °C and dwelling for 15 min). The temporal changes of the amount of  $\text{Ca}_{\text{surf}}$  (Fig. 2a) are plotted as thick solid lines (y-axis on the right) together with  $Z_{\text{norm}}$ .



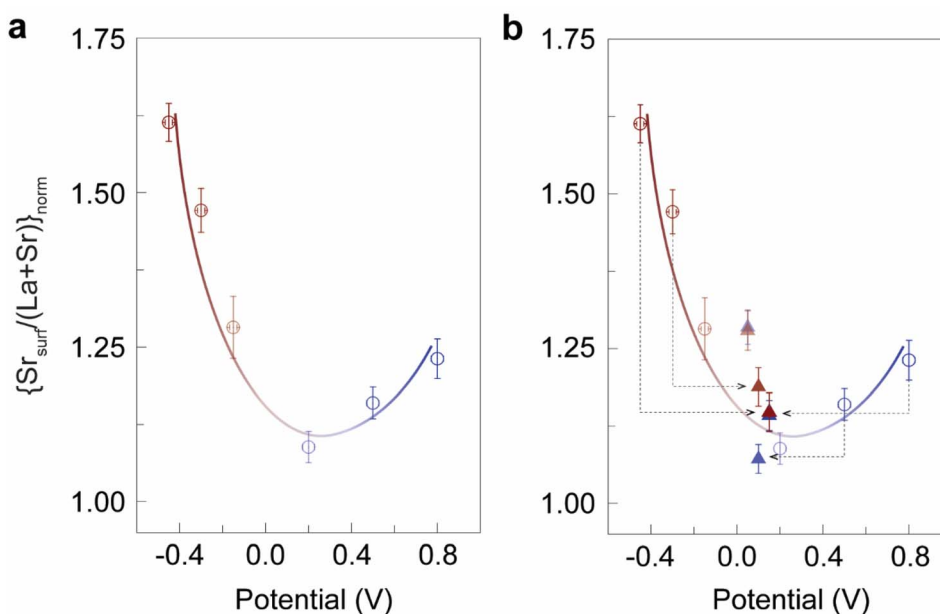


Fig. 7 (a) Segregation vs. potential on  $\text{La}_{0.8}\text{Sr}_{0.2}\text{MnO}_3$  (LSM). (b) Another set of LSM samples were annealed with the two-step approach to investigate the reversal of Sr segregation; black dashed arrows indicate the change of the applied local potential during the two-step polarization experiment. The amounts of the resulting  $\{\text{Sr}_{\text{surf}}/(\text{La} + \text{Sr})\}_{\text{norm}}$  after changing the potential to 0 to +0.2 V are plotted as triangles and overlapped with the data points in (a).

a different anodic potential ( $-0.15 \rightarrow +0.3$  V) also showed similar SEM results (Fig. S5a†).

When  $-0.6$  V was applied to LCM (brown star in Fig. 2a, y-axis on the right), more than 6-times higher amount of  $\text{Ca}_{\text{surf}}/(\text{La} + \text{Ca})$  was formed at the surface than the reference sample. Unlike the rectangular precipitates observed at the surface of the sample initially polarized at  $-0.15$  V (Fig. 2b), the surface of the sample initially polarized under  $-0.6$  V seems to be evenly covered by  $\text{CaO}_x$  layers (Fig. 2e). Such different surface morphologies of LCM formed under different potentials were also reported in our previous report.<sup>34</sup> Applying anodic potential could also reverse the dopant segregation that was induced under  $-0.6$  V, as evidenced by the decrease in the XPS signal from the  $\text{Ca}_{\text{surf}}$  component (Fig. S3b†). However, the amount of  $\text{Ca}_{\text{surf}}$  on this sample (brown circles in Fig. 2a) decreased at a slower rate than the sample polarized under  $-0.15$  V. Only after the prolonged application of anodic potential for  $>1$  h, the surface morphology of LCM started to change as the  $\text{CaO}_x$  layer was removed (Fig. 2g). The samples polarized at a different anodic potential ( $-0.6 \rightarrow +0.45$  V) also showed similar SEM results (Fig. S5b†). After applying anodic potential for 9 h, the amount of the  $\text{CaO}_x$  decreased down to one third of its initial amount. Different from the samples treated initially under  $-0.15$  V, the samples treated first at  $-0.6$  V gave different final amounts of  $\text{Ca}_{\text{surf}}$  when different anodic potentials were applied (Fig. 2a). Also, the  $\text{Ca}_{\text{surf}}$  amount was highest at  $+0.3$  V and lowest at  $+0.9$  V after 60 min, but such a trend was inverted at 540 min. We think such a different dopant re-incorporation behavior is attributed to the formation of microcracks in the sample pretreated under  $-0.6$  V. Further discussion on this is given in the later section.

