



Cite this: *RSC Adv.*, 2018, 8, 26448

A first-principles study on divergent reactions of using a Sr₃Fe₂O₇ cathode in both oxygen ion conducting and proton conducting solid oxide fuel cells†

Wenzhou Tan,^a Daoming Huan,^a Wenqiang Yang,^a Nai Shi,^a Wanhua Wang,^a Ranran Peng,^{id} Xiaojun Wu^{id} and Yalin Lu^{*abcd}

Exploring mechanisms for sluggish cathode reactions is of great importance for solid oxide fuel cells (SOFCs), which will benefit the development of suitable cathode materials and then accelerate cathode reaction rates. Moreover, possible reaction mechanisms for one cathode should be different when operating in oxygen ion conducting SOFCs (O-SOFC) and in proton conducting SOFCs (P-SOFCs), and therefore, they lead to different reaction rates. In this work, a Ruddlesden–Popper (R–P) oxide, Sr₃Fe₂O₇ (SFO), was selected as a promising cathode for both O-SOFCs and P-SOFCs. Using the first-principles approach, a microscopic understanding of the O₂ reactions over this cathode surface was investigated operating in both cells. Compared with La_{0.5}Sr_{0.5}Co_{0.25}Fe_{0.75}O₃ (LSCF), the low formation energies of oxygen vacancies and low migration energy barriers for oxygen ions in SFO make oxygen conduction more preferable which is essential for cathode reactions in O-SOFCs. Nevertheless, a large energy barrier (2.28 eV) is predicted for oxygen dissociation reaction over the SFO (001) surface, while there is a zero barrier over the LSCF (001) surface. This result clearly indicates that SFO shows a weaker activity toward the oxygen reduction, which may be due to the low surface energies and the specific R–P structure. Interestingly, in P-SOFCs, the presence of protons on the SFO (001) surface can largely depress the energy barriers to around 1.46–1.58 eV. Moreover, surface protons benefit the oxygen adsorption and dissociation over the SFO (001) surface. This result together with the extremely low formation energies and migration energy barriers for protons seem to suggest that SFO could work more effectively in P-SOFCs than in O-SOFCs. It's also suggested that too many protons at the SFO surface will lead to high energy barriers for the water formation process, and thus that over-ranging steam concentrations in the testing atmosphere may have a negative effect on cell performances. Our study firstly and clearly presents the different energy barriers for one cathode performing in both O- and P-SOFCs according to their different working mechanisms. The results will be helpful to find the constraints for using cathodes toward oxygen reduction reactions, and to develop effective oxide cathode materials for SOFCs.

Received 13th May 2018
 Accepted 16th July 2018

DOI: 10.1039/c8ra04059a

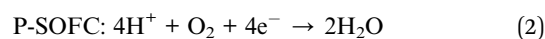
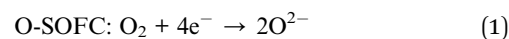
rsc.li/rsc-advances

Introduction

As green and efficient energy conversion devices, solid oxide fuel cells (SOFCs) have attracted much attention nowadays. According to the conducting ions in their electrolytes, SOFCs can be divided

into two categories: oxygen ion conducting SOFCs (O-SOFCs) and proton conducting SOFCs (P-SOFCs). Interestingly, in spite of their conducting ions in electrolytes (protons or oxygen ions), lots of experiments have also suggested that polarization resistances corresponding to the sluggish cathode reactions are one of the main factors to limit the electrochemical performance of SOFCs. As a result, in the past decades, great efforts have been devoted to develop suitable cathode materials to accelerate the cathode reaction rate and to improve the cell performance.^{1,2}

As shown in eqn (1) and (2), cathode reactions for O-SOFCs and P-SOFCs are quite different and thus have different requirements on their cathode materials.



^aCAS Key Laboratory of Materials for Energy Conversion, Department of Materials Science and Engineering, University of Science and Technology of China, Hefei, 230026 Anhui, China. E-mail: pengrr@ustc.edu.cn; yllu@ustc.edu.cn

^bSynergetic Innovation Center of Quantum Information & Quantum Physics, University of Science and Technology of China, Hefei, Anhui 230026, China

^cSynchrotron Radiation Laboratory, University of Science and Technology of China, Hefei 230026, P. R. China

^dHefei National Laboratory of Physical Science at the Microscale, University of Science and Technology of China, Hefei, 230026 Anhui, China

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8ra04059a



For O-SOFCs, good oxygen ion and electron conduction are essential to accelerate cathode reactions. To meet such demands, lots of efficient composite cathodes, for example $\text{La}_{0.8}\text{Sr}_{0.2}\text{MnO}_{3-\delta}\text{-Y}_{0.88}\text{Zr}_{0.12}\text{O}_{2-\delta}$ (LSM-YSZ),³ $\text{La}_{0.6}\text{Sr}_{0.4}\text{Co}_{0.2}\text{-Fe}_{0.8}\text{O}_{3-\delta}\text{-Sm}_{0.2}\text{Ce}_{0.8}\text{O}_{1.9}$ (LSCF-SDC),⁴ and $\text{Ba}_{0.5}\text{Sr}_{0.5}\text{Co}_{0.8}\text{Fe}_{0.2}\text{-O}_{3-\delta}\text{-Sm}_{0.2}\text{Ce}_{0.8}\text{O}_{1.9}$ (BSCF-SDC),⁵ have been investigated, where the electrolytes were added to improve oxygen ion conduction or to enlarge length of triple phase boundaries. While for P-SOFCs, proton conduction in addition to electron conduction is urgently needed, and conduction of oxygen ions seems not as vital as in O-SOFCs where oxygen ions need to diffuse to electrolytes and finally to the anode. For P-SOFCs, in addition to those composite cathodes, such as $\text{La}_{0.6}\text{Sr}_{0.4}\text{Co}_{0.2}\text{-Fe}_{0.8}\text{O}_{3-\delta}\text{-BaZr}_{0.1}\text{Ce}_{0.7}\text{Y}_{0.2}\text{O}_{3-\delta}$ (LSCF-BZCY),⁶ $\text{Sm}_{0.5}\text{Sr}_{0.5}\text{-CoO}_{3-\delta}\text{-BaZr}_{0.1}\text{Ce}_{0.7}\text{Y}_{0.2}\text{O}_{3-\delta}$ (SSC-BZCY),⁷ single phase cathodes, including $\text{BaCo}_{0.4}\text{Fe}_{0.4}\text{Zr}_{0.1}\text{Y}_{0.1}\text{O}_{3-\delta}$,⁸ $\text{BaCo}_{0.4}\text{Fe}_{0.4}\text{-Zr}_{0.2}\text{O}_{3-\delta}$,⁹ $\text{BaCe}_{0.5}\text{Fe}_{0.5}\text{O}_{3-\delta}$,¹⁰ $\text{BaZr}_{0.6}\text{Co}_{0.4}\text{O}_{3-\delta}$,¹¹ and $\text{Sr}_3\text{Fe}_2\text{-O}_{7-\delta}$,¹² were also designed and expected to function better than those composites cathodes because of their potential proton-electron mixed conduction which could enlarge the active reaction area to the cathode surface instead of TPB. Proton conduction in these materials are formed *via* the incorporation of steam molecules into oxygen vacancies, as shown in eqn (3).



In the past, acceptable performances were achieved with above cathodes. For example, applying $\text{BaCo}_{0.4}\text{Fe}_{0.4}\text{Zr}_{0.2}\text{O}_{3-\delta}$, peak power densities of 405 mW cm^{-2} were achieved at $500 \text{ }^\circ\text{C}$ for P-SOFCs when using a $\text{BaCe}_{0.7}\text{Zr}_{0.1}\text{Y}_{0.1}\text{Yb}_{0.1}\text{O}_{3-\delta}$ electrolyte.⁹ Nevertheless, despite of the improved cell performances, in the microscopic view detailed cathode reaction mechanisms and the influences are still frustrating for both O-SOFCs and P-SOFCs. Especially, no discussions with regarding to use one cathode material in both O-SOFCs and P-SOFCs have ever been proposed yet.