Above results from XPS and SEM clearly show that the dopant precipitates at the LCM surface formed under cathodic potential can be removed by applying anodic potential. Also, the magnitude of the initial cathodic potential had a significant impact on the rate of dopant re-incorporation under anodic potential. Then, a question arises as to which chemical reactions induce the re-incorporation of the dopant precipitates into the perovskite phase. We believe that understanding the oxidation behavior of the LCM surface is key to answer this question, so we investigated the dominant redox-active species of the LCM surface under anodic potential. Then, we analyzed how the initial cathodic potential and the following anodic potential affect the oxidation kinetics of the LCM surface.

### Redox-active species of the LCM surface

To identify the dominant redox-active species in relation to Ca segregation at the LCM surface, we performed X-ray absorption spectroscopy (XAS) measurement in total-electron-yield (TEY) mode on the LCM thin films. The samples were annealed in 1 atm  $\text{O}_2$  at  $770$  °C for 1 h under potentials ranging from  $-0.75$  to  $+0.3$  V using conventional polarization method. Fig. 3a shows the O K-edge XAS spectra (*ex situ*) of these LCM samples. The peak at  $\sim 530$  eV, known as the hybridized Mn 3d–O 2p state,<sup>49,50,57,58</sup> indicates the presence of the oxygen holes in the valence band of LCM.<sup>57</sup> The inset of Fig. 3a shows that the intensity of this peak increases as higher anodic potential is applied. This means that the lattice oxygen at/near the LCM surface is oxidized under anodic potential, thus creating more oxygen holes. By contrast, the Mn L-edge XAS spectra in Fig. 3b show no noticeable peak shifts beyond the instrument error ( $\sim 0.1$  eV) or changes in the spectral shape. These results



indicate that O is the predominantly redox-active species in the oxidation of the LCM surface. When we measured the bulk of LCM in partial-fluorescence-mode (PFY) mode, the Mn L-edge spectra showed a shift of  $\sim 0.2$  eV as potential changed from  $-0.75$  to  $+0.3$  V, while O K edge did not show notable changes (Fig. S6†). Thus, the conventional understanding that the transition metal, Mn, is the redox-active species of manganite perovskite oxides is valid in the bulk of LCM,<sup>59</sup> but the surface of this perovskite oxide had oxygen as the redox-active species, consistent with another perovskite oxide surface reported earlier.<sup>50</sup>