It has been generally accepted that the first-principles calculation could provide useful information on revealing and understanding detailed mechanism for catalytic reactions. For instance, Y. Choi *et al.* have studied oxygen reduction barriers for $\text{La}_{0.5}\text{Sr}_{0.5}\text{MnO}_3$ (LSM0.5) and proposed a fast O_2 reduction on LSM0.5 with nonexistence of transition-state barriers.¹³ Yueh-Lin Lee *et al.* have demonstrated that the experimentally measured area specific resistance and oxygen surface exchange of solid oxide fuel cell cathode strongly correlate with the theoretical calculation data.¹⁴ Sihyuk Choi *et al.* have studied the possible elementary pathway for the oxygen reduction reaction on $\text{PrBa}_{0.5}\text{Sr}_{0.5}\text{Co}_{2-x}\text{Fe}_x\text{O}_5$ by DFT analysis, and suggested that the most attractive properties of these materials are the presence of pore channels in the [PrO] and [CoO] planes that provide paths for fast oxygen transport that accelerates the surface oxygen exchange kinetics.¹ We have also explored bulk proton transporting and oxygen reduction behaviors on $\text{BaZr}_{0.75}\text{Co}_{0.25}\text{O}_3$ cathodes and found that the $\text{BaZr}_{0.75}\text{Co}_{0.25}\text{O}_3$ a promising cathode for P-SOFCs because of its low proton formation energy, low diffusion energy barrier and its low energy barriers toward cathode reactions at the presence of protons.¹⁵ Recently, we found that $\text{Sr}_3\text{Fe}_2\text{O}_7$ (Ruddlesden-

Popper oxides) has low formation energies and diffusion energy barriers for both oxygen vacancies and protons in bulk model, which make it suitable for both O-SOFCs and P-SOFCs.^{12,16}

In this work, we performed a first-principles study on the oxygen reduction process over the $\text{Sr}_3\text{Fe}_2\text{O}_7$ surfaces which acts as a single phase cathode in both O-SOFCs and P-SOFCs, as shown in Fig. 1. Feasible reaction paths over the $\text{Sr}_3\text{Fe}_2\text{O}_{7-\delta}$ cathode were explored in both operating modes, and effects of proton presentation over oxygen adsorption and dissociation reactions were evaluated. Also, the impact of R-P structure on the oxygen reduction reactions were compared with that of perovskite structure using $\text{La}_{0.5}\text{Sr}_{0.5}\text{Co}_{0.25}\text{Fe}_{0.75}\text{O}_3$, a classic cathode material, as a reference.

Computational methods

Theoretical study was conducted to reveal the O_2 reduction reaction process over $\text{Sr}_3\text{Fe}_2\text{O}_7$ (001) surface using the projector augmented wave (PAW) method, implemented in the Vienna ab initio simulation package (VASP).¹⁷⁻¹⁹ Perdew-Berke-Ernzerhoff (PBE) generalized gradient approximation was adapted to treat the exchange-correlation effects.²⁰ Kinetic energy cutoff set was 450 eV and energy convergence criterion was set to $10^{-5} \text{ eV per atom}$. Structural optimization was carried out until the Hellmann-Feynman force on each atom was lower than $0.03 \text{ eV } \text{\AA}^{-1}$. All calculations were spin-polarized,^{21,22} and the ferromagnetic (FM) states instead of antiferromagnetic (AFM) were used as indicated by Tadashi Ota *et al.*²³ To predict the electronic structures more precisely, DFT + U method²⁴ was applied in all our calculations. Here, the value of $U_{\text{eff}} = U - J = 5.3 \text{ eV}$ was applied for Fe ions, and the Coulomb (U) and exchange (J) parameters were not taken into account separately.^{12,25} An elementary crystal structure containing 24-atoms was established for SFO, as shown in Fig. 2(a). Parameters of the tetragonal unit cell are calculated as $a = b = 3.8873 \text{ \AA}$ and $c = 20.2442 \text{ \AA}$, in good agreement with experimental results (space group $I4/mmm$, $a = b = 3.8668 \text{ \AA}$ and $c = 20.1737 \text{ \AA}$ (ref. 26)).

As previously reported, (001) surface is more stable than (100), (110), (111), and (011) surfaces in SFO,¹⁶ and therefore, five ten-layer (001) slabs, as shown in Fig. S1,† were cleaved to simulate the terminal surface and to intensively seek the suitable terminal surface for later reaction mechanism investigation. Here, a $5 \times 5 \times 1$ k -points mesh was adapted using the Monkhorst-Pack scheme to sample the Brillouin zone integration.²⁷ To investigate

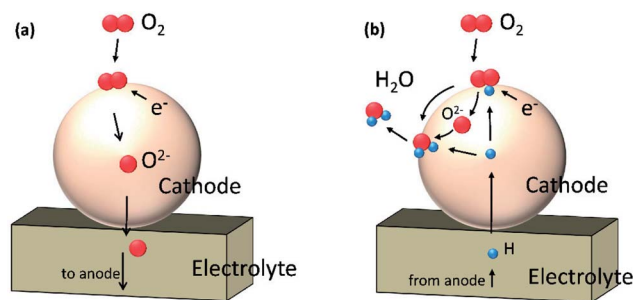


Fig. 1 Schematic diagrams of cathode reactions on $\text{Sr}_3\text{Fe}_2\text{O}_{7-\delta}$ operating in O-SOFCs (a) and P-SOFCs (b), respectively.



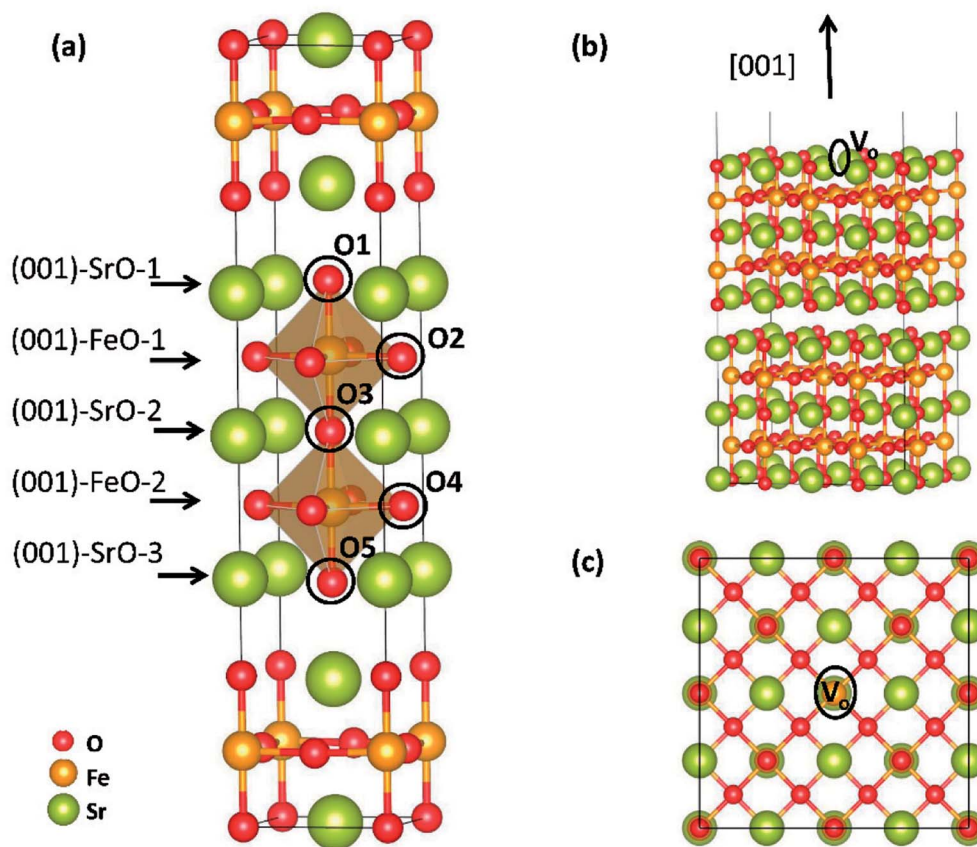


Fig. 2 (a) Elementary unit cell structure of SFO with 24 atoms, (b) side and (c) top views of the Sr₃Fe₂O₇ (001) surface with oxygen vacancy. The O1–O5 refer to possible locations where oxygen atoms or proton pass through during the surface diffusion.

O₂ adsorption and dissociation reactions, a $2\sqrt{2} \times 2\sqrt{2}$ surface (001) slab was built to make a bigger surface system with 192 atoms (as shown in Fig. 2(b) and (c)). To save the computing resources, these calculations were performed with $2 \times 2 \times 1$ *k*-points grid. Transition states (TSs) of oxygen dissociation and ion migration were calculated through the climbing image nudged elastic band (CI-NEB) method.²⁸ All the slabs were separated perpendicularly by a 15 Å vacuum space, and the bottom-two layers were fixed to their bulk position, while the other atoms were allowed to fully relax. Dipole correction was applied to the direction perpendicular to the surface.

Adsorption energy of reactants was calculated using $E_{\text{ads}} = E_{[\text{slab}+\text{reactant}]} - E_{[\text{slab}]} - E_{[\text{reactant}]}$, where the $E_{[\text{slab}+\text{reactant}]}$, $E_{[\text{slab}]}$ and $E_{[\text{reactant}]}$ denoted the total energy of the reactant adsorbed surface, the pristine surface and the gas phase reactant, respectively. So, a positive value of E_{ads} suggests that the adsorption reaction is endothermic, while a negative one indicates an exothermic reaction. Formation energy (E_{vac}) of an oxygen vacancy was calculated according to $E_{\text{vac}} = E_{[\text{defective}]} + 1/2E_{[\text{O}_2]} - E_{[\text{perfect}]}$, where $E_{[\text{defective}]}$ and $E_{[\text{perfect}]}$ were the total energy of defective and perfect Sr₃Fe₂O₇ surfaces, respectively. Proton formation energy (E_{hyd}) was calculated using the equation: $E_{\text{hyd}} = 2E_{\text{OH}} - E_{\text{vac}} + E_{\text{pt}}$, where E_{OH} was the energy associated with substitution of O²⁻ with an OH⁻ group, E_{vac} was the energy needed to create an oxygen vacancy, and E_{pt} was the energy of the gas phase reaction: $\text{H}_2 + 1/2\text{O}_2 = \text{H}_2\text{O}$. E_{pt} was

calculated to be -2.52 eV. The ground state triplet O₂ was used for oxygen adsorption and dissociation, and the value of evaluated bond length was fit as 1.233 Å, which agreed with the experimental results of 1.207 Å.

Results and discussion

Properties of Sr₃Fe₂O₇

Electronic density of state (DOS) of SFO, with the smear value of 0, and the sigma value of 0.1 in VASP, are displayed in Fig. 3. Fe 3d electrons arise significantly around the Fermi energy level and hybridize with the 2p electrons of the O, indicating an increase of the charge-carrier concentration. The conducting characteristics of Sr₃Fe₂O₇ is half-metallic, in good agreement with previous experimental and theoretical investigations on Sr₃Fe₂O₇, facilitating the electron conduction.^{12,23,29,30}

To explore stable (001) surface for O₂ reduction investigation, five types of (001) terminal surfaces can be claved, as illustrated in Fig. 2 and S1† and noted as SrO-1, SrO-2, SrO-3, FeO-1 and FeO-2 terminal surfaces, respectively. Surface energies (E_{sur}) of SrO-1, SrO-2, SrO-3, FeO-1 and FeO-2 terminal surfaces are calculated as 7.69, 12.98, 19.06, 18.90 and 13.94 eV Å⁻², respectively. Among these surfaces, SrO-1 terminal surface has much lower E_{sur} than the others, indicating that it is the most stable one. And therefore, our later investigations on O₂ reduction reaction are mainly focused on SrO-1 terminal surface.



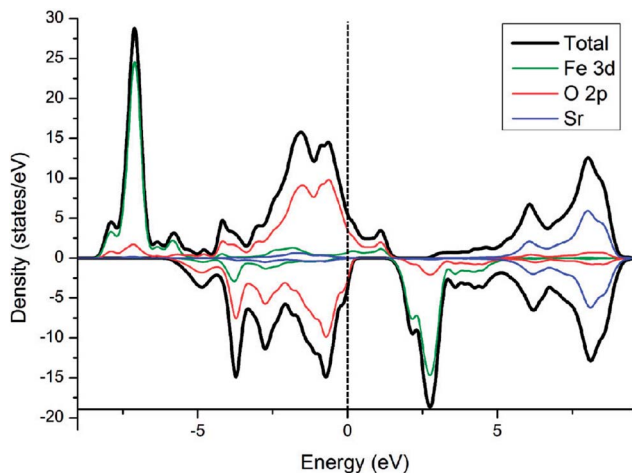


Fig. 3 Electronic density of states of $\text{Sr}_3\text{Fe}_2\text{O}_7$.

Oxygen vacancy formation energy (E_{vac}) and proton formation energy (E_{hydrd}) on SrO-1 surface (at O1 site, as indicated in Fig. 2(a)) are calculated as 0.77 and -2.26 eV, respectively. The extremely negative value of E_{hydrd} on SrO-1 terminal surface indicates that protons tend to automatically form in wet atmosphere with no need of additional energy input. E_{vac} and E_{hydrd} at SrO-2 and FeO-2 terminal surfaces (with relatively low surface energies) are also calculated. As summarized in Table 1, within the investigated terminal surfaces, SrO-1 terminal surface has the lowest E_{hydrd} value, while SrO-2 has the lowest E_{vac} value. These results are consistent with our previous investigations on E_{vac} and E_{hydrd} calculated at different O positions (as shown in Fig. 2(a)) in SFO bulk,¹² suggesting that protons are more easily to locate at rock-salt layer while oxygen vacancies at the central of perovskite layers. Moreover, compared with those in $\text{Sr}_3\text{Fe}_2\text{O}_7$ bulk model,^{12,16} values of E_{vac} and E_{hydrd} are much lower for surfaces, suggesting that oxygen vacancies and protons are more easily to form at surface than in bulk.

As comparison, (001) terminal surface of $\text{La}_{0.5}\text{Sr}_{0.5}\text{Co}_{0.25}\text{Fe}_{0.75}\text{O}_3$ (LSCF) are also built, as shown in Fig. S2.† Calculation parameters of LSCF surface are set based on literature.³¹ It should be noticed that in some recent studies, SrO terminal surfaces were applied for LSCF on account of Sr^{2+} segregation.^{32,33} Yet, such Sr segregation were usually experimentally observed after heating at intermediate temperatures (600–800 °C) for at least tens of hours and could be aggravated at the presence of Cr_2O_3 or electrochemical forces.^{32–36} While on fresh LSCF sintered at 1350–1400 °C for 4–5 hours, no Sr segregation

were spotted.^{32–34} Moreover, XPS analysis on the surface composition of pretested LSCF cathode indicated that its Sr/(La + Sr) ratios (~ 0.43) were reasonably close to the desired values (0.4).³⁶ In this work, emphasis is put on investigating the native catalytic ability of SFO cathode toward oxygen reduction reaction in O-SOFC and in P-SOFC with LSCF as a comparison. And therefore, LSCF computational model is built based on its original composition without considering the Sr segregation as some studies suggested.^{32,33} Moreover, to get good compare with SrO-1 terminal surface (most stable surface) in $\text{Sr}_3\text{Fe}_2\text{O}_7$, LaSrO (001) terminal surface of LSCF is adopted for catalytic activity investigation.^{37,38} LSCF (001) surface has a surface energy of 1.15 eV, higher than those of SFO (001) surface, as shown in Table 1. Importantly, E_{vac} and E_{hydrd} on this surface are 2.66 and 3.59 eV, respectively, much higher than those of SFO, indicating that these defects are harder to generate in LSCF than in SFO. The large E_{vac} and E_{hydrd} values in LSCF may root in its high La atom content in surface²² and its perovskite structure. Moreover, the extremely high value of E_{hydrd} on LSCF surface suggests that it is impossible to form proton defects even at high temperatures. And therefore, in P-SOFCs, LSCF has to work as composite cathodes to effectively accelerate cathode reactions.