To quantify the oxidation of the LCM surface, the area of the Mn–O peaks is calculated (see Fig. S7† for the details of the peak integration) and plotted with respect to the effective  $pO_2$  applied to the LCM thin films by potential as shown in Fig. 3c (colored circles, y-axis on the right). Each potential value was converted to the effective  $pO_2$  by using the Nernst equation (high effective  $pO_2$  is applied at anodic potential). We then compared the resulting curve to the oxygen non-stoichiometry curve of the LSM bulk (black dots and line in Fig. 3c) with the same amount of dopant as a reference. We assumed here that LCM would have similar oxygen non-stoichiometry with LSM. It can be found that the oxidation of the LCM surface (colored circles) shows similar  $pO_2$  dependence with the oxidation of the LSM bulk (black dots), having two steeply ascending regions and two plateau regions. The oxidation of the LSM bulk in the first ascending region at low  $pO_2$  ( $-20 < \log(pO_2) < -15$ ) is explained by the formation/annihilation of the oxygen vacancies. The oxidation of the LCM surface also increased at low  $pO_2$  presumably as oxygen vacancies are annihilated, but this occurred at higher  $pO_2$  ( $-15 < \log(pO_2) < -5$ ) than the LSM bulk. This mismatch between the two is thought to be because the surface of manganite perovskite oxides is more reducible than the bulk;<sup>60</sup> the reduction of the LCM surface occurs at higher  $pO_2$  than its bulk. After passing the first plateau region, both the LSM bulk and LCM surface start to exhibit the second ascending region at high  $pO_2$ ;  $-5 < \log(pO_2) < 0$  for the LSM bulk and  $0 < \log(pO_2) < 5$  for the LCM surface. It is known that the LSM bulk starts to accommodate oxygen excess at this high  $pO_2$  by creating cation vacancies,  $V''_{La}$  and  $V''_{Mn}$  (eqn (2)).<sup>43</sup> Based on the chemical similarity between LSM and LCM, it is reasonable to assume that the LCM surface also undergoes similar oxidation process, *i.e.*, creation of cation vacancies, during oxidation at this high  $pO_2$  ( $0 < \log(pO_2) < 5$ ). It is hence expected that the formation of the cation vacancies at the LCM surface would be related to the dopant re-incorporation reaction on the LCM surface under anodic potential ( $\log(pO_2) > 0$ , blue-colored region in Fig. 3c). As the LSM bulk and LCM surface get further oxidized, the creation of additional cation vacancies becomes energetically more demanding due to the significant repulsive forces among these defects.<sup>43</sup> Therefore, perovskite oxides can hardly undergo further oxidation and this is manifested as the second plateau in the non-stoichiometry curve of the LSM bulk ( $0 < \log(pO_2) < 7.5$ ) and the oxidation curve of the LCM surface ( $2.5 < \log(pO_2) < 6$ ) (Fig. 3c, blue-colored region). Based on this result, we performed mechanistic studies and quantitative analysis on the dopant re-

incorporation at the LCM surface, and the results are given in the following section.

### Mechanism behind dopant re-incorporation under anodic potential

Examining the relation between dopant re-incorporation rates and oxidation rates is helpful in resolving the mechanism of dopant incorporation under anodic potential. Hence, we measured *ex situ* XAS on the LCM samples used in Fig. 2 to investigate their oxidation kinetics. Fig. 4a and b show the resulting O K-edge XAS spectra of the LCM samples in total-electron-yield (TEY) mode. The sample annealed first under  $-0.15$  V (Fig. 4a) shows a fast increase in the Mn 3d–O 2p peak intensity shortly after applying  $+0.9$  V without further changes afterwards. Based on the results in Fig. 3, this can be interpreted that the sample underwent fast oxidation under  $+0.9$  V within a few minutes without further oxidation afterwards. On the other hand, the LCM film pre-annealed under  $-0.6$  V shows a gradual increase in the Mn 3d–O 2p peak intensity over 9 h under the same  $+0.9$  V potential (Fig. 4b). This means that this sample took much longer time to be oxidized compared to the one pre-annealed under  $-0.15$  V. Similar results are observed also under different anodic potentials (see Fig. S8 in ESI†). The temporal changes of the integrated area of the Mn 3d–O 2p peak are summarized in Fig. 4c (pink and brown data points, y-axis on the left), clearly showing that the samples kept initially under  $-0.15$  V (pink data points) underwent much faster oxidation than those kept under  $-0.6$  V (brown data points). Such a difference in the oxidation rates of these two sample sets is very similar to the rate of dopant re-incorporation behaviors shown in Fig. 2a. For easy comparison, the graph in Fig. 2a showing the temporal changes of the amount of  $Ca_{surf}$  is also plotted in Fig. 4c (thick solid lines, y-axis on the right). The resulting plots show that the oxidation of the LCM surface and dopant re-incorporation reaction on it have very similar time dependence. This implies that the oxidation of the LCM surface and the re-incorporation of  $CaO_x$  are coupled and take place simultaneously.