O_2 reduction on the SrO-1 terminal surface in O-SOFCs

Based on above results, we can find that SrO-1 terminal surface is the most stable surface, and can energetically benefit the formations of oxygen vacancies and protons defect. And therefore, SrO-1 terminal surface is chosen for our later investigations on oxygen reduction reactions in both O-SOFCs and P-SOFCs.

In O-SOFCs, cathode-reaction processes begin with oxygen adsorption and dissociation on the surface, as illustrated in Fig. 1(a). On defect SrO-1 terminal surface which contains one oxygen vacancy (as shown in Fig. 2(b)), there are three possible active sites for O_2 adsorption, as shown in Fig. 4, which can be indicated as Sr-top1, Sr-top2 and $\text{O}_{\text{vac-top}}$, respectively. The adsorption energies, O–O bond lengths and atomic charges of adsorbed oxygen species are calculated and summarized in Table 2. As shown in Table 2, values of O_2 adsorption energies on Sr-top1, Sr-top2 and $\text{O}_{\text{vac-top}}$ are about -0.41 , -0.95 and -1.21 eV, respectively. These negative values indicate that adsorption of O_2 molecules on the $\text{Sr}_3\text{Fe}_2\text{O}_7$ (001) surface is an exothermal reaction. Especially, adsorption energy at $\text{O}_{\text{vac-top}}$ is much larger than those at Sr sites, suggesting that surface oxygen defect is energetically more favorable for O_2 adsorption than Sr ions. Adsorption energies of O_2 over perfect SrO-1 terminal surface are also conducted as

Table 1 Surface energies (E_{sur}), oxygen vacancy formation energies (E_{vac}) and proton formation energies (E_{hydrd}) of SrO-1 (O1), SrO-2 (O3) and FeO-2 (O4) terminal surface in SFO (as indicated in Fig. 2a). E_{vac} and E_{hydrd} at O1, O3, and O4 sites in SFO bulk are also given for comparison.^{12,16} The suffix –surf and –bulk denotes the values acquired on the terminal surface model and the corresponding bulk model, respectively. E_{vac} and E_{hydrd} at LSCF (001) surface are also calculated as comparison

	E_{sur} (eV)	$E_{\text{vac-surf}}$ (eV)	$E_{\text{vac-bulk}}$ (eV)	$E_{\text{hydrd-surf}}$ (eV)	$E_{\text{hydrd-bulk}}$ (eV)
SrO-1/O1	0.48	0.77	1.57	-2.26	-1.44
SrO-2/O3	0.81	0.47	0.53	-1.52	-0.23
FeO-2/O4	0.87	1.00	1.10	-0.65	-0.34
LSCF(001)	1.15	2.66	—	3.59	—



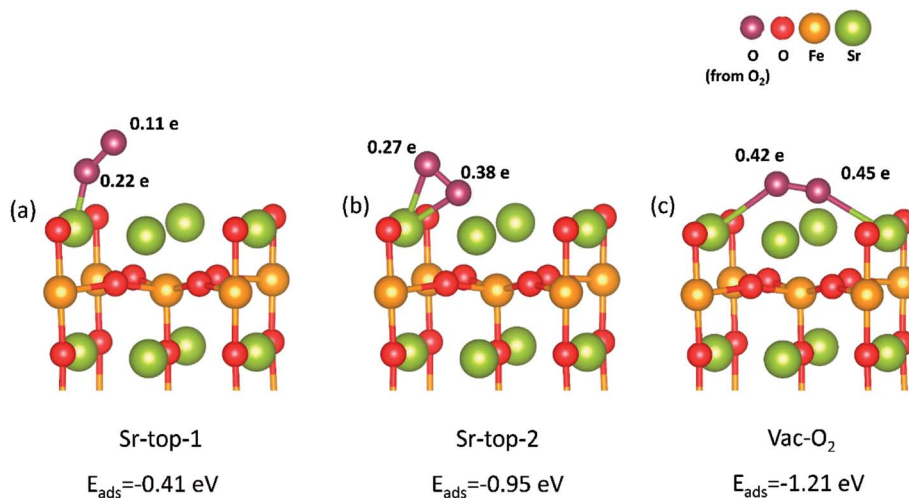


Fig. 4 Optimized O_2 adsorption structures on the SrO-1 (001) terminal surface with their respective adsorption energy. Values close to each oxygen atoms indicate the atomic charge of the adsorbed oxygen species obtained from surface.

shown in Fig. S3,[†] of which the largest adsorption energy is only -0.47 eV. The low adsorption energy on perfect surface further suggest that existence of surface oxygen vacancy is very important for O_2 adsorption.

Bader charges⁴⁰ of the adsorbed O_2 species are summarized in Table 2, which could provide qualitative analysis on their bonding strength to the surface. Different electrons are obtained for the adsorbed O_2 species which depend largely on their adsorption sites and their relative positions. Largest charge ($0.87 e$) of oxygen atom is observed when O atoms bonding to Sr atom over the $O_{vac-top}$ site (Fig. 4(c)). This is consist with the above adsorption energy results, and indicates further the great promotion effect of oxygen vacancy on O_2 adsorption reaction. When the oxygen adsorbs on LSCF (001) surface, the calculated charge of oxygen atom is $1.46 e$. Since these charges go into the antibonding orbital of the oxygen molecule, the O-O is easy to break in this case. This means that it's easier for oxygen to dissociate on LSCF than on SFO surface.

Minimum energy pathway for O_2 dissociation on the defect SFO (001) surface are then explored based on the Climbing

Image Nudged Elastic Band (CI-NEB). As shown in Fig. 5(a), O_2 adsorbs automatically over oxygen vacancy on SrO-1 surface, releasing 1.21 eV heat; then, the adsorbed O_2 dissociates with one oxygen atom incorporating the neighboring oxygen vacancy, and in this process an energy barrier of 2.28 eV needs to overcome; and finally, the oxygen atom bonding above Sr sites migrate to a more stable site without overcoming any energy barrier. In the whole process, O_2 dissociation should be the rate-determining step because of its large energy barrier.

Notably, when O_2 adsorbs over similar LSCF (001) terminal surface, no energy barrier need to be overcome within the whole reactions (as shown in Fig. 5(b)), indicating that O_2 dissociation over LSCF surface is very fast and should not be the rate-limiting steps. Previous studies on defect LSCF surface indicates that La and Sr ions have similar activity towards O_2 adsorption and dissociation, which are much lower than those of Fe and Co ions.⁴¹ Since no active ions such Fe or Co on both LSCF (001) surface and SrO-1 terminal surface, such different energy barrier for O_2 dissociation in these two surfaces may be related to their crystal structures, surface energies and defects formation energies. It is

Table 2 The calculated adsorption energies (E_{ads}), O–O bond lengths (r_{O-O}), and atomic charges of adsorbed oxygen. There are two possible configurations for oxygen species (superoxide and peroxide), of which the O–O bond is vertical or parallel to the surface, respectively.^{13,39} The x-H means the numbers of protons on the specific surface

Species	E_{ads} (eV)	r_{O-O} (Å)	Atomic charge (e)			Assignment	
			O1	O2	O_{sum}		
With oxygen vacancy	SFO-0H	Sr-top-1	-0.41	0.22	0.11	0.33	Superoxide
		Sr-top-2	-0.95	0.27	0.38	0.65	Superoxide
		Vac- O_2	-1.21	0.45	0.42	0.87	Peroxide
	SFO-2H	Sr-top	-0.92	0.38	0.30	0.68	Superoxide
		Vac- O_2	-1.29	0.40	0.51	0.91	Peroxide
		SFO-4H	Sr-top	-0.99	0.41	0.35	0.76
	LSCF	Vac- O_2	-1.39	0.41	0.52	0.93	Peroxide
		Sr-top	-1.52	0.72	0.74	1.46	Peroxide
		Vac- O_2	-1.52	0.72	0.74	1.46	Peroxide
Without oxygen vacancy	SFO-0H	Sr-top-1	-0.22	0.06	0.21	0.27	Superoxide
		Sr-top-2	-0.47	0.13	0.18	0.31	Superoxide
	SFO-2H	Sr-top-1	-0.37	0.15	0.14	0.29	Superoxide
		Sr-top-2	-0.54	0.36	0.19	0.55	Superoxide



suggested that to improve catalytic activity of R-P cathodes towards O_2 dissociation will be very effective to accelerate O_2 reduction process in O-SOFCs.

It should be noted that after O_2 dissociation, the formed oxygen ions need to migrate through surface or bulk of cathode to electrolyte to release the oxygen vacancy and to fulfill the

cathode reactions. Surface effects on the diffusions of oxygen ions (in opposite diffusion direction with oxygen vacancies) are investigated, as shown in Fig. S4† ([001] direction) and Fig. S5† ([100] direction). Data in Table S1† clearly indicates that SrO-1 surface has a possible effect on the diffusion of oxygen ions, which largely lowers the energy barrier for O ions from 1.40 eV

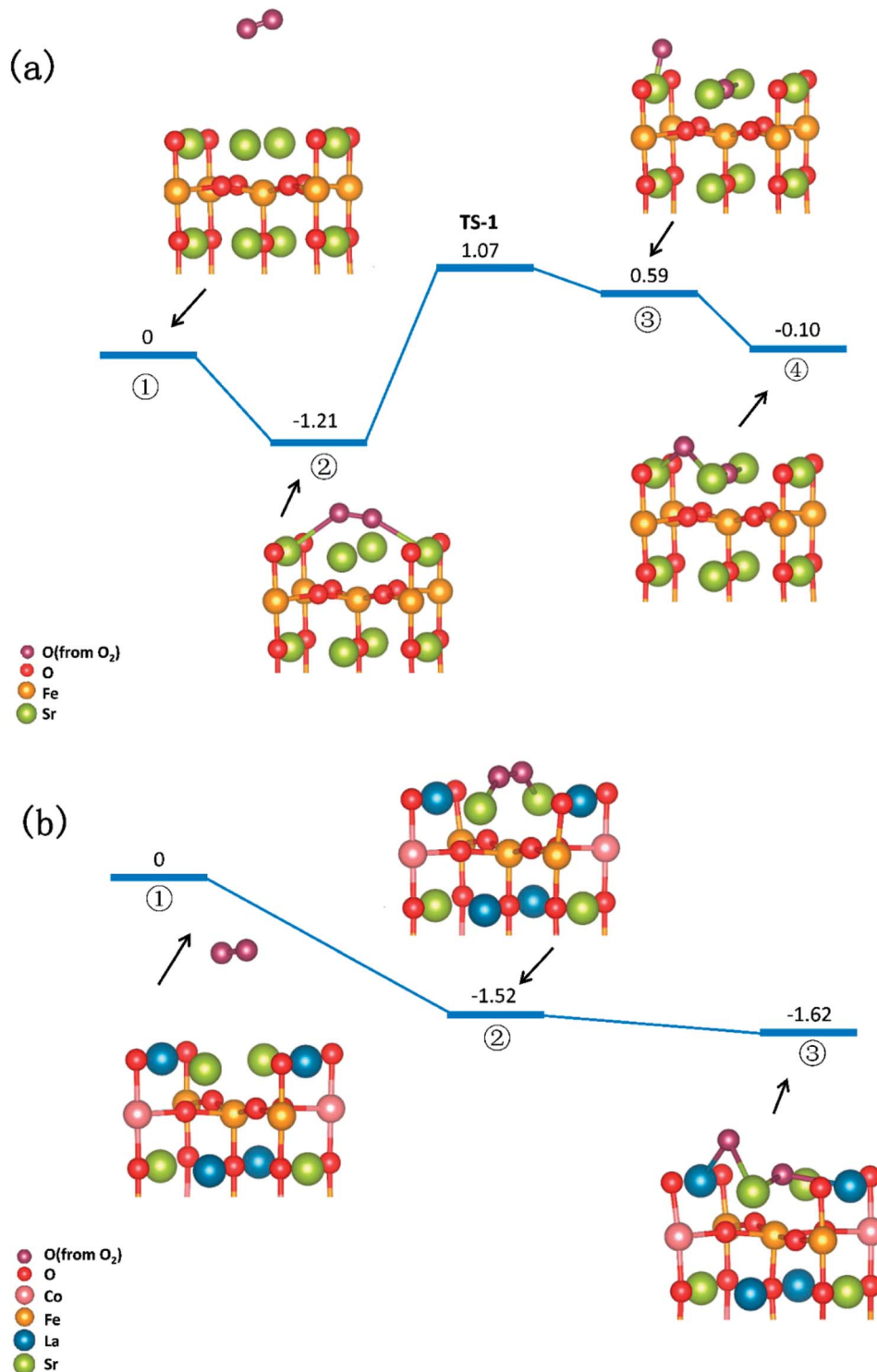


Fig. 5 Potential energy diagram for oxygen reduction on (a) SrO-1 terminal surface of $Sr_3Fe_2O_7$ cathode and (b) LSCF (001) terminal surface in O-SOFCs.



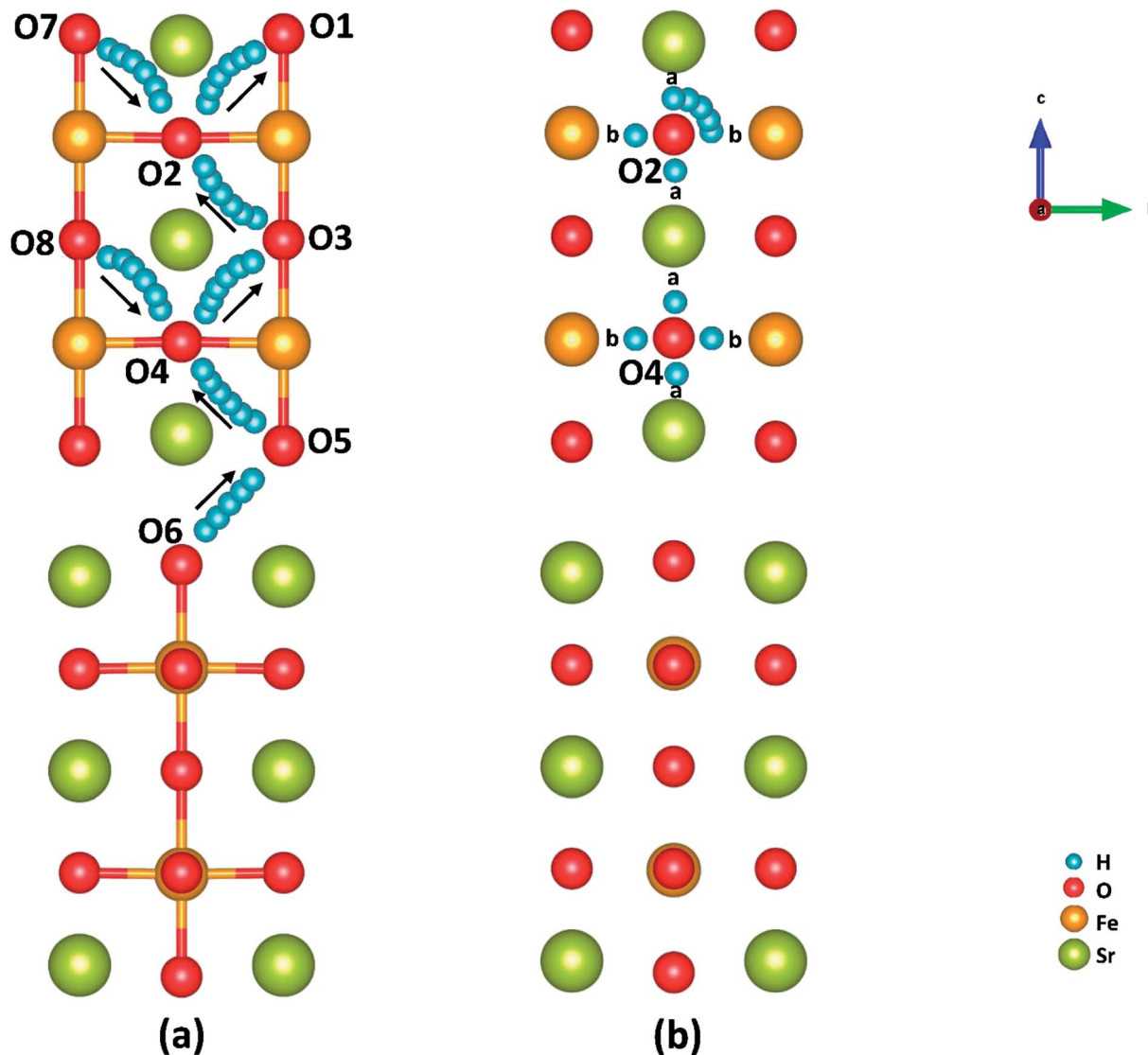


Fig. 6 Illustration of (a) proton intraoctahedral hopping pathways and (b) the proton reorientation around O.