One possible question here would be, why the oxidation of the LCM surface should occur by re-incorporating the dopant precipitates, rather than it undergoing the typical oxidation reaction of manganite perovskite oxides (eqn (2)). We think this is because the near-surface region of LCM is highly A-site deficient after the dopant segregation under cathodic potential.<sup>61</sup> The oxidation process through eqn (2) accompanies the formation of A- and B-site vacancies and new unit cells at the surface. However, the formation of additional unit cells at the already A-site deficient surface would be highly unfavorable. Therefore, the surface would create only B-site vacancies while incorporating Ca, which is readily available in the surface  $CaO_x$  precipitates, into the A sites of the new unit cells. The relevant defect equation is given in eqn (3) in the next section.

In Fig. 2 and 4, we showed that the magnitude of the initial cathodic potential had a significant effect on the dopant re-incorporation and oxidation rates of the LCM surface when reversed to anodic potential. Both reactions occurred very

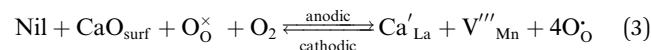


slowly at anodic potentials after the LCM samples were pre-annealed under high cathodic potential ( $-0.6$  V). We think this was because the dissolution of the  $\text{CaO}_x$  was kinetically slow on these samples, causing also a slow reoxidation of the LCM surface. The magnitude of the initial cathodic potential would affect the amount, size, and shape of the  $\text{CaO}_x$  precipitates (Fig. 2b and e), and their interfacial energy with the perovskite phase. These factors might have contributed to the sluggish dissolution of the large amount of  $\text{CaO}_x$  precipitates formed under  $-0.6$  V. The magnitude of the anodic potential did not have a significant impact on the dopant re-incorporation reaction when the LCM film was pretreated under  $-0.15$  V (Fig. 2a). This can be explained by the oxidation behavior of the LCM surface shown in Fig. 3c. The oxidation of the LCM surface becomes energetically demanding at high  $p\text{O}_2$  (blue points in Fig. 3c), and thus applying higher anodic potential than  $+0.3$  V would not lead to further oxidation of the LCM surface. Therefore, the amount of the  $\text{CaO}_x$  being re-incorporated into the perovskite phase through the oxidation reaction in eqn (3) does not change significantly with increasing the anodic potential. In contrast, the samples treated first at  $-0.6$  V gave different final amounts of  $\text{Ca}_{\text{surf}}$  when different anodic potentials were applied. Also, the trend of the  $\text{Ca}_{\text{surf}}$  amount at different anodic potentials was inversed as time increased. We think that this is because the amount of  $\text{Ca}_{\text{surf}}$  can also be largely affected by the kinetics of the dopant re-incorporation reaction or microstructural changes in the film. The film treated under  $+0.9$  V after  $-0.6$  V initially showed the lowest amount of  $\text{Ca}_{\text{surf}}$  at 60 min because of the fast dopant re-incorporation reaction induced by the large anodic potential,  $-0.6$  V  $\rightarrow$   $+0.9$  V. However, due to the large potential change, microcracks gradually formed in this film and dopant re-incorporation reaction no longer occurred over the whole film surface. This is manifested by the nominal decrease in the dopant re-incorporation rate over time seen in Fig. 2a (brown triangles). In the LCM film treated under  $+0.3$  V after  $-0.6$  V (brown circles), the number of microcracks, if any, was far less, and hence, the dopant re-incorporation rate was almost constant over 9 h. The LCM film treated under  $+0.45$  V after  $-0.6$  V (brown squares) showed intermediate behavior. Such a change of trend in the  $\text{Ca}_{\text{surf}}$  amount was not observed when  $-0.15$  V was applied first because (1) the magnitude of the potential change was not huge (max.  $-0.15$  V  $\rightarrow$   $+0.9$  V) and (2) dopant re-incorporation occurred much faster within an hour before the formation microcracks, if any, gave any noticeable effects.