to 1.04 eV. Moreover, an energy barrier as low as 0.35 eV is observed for oxygen diffusion within SrO-1 terminal surface, suggesting that oxygen ions are anisotropic in SFO. Notably, energy barriers for oxygen diffusion in LSCF is about 1.96 eV, higher than those in SFO. The relatively lower oxygen vacancies formation energy and lower energy barrier for oxygen diffusion makes SFO have a better oxygen ion conduction than LSCF, which is in good agreement with experimental investigations.¹⁶

In conclusion, compared with perovskite oxides, SFO cathode has low oxygen vacancies formation energies and low ion diffusion energy barriers, which can bring forth excellent oxygen ion conduction. Nevertheless, its high energy barrier for O₂ dissociation may restrict its application as an excellent cathode in O-SOFCs. To improve the catalytic activity of SFO *via* structure or composition modifications will be the key. Also, the different conducting properties and catalytic activities of SFO and LSCF clearly indicate their different rate-limiting steps in cathode reactions in O-SOFCs.¹⁴

Cathode reaction in P-SOFCs

In P-SOFCs, cathode reactions may include the following steps: (1) protons generated at anode transfer to the surface of SFO cathode *via* diffusion; (2) O₂ molecules adsorb on the SFO cathode surface, and then dissociate with the presence of protons; (3) protons react with the dissociated O₂ to form water molecules; and finally, (4) the formed water molecules are released from the cathode surface to the gas atmosphere. As shown in eqn (2), four protons are needed to react completely with one oxygen molecule. To theoretically investigate the cathode reaction paths, two surfaces containing two or four protons are used, respectively, as shown in Fig. 8 and 9. For the former, the other two protons needed in reactions are located at the second layer of the surface. And to distinguish the amount of surface protons, SrO-1-xH, where the *x* indicates the number of protons on the surface.



Table 3 Energy barriers for proton intraoctahedral hopping in SrO-1 surface model and those acquired in the similar hopping process in SFO bulk model.¹² Energy barriers for proton intraoctahedral hopping in BZCO bulk¹⁵ is also listed

Octahedron transfer path	Energy barriers/eV		
	Surface	Bulk	BaZr _{0.75} Co _{0.25} O ₃
O6 to O5	0.67	0.62	0.03–0.63
O5 to O4	0.34	0.53	
O4 to O3	0.04	0.05	
O3 to O2	0.31	0.25	
O2 to O1	0.14	0	
O7 to O2	0.62	0.53	
O8 to O4	0.28	0.25	

Table 4 Energy barriers for proton reorientation in SrO-1 surface model and those acquired in the similar reorientation process in SFO bulk model.¹² Energy barriers for proton reorientation in BZCO bulk¹⁵ is also listed

Reorientation path	Energy barriers/eV		
	Surface	Bulk	BaZr _{0.75} Co _{0.25} O ₃
O2(a) to O2(b)	0.44	0.32	0.05–0.26
O2(b) to O2(a)	0.07	0.25	
O4(a) to O4(b)	0.24	0.27	
O4(b) to O4(a)	0.12	0.08	

The proton conduction in the Sr₃Fe₂O₇ cathode

Cathode reaction in P-SOFCs starts from the proton diffusion, which are also suggested as one of the rate-limiting steps in composite cathodes.¹² Proton diffuse to SFO-1 surface *via* the combination of two typical paths: the proton intra octahedral hopping and the proton reorientation around O ions,⁴² as shown in Fig. 6(a) and (b), respectively. Energy barriers for the seven possible unique intraoctahedral hopping and four proton reorientations are summarized in Tables 3 and 4. Protons

transferring across the rock-salt layer (O6 to O5) is most difficult process, which needs to overcome the highest energy barrier of 0.67 eV. This value is in the same range for proton conduction in proton conducting electrolytes, such as Y-doped BaZrO₃ (0.08–1.35 eV)⁴³ and BaZr_{0.75}Co_{0.25}O₃ (0.01–0.63 eV).¹⁵ The low energy barrier for proton migration and the low proton formation energy strongly suggest the SFO system is good proton conduction containing cathode material.

As shown in Fig. 6 and Table 3, the energy barriers for protons hopping along the [010] direction (O8 → O4 → O3) are far smaller than those for [001] direction (O6 → O5 → O4 → O3 → O2 → O1),

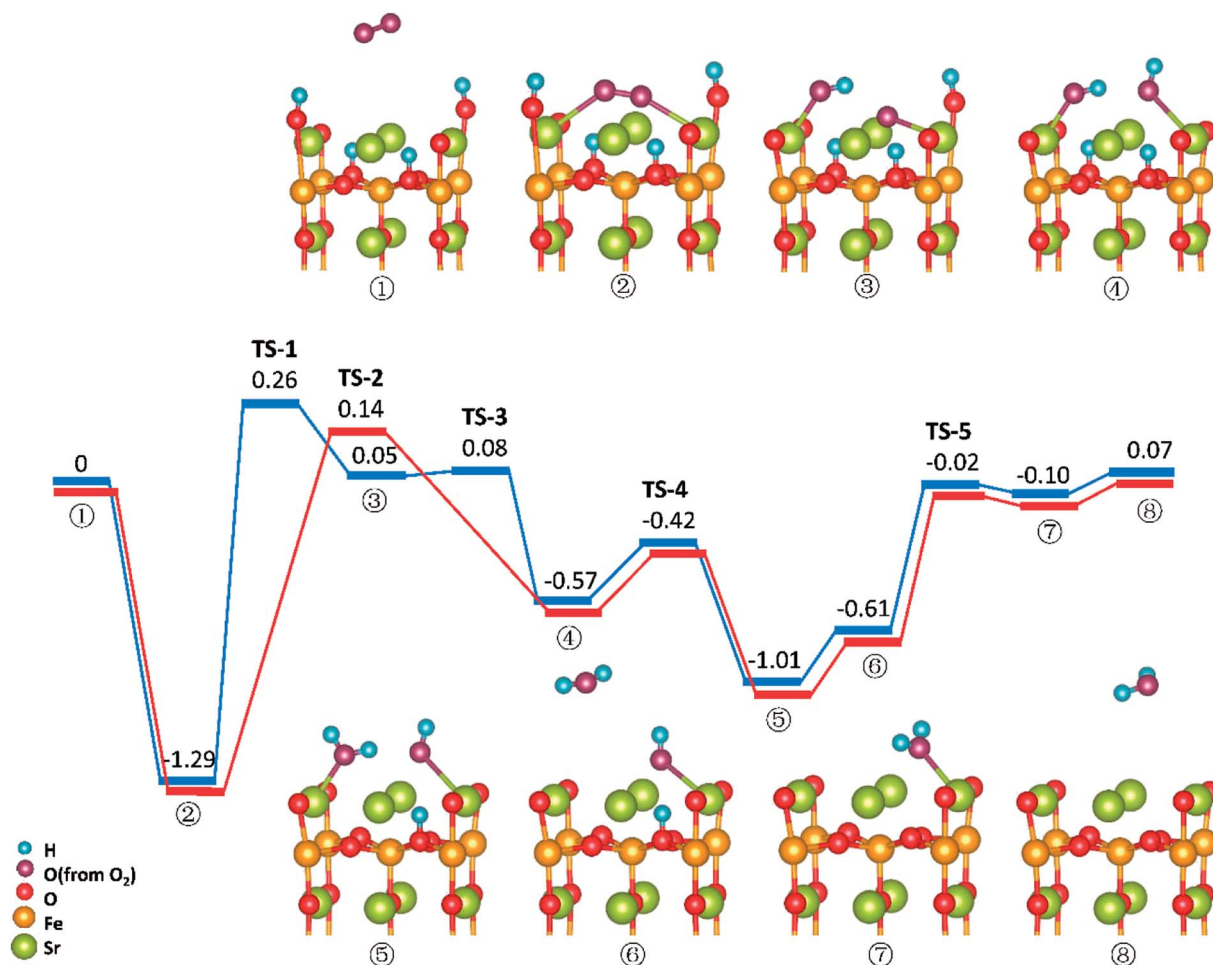


Fig. 7 Potential energy diagram for reactions on the SrO-1 surface with two protons present.