We have also investigated the bulk oxidation of the LCM samples initially polarized under  $-0.6$  V, to see if it has any relation to their dopant re-incorporation reactions. In their bulk Mn spectra (Fig. S9g and h $\dagger$ ), we observed a noticeable peak shift of around  $0.2$ – $0.3$  eV upon the application of anodic potential. The amount of the shift is almost identical to the peak position difference observed between the LCM sample annealed with single polarization approach under  $-0.75$  and that under  $0.3$  V (Fig. S6a $\dagger$ ), thus meaning that the LCM bulk was fully oxidized. Also, it should be noted that the shift occurred much faster ( $<1$  h) than both the oxidation rate of the surface O ( $\sim 9$  h) and the rate of  $\text{CaO}_x$  being removed. This result

indicates that the bulk oxidation of these LCM samples was not related to the sluggish oxidation and dopant re-incorporation kinetics at the surface. Also, this further supports our above argument that the oxidation of the LCM surface does not undergo the typical oxidation process (eqn (2), where Mn is the redox-active species), but rather involves the reincorporation of the aliovalent dopant, Ca. If the reaction in eqn (2) had occurred at the surface, the surface oxidation should have been as fast as the bulk, and independent from the dopant incorporation rate.

Based on the results presented above, we formulated a surface defect reaction that explains dopant segregation under cathodic and re-incorporation under anodic potential:



( $\text{Ca}'_{\text{Mn}} = \text{Ca}$  on the La lattice site,  $\text{O}_{\text{O}}^{\times} =$  oxygen hole in the Mn 3d–O 2p band)

Eqn (3) comprises defect reactions that lanthanum manganites undergo under anodic conditions: formation of cation vacancies ( $\text{V}'''_{\text{Mn}}$ ) and oxygen holes ( $\text{O}_{\text{O}}^{\times}$ ) and re-incorporation of the dopant precipitates ( $\text{CaO}_{\text{surf}}$ ) into the perovskite phase. It is derived from a linear combination of eqn (1) and (2), and modification of the redox-active species from Mn to O (eqn (S1)–(S4) in ESI $\dagger$ ). Basically, it describes that dopant re-incorporation under anodic potential occurs by forming new perovskite unit cells that incorporate  $\text{CaO}_x$  and oxygen molecules into the A and O sites and that have B-site vacancies (Fig. 4d). In this scenario, the oxidation of the LCM surface can be impeded if the dissolution kinetics of  $\text{CaO}_{\text{surf}}$  is slow as explained above. The formation of the B-site vacancies ( $\text{V}'''_{\text{Mn}}$ ) under anodic conditions is in line with a recent theoretical and computational work from our group which predicted the formation of a large amount of B-site vacancies in  $\text{La}_{0.8}\text{Sr}_{0.2}\text{MnO}_3$  under anodic conditions.<sup>38</sup> In other papers, it was reported that undoped  $\text{LaMnO}_3$  contained similar amounts of A- and B-site vacancies under oxidizing conditions,<sup>44,62</sup> and also, some other perovskites are known to exhibit B-site deficiency such as  $\text{Pr}_{0.5}\text{Sr}_{0.5}\text{Co}_{1-y}\text{O}_3$  (10%)<sup>63</sup> and  $\text{Sr}_{0.9}\text{La}_{0.1}\text{Ti}_{1-y}\text{O}_3$  (6%).<sup>64</sup> Also, we point out the redox-active species on the LCM surface in eqn (3). Mn is thought to be the redox active species in doped  $\text{LaMnO}_3$ ,<sup>44,65</sup> and we have also assumed that in our previous theoretical work connecting defect reactions to surface segregation.<sup>38</sup> This knowledge comes from bulk defect equilibria studies on the manganite system.<sup>44,65</sup> From the XAS measurements, we found that Mn is indeed redox-active in the bulk, but at the surface it is the lattice oxygen that was redox-active.