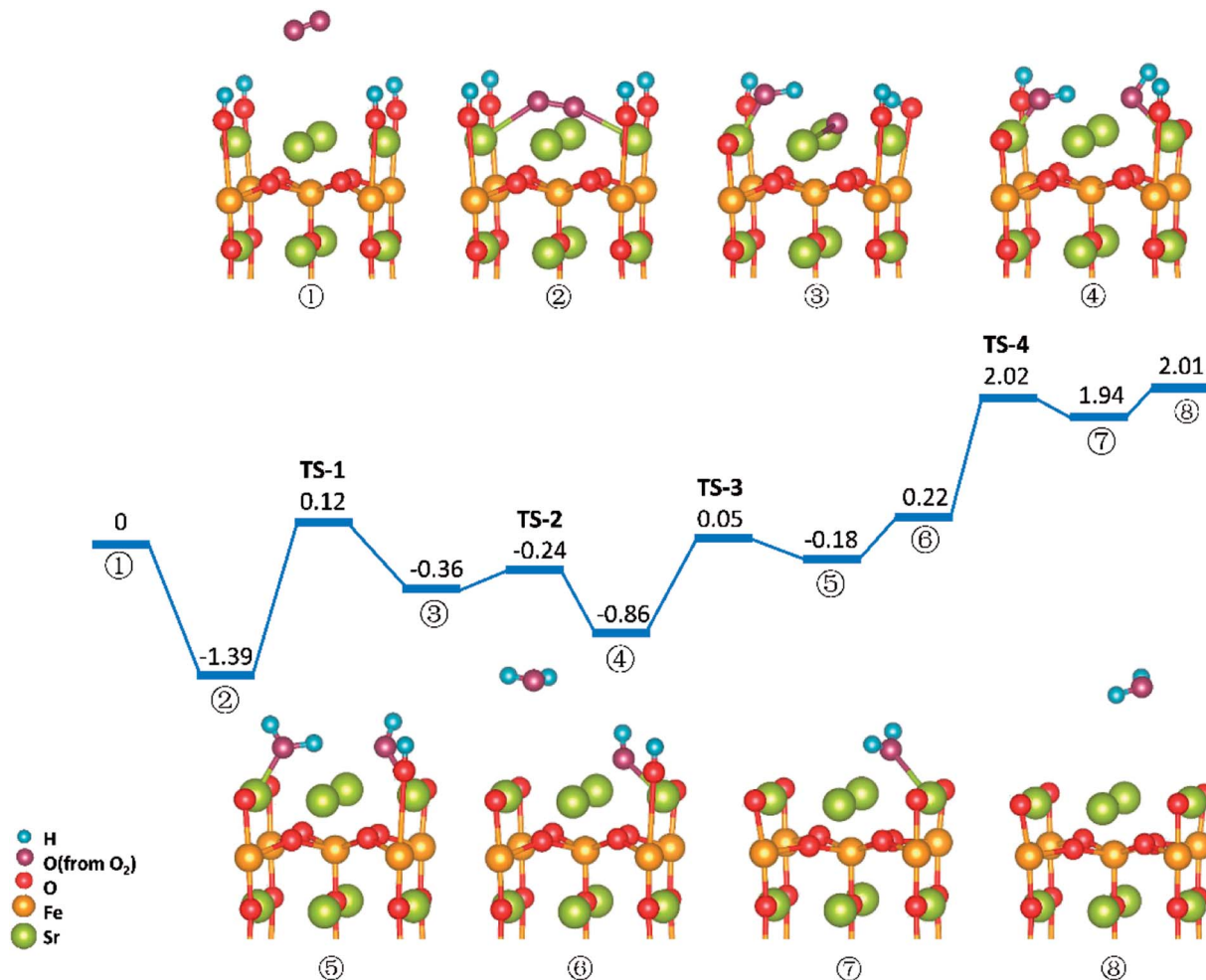


Fig. 8 Potential energy diagram for reactions on SrO-1-4H surface.

indicating that proton transferring in surface mode is also anisotropic as that in bulk.¹⁵ However, the energy barriers for proton hopping in surface model are similar to those obtained in bulk model, indicating that the surface effect on proton transferring is not as obvious as that on oxygen diffusion (as shown in Tables S2 and S3†). Moreover, when the proton is hopping away from an oxygen ion locating at Sr-O layer, the energy barrier is relatively high, while relatively low in the opposite process. It's found that the effective charge of H atoms ($-0.34 e$ in SrO layer and $-0.22 e$ in FeO layer) and the bond length of O-H (0.96 \AA in SrO layer and 1.08 \AA in Fe-O layer) differ with the positions of proton defects. These suggest a relatively strong bonding of proton defects with O in Sr-O layer, which attributes to the trapping effect⁴⁴⁻⁴⁶ of the SrO layer on proton transferring as other studies.¹² This property makes protons hop to outer surface with a small energy barrier (0.14 eV), which provides support for our SrO-1-2H model.

Reactions on the SrO-1 terminal surface with two surface proton present (SrO-1-2H)

Similar to those in O-SOFCs, O_2 adsorbing over the oxygen vacancy is the most stable configuration of O_2 on SrO-1-2H surface with one oxygen vacancy, as shown in Fig. 7-② and

Fig. S6.† The adsorption energy of such configuration is -1.29 eV , larger than that without presence of protons (-1.21 eV in Fig. 4). Meanwhile, the O_2 adsorption energies on the SrO-1-2H surface with no oxygen vacancy are simulated. As shown in Table .2, the largest O_2 adsorption energy is -0.54 eV , also larger than that on perfect SrO-1 surface (-0.47 eV) with no protons present. These results clearly suggest that presence of protons on SrO-1 surface could promote the adsorption of oxygen molecule.

In regards to O_2 dissociation and water formation process, two potential paths are mapped on SrO-1-2H surface denoted by red and blue line, respectively, as shown in Fig. 7. In path-1 (blue line), detailed reaction processes can be described as: (1) the first proton (left side in ②) migrates to the adsorbed O_2 molecule and activate the dissociation of O_2 molecule to form one hydroxyl (③); (2) the second proton (right side in ③) attaches to the right oxygen ion forming a new hydroxyl incorporating in the former address of oxygen vacancy (④) with a negligible energy barrier of 0.03 eV (TS-3) and releases 0.62 eV heat; 3) the proton locating at the second layer transfer to the surface and then react with the first hydroxyl to form an adsorbed water molecule (⑤) which then desorbs from the surface to gas atmosphere (⑥); 4) and finally, the second water molecular forms *via* the reaction of the