Lastly, we investigated if dopant re-incorporation under anodic conditions can occur not only by applying anodic potential, but also by exposing the LCM surface to oxidizing gas environment. First, we observed that a significant amount of  $\text{Ca}_{\text{surf}}$  is formed at the LCM surface after annealing it in 5%  $\text{H}_2/\text{N}_2$  (reducing condition) for 1 h (Fig. S10 $\dagger$ ). This matches well with the above result in Fig. 2 that Ca segregation is induced under cathodic potential. To test dopant re-incorporation under oxidizing environment, we annealed another LCM film in 5%  $\text{H}_2/\text{N}_2$  for 1 h, switched the gas to synthetic air, and annealed it additionally for 3 h. As a result, this sample showed



a significantly reduced amount of  $\text{Ca}_{\text{surf}}$  compared to the one annealed only in  $\text{H}_2$ . This result clearly shows that dopant re-incorporation can also occur chemically at oxidizing conditions, and its applicability is not limited to the electrochemically anodic conditions.

### Activation of the LCM surface under anodic potential

We then investigated if the removal of  $\text{CaO}_x$  by reversing dopant segregation under anodic potential activates the oxygen exchange kinetics on the LCM surface. For this, we performed electrochemical impedance spectroscopy (EIS) measurements on LCM films under the same experimental conditions as described above in Fig. 2 and 4. Black dashed lines in Fig. 5a and b are the EIS spectra of the as-prepared LCM film under 0 V which comprise two semicircles with the peak frequencies of 6 and 1000 Hz. Based on the previous EIS studies on LSM, the low-frequency arc is attributed to the oxygen exchange reaction at the surface ( $\text{O}_{2,\text{gas}} + 4e^- \leftrightarrow 2\text{O}^{2-}_{\text{oxide}}$ ) and the high frequency arc is attributed to the oxygen ion transport processes along the grain boundaries and/or bulk.<sup>36,39,41,66–68</sup> After cathodic potentials ( $-0.15$  or  $-0.6$  V) are applied to the LCM film, its total impedance measured under 0 V grows gradually over time as can be seen in Fig. S11a.† This result coincides well with the XPS and SEM results in Fig. 2a, b and e, showing the formation of insulating  $\text{CaO}_x$  after applying cathodic potential;  $\text{CaO}_x$  layers at the surface blocks the active site for surface oxygen exchange reactions and increases the impedance of the sample. After applying cathodic potential for 1 h, anodic potential was then applied to remove the  $\text{CaO}_x$  and the impedance of the cell was measured over time consistently under 0 V. Fig. 5a and b (blue dashed lines) show that the impedances of both samples, initially annealed under  $-0.15$  and  $-0.6$  V, decrease as  $\text{CaO}_x$  is removed under anodic potential. This clearly indicates that applying anodic potential recovers the degraded oxygen exchange activity of the LCM surface under cathodic potential.

The polarization resistance of the LCM films ( $Z$ , difference between the low- and high-frequency intercepts of the semicircles) was normalized by their initial polarization resistance before applying any potentials ( $Z_0$ ) and the resulting  $Z_{\text{norm}}$  ( $= Z/Z_0$ ) was plotted with respect to the time duration of anodic potential (pink and brown data points in Fig. 6). In the case of the samples annealed initially under  $-0.15$  V (pink data points),  $Z_{\text{norm}}$  sharply decrease close to unity (the impedance of LCM before applying any potentials) within a few minutes after applying anodic potentials. Interestingly, even after a prolonged time period of degradation process (10 h), applying anodic potential recovered 50% of the LCM's initial oxygen exchange activity just in a few minutes (Fig. S12†). On the other hand, for the samples initially polarized under  $-0.6$  V (brown data points),  $Z_{\text{norm}}$  decrease in a far slower rate under anodic potential than those polarized under  $-0.15$  V, and they do not fully recover their initial impedance even after 10 h. Such difference observed in the two sets of the LCM samples matches very well with the different rates of dopant re-incorporation shown in Fig. 2a. For easier comparison, the plots in Fig. 2a are overlapped with  $Z_{\text{norm}}$  in Fig. 6 (thick solid lines, y-axis on

the right). The conformity between the temporal changes of  $Z_{\text{norm}}$  and  $\text{Ca}_{\text{surf}}$  clearly indicates that the removal of  $\text{CaO}_x$  under anodic potential leads to the activation of the oxygen kinetics in LCM.

Lastly, we investigated the reversibility of the passivation and activation processes by the formation and removal of  $\text{CaO}_x$  layers, respectively. Four to five potential cycles under  $-0.15$  V  $\leftrightarrow$   $+0.9$  V,  $-0.15$  V  $\leftrightarrow$   $+0.3$  V, and  $-0.6$  V  $\leftrightarrow$   $+0.3$  V were applied to LCM films and the resulting EIS spectra are given in Fig. S13.† In all cases, the impedance changes were reversible in terms of both the shape and magnitude of the EIS spectra. However, when the change of potential was large, e.g.,  $-0.6$  V  $\leftrightarrow$   $+0.9$  V, the EIS spectrum became irreversible. Fig. S14† shows the irreversible changes of EIS spectrum after finishing the first potential cycle of  $-0.6$  V (1 h)  $\leftrightarrow$   $+0.9$  V (10 h). According to the previous studies on LSM thin films, microcracks or holes can form under high anodic potential because of the high effective  $p\text{O}_2$  applied to the film, and this can lead to the decrease in the impedance of the film as the cracks or holes serve as additional surface and diffusion pathways.<sup>5,36,69</sup> Likewise, we could also observe the formation of microcracks or delamination of the LCM films (Fig. S15†) after undergoing a large potential change from  $-0.6$  V to  $+0.45$  V or  $+0.9$  V.

### Comparison to Sr-substituted $\text{LaMnO}_3$

Though LCM was used as a primary model system in this work, it is important to investigate whether the results can be transferred to LSM which is much more widely used as the air electrode in SOFC/EC cells. To this end, we investigated the dopant segregation and re-incorporation behaviors of LSM in a similar way that we did above for LCM in Fig. 1b. Again, LSM thin-film samples were annealed at 770 °C in 1 atm  $\text{O}_2$  for 1 h under a wide range of potentials by using the lateral polarization method;  $-0.45$ ,  $-0.3$ ,  $-0.15$ ,  $+0.3$ ,  $+0.6$ , and  $+0.9$  V. XPS core-level spectra of Sr, specifically Sr 3d, were deconvoluted into the surface ( $\text{Sr}_{\text{surf}}$ , red envelopes in Fig. S16†) and lattice components ( $\text{Sr}_{\text{latt}}$ ) to quantify the surface  $\text{SrO}_x$ -like phase which was formed as a result of dopant segregation. The amount of  $\text{Sr}_{\text{surf}}$  was then divided by the total A-site cations and the resulting  $\text{Sr}_{\text{surf}}/(\text{La} + \text{Sr})$  was normalized by the  $\text{Sr}_{\text{surf}}/(\text{La} + \text{Sr})$  of the as-prepared LSM sample. The resulting  $\{\text{Sr}_{\text{surf}}/(\text{La} + \text{Sr})\}_{\text{norm}}$  are plotted with respect to the applied local potential in Fig. 7a (open circles). The resulting segregation behavior of LSM is consistent with our previous results;<sup>34</sup> dopant segregation occurs also under anodic potential when the dopant size is larger than the host cation, La ( $R_{\text{Ca}} \approx R_{\text{La}} < R_{\text{Sr}}$ ), and hence the dopant has a high  $E_{\text{ela}}$ . In order to examine the dopant re-incorporation in LSM, another set of samples were annealed with the two-step polarization approach. The same potentials as above, from  $-0.45$  to  $+0.9$  V, were first applied for 1 h, followed by application of 0.0 to  $+0.2$  V for another hour. This value of anodic potential corresponds to the minimum of the Sr segregation graph in Fig. 7a. The resulting  $\{\text{Sr}_{\text{surf}}/(\text{La} + \text{Sr})\}_{\text{norm}}$  were again quantified by XPS (Fig. S16†) and plotted together with the data points in (a) (Fig. 7b, triangles). Notably, applying 0.0 to  $+0.2$  V could remove the dopant segregation in LSM induced



under both high cathodic and anodic potentials (dashed arrows in Fig. 7b). The XPS results were further supported by SEM measurements on the LSM films (Fig. S17 and S18†); surface particles, assumed as  $\text{SrO}_x$ , are removed after annealing under 0.0 to +0.2 V. The observed two different types of segregation (cathodic and anodic) and re-incorporation at 0 to +0.2 V, might explain the contradicting segregation behaviors of LSM reported in the literature; Sr segregation under both cathodic and anodic potential, and Sr re-incorporation under both anodic and cathodic potential. We also performed similar experiments with  $\text{La}_{0.8}\text{Ba}_{0.2}\text{MnO}_3$  (LBM) and showed that Ba segregation on LBM could also be removed under 'relatively' anodic potential (Fig. S19†). The results from LSM and LBM were reproducible with conventional polarization method as well (Fig. S20 and S21†).