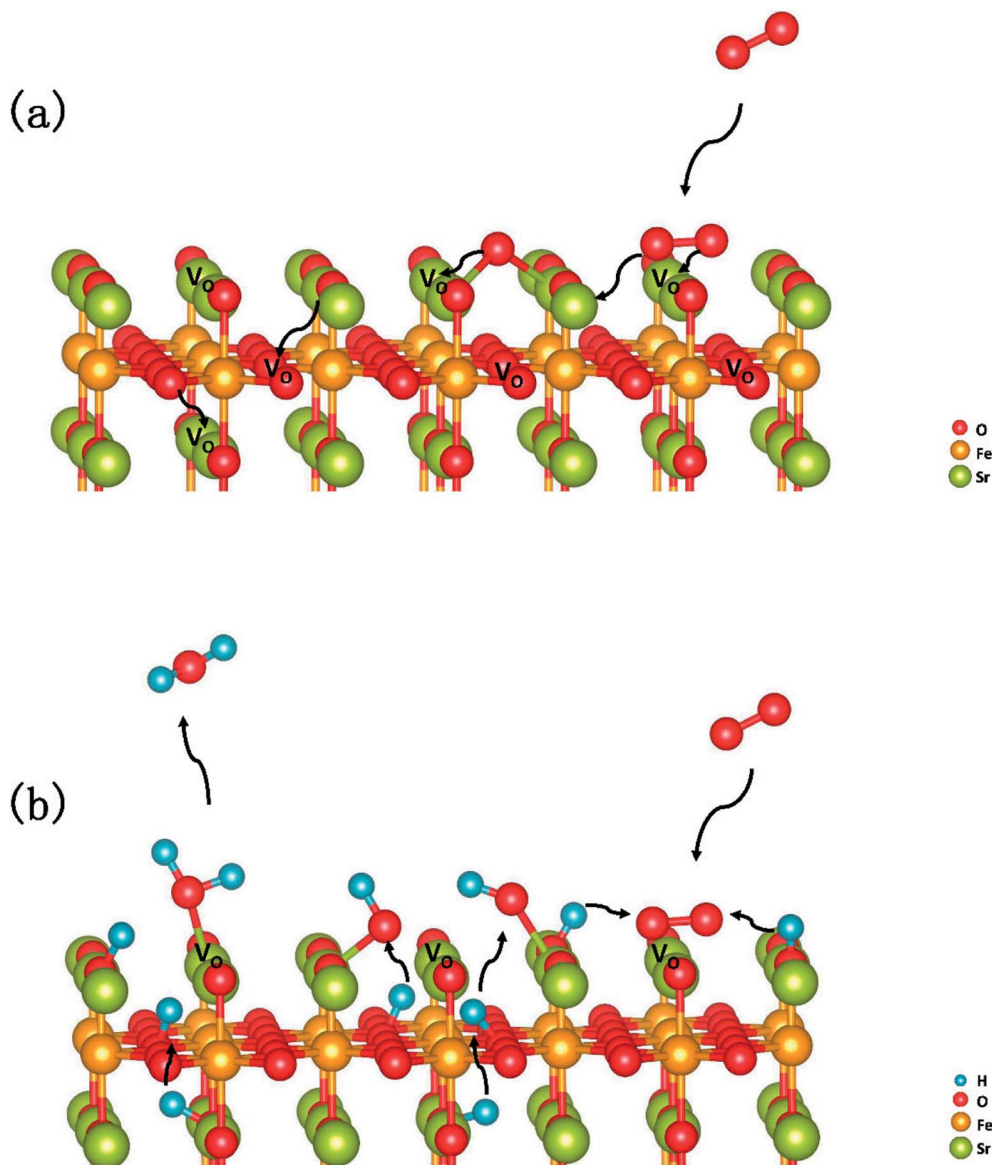


Fig. 9 Schematic diagram of the most feasible reaction model over the $\text{Sr}_3\text{Fe}_2\text{O}_{7-\delta}$ cathode for oxygen conducting SOFCs (a) and proton conducting SOFCs (b).

last proton with another hydroxyl (OH), and is released to the gas (H_2O). Throughout the whole reaction process, O_2 dissociation is still the one needing to overcome the largest energy barrier of 1.55 eV (TS-1). Fortunately, compared to those in O-SOFCs (energy barrier of 2.28 eV), the energy barrier here is much depressed, indicating that the hydrogenated surface can obviously accelerate the O_2 reduction processes.

The main divergence between path-1 and path-2 lies in the way for protons on surface to react with the O_2 . In path-2 (red line), the cathode reaction begins with two protons simultaneously attaching to the oxygen molecule to dissociate O_2 , and thus, generating two hydroxyls concurrently. In this process, an energy barrier of 1.43 eV need to be overcome, slightly lower than that in path-1. This seems to suggest that two protons react with the oxygen molecule at the same time is more effective than that react one by one.

It is worth mentioning that direct formation of the first water *via* the reaction of adsorbed oxygen ion and the two surface protons is not observed in our DFT calculation process. This may root in its unstable structure in energy because of the contest of the two dissociated O atoms for the protons.

Reactions on the SrO-1 terminal surface with four surface proton defects present (SrO-1-4H)

O_2 reduction reactions on SrO-1-4H surface are shown in Fig. 8. In this model, all four protons required in oxygen reduction reaction (as shown in eqn (2)) locate at SrO-1 terminal surface. And reaction processes in this path (path-3) occur in sequence as: (1) O_2 adsorbs on the $\text{O}_{\text{vac-top}}$ site (O_2); (2) one proton diffuses to the adsorbed O_2 molecule and forms a hydroxyl (OH) by surmounting an energy barrier of 1.51 eV, similar to that in



path 1; (3) the second proton (on the opposite site of the first hydroxyl) transfers to the dissociated oxygen ion to form another hydroxyl (④); (4) the first water molecule forms *via* the reaction of the third proton with the first hydroxyl (⑤), which is then released to gas (⑥); (5) the second water molecule forms with a large energy barrier of 1.80 eV (TS-4) to be overcome, and then desorbs to gas phase.

In this path, the dissociation of O₂ and the formation of second water molecule, need to overcome large energy barriers. Moreover, compared with the energy map for SrO-1-2H surface (Fig. 7), the formation energy barriers for the water molecule, especially the second one, are extremely high on SrO-1-4H surface, suffering from the competition of neighboring surface oxygen ions for protons. These seem to suggest that existence of too many protons at surface is unfavorable to the formation of water. This along with that presence of oxygen vacancy favoring for O₂ adsorption implies that steam pressure in testing atmosphere should not be too much to achieve high performance, and that coexistence of suitable concentrations of protons, oxygen vacancies and electron defects (triple conducting) should be of key importance for cathode reactions in P-SOFC.

Comparing the SFO performance in O-SOFC and P-SOFC, our DFT calculation also reveals that SFO shows a large energy barrier (~2.28 eV) for oxygen dissociation reaction in O-SOFC, while in P-SOFCs, such energy barrier has been greatly depressed at the presence of protons (1.43–1.57 eV). This is in good agreement with experimental observation that SFO is more effective than both LSCF and LSCF-BZCY as a cathode for P-SOFCs; while for O-SOFCs, opposite is the case.^{12,16}

Moreover, DFT investigations suggest that the high proton formation energy of LSCF makes it unfavorable for proton formation, and therefore, composite cathodes consisting LSCF and proton conductors (for example, BZCY) have to be made up to effectively accelerate cathode reactions.

Conclusions

In summary, a comprehensive DFT calculation was performed, which studied the O₂ reduction on the SFO (001) surface working in both O-SOFCs and P-SOFCs. Most feasible paths for cathode reaction over the SrO-1 surface are summarized in Fig. 9(a) for O-SOFCs and Fig. 9(b) for P-SOFCs.

In O-SOFCs, adsorption of O₂ on the SFO surface goes smoothly, while O₂ dissociation occurs with a high energy barrier (2.19 eV) that needs to be overcome, which greatly limits the cathode reaction rates of SFO. And therefore, to improve cathode performance of SFO, it is key to improve its catalytic activity toward O₂ dissociation. In P-SOFCs, existence of protons benefits the dissociation of O₂, which effectively lowers the energy barriers to about 1.43 eV. Yet, when four protons needed in cathode reactions all locate at the SFO surface, the energy barrier for steam formation increased largely from 0.59 eV to 1.80 eV. This result seems to indicate that overranging steam concentrations in the testing atmosphere may have a negative effect on cell performance for the reduction of oxygen vacancies and the high energy barrier for water formation.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was financially supported by the National Key Research and Development Program