The two different types of dopant segregation/precipitation and re-incorporation on LSM shown in Fig. 7 can be explained by the defect reactions in eqn (1) and (3), which become predominant at different potential regimes. In the anodic-potential regime (+0.2 to +0.8 V),  $\text{SrO}_x$  precipitation and re-incorporation can be explained by the defect reaction in eqn (1). When high anodic potential is applied (oxidizing conditions), the formation of  $\text{SrO}_x$  is promoted according to the backward reaction of eqn (1) (positive-slope branch of the segregation graph in Fig. 7a). When 'small anodic, and relatively cathodic' potential (0 to +0.2 V) is applied thereafter, dopant re-incorporation reaction occurs as explained by the forward reaction of eqn (1) (Fig. 7b). On the other hand, in the strongly cathodic-potential regime (−0.45 to +0.2 V), the defect reaction in eqn (3) becomes predominant. When high cathodic potential is applied (reducing conditions), the formation of  $\text{SrO}_x$  is promoted according to the backward reaction of eqn (3) (negative-slope branch of the segregation graph in Fig. 7a). When low anodic potential (0 to +0.2 V) is applied thereafter, dopant re-incorporation reaction occurs as explained by the forward reaction of eqn (3) (Fig. 7b). In our previous studies,<sup>19,34</sup> we used the electrostatic and elastic energy of dopant to explain the predominant driving force of the dopant segregation under reducing and oxidizing conditions. However, these energy-based drivers of segregation rather limit us to the 'enrichment' of the dopant at/near the surface, and does not account for how dopant oxide 'precipitation' and redissolution can occur. Here, we show that we must consider the predominant defect reactions (eqn (1) and (3)) in the reducing and oxidizing regimes to account for dopant oxide phase precipitation and redissolution, as seen on LCM and LSM.

## Summary

In summary, we established that the mechanism of the dopant re-incorporation at a doped manganite surface under anodic potential involves the dominant defect equilibria at highly oxidizing conditions. Based on XPS and XAS results and by taking LCM as a model system, we have shown that the dopant re-incorporation reaction on LCM is coupled with its surface oxidation by creating B-site cation-deficient perovskite unit cells under high- $p\text{O}_2$  conditions. In addition, oxidation of the LCM

surface was mediated by the redox-active surface oxygen anions rather than the surface Mn cations. The EIS experiments showed that applying anodic potential improved the oxygen exchange kinetics of LCM in a short time (<5 min) as it removed the insulating  $\text{CaO}_x$  at the surface. We also showed that the results from LCM can be transferred to LSM which is one of the state-of-the-art air electrodes in SOF/EC. Importantly, the new defect reaction proposed in this paper reconciles the contradicting experimental results on dopant segregation and re-incorporation reactions in the literature. We believe that this anodic activation pathway of lanthanum manganites would be applicable to other perovskite oxides which also exhibit a large cation deficiency under oxidizing conditions.<sup>70</sup> Our findings indicate that operating perovskite oxides in a reversible solid oxide cell<sup>5</sup> would be advantageous in achieving their long-term stability by periodically removing dopant segregation under alternating polarizations.

## Author contributions

D. K. and B. Y. conceived the original ideas. D. K. and B. Y. designed the research. D. K. performed all the experiments. A. H. and I. W. helped the beamline experiments at beamline 23-ID-2 of the National Synchrotron Light Source II in BNL. D. K. wrote the manuscript with contributions from all authors.

## Conflicts of interest

There are no conflicts of interest to declare.

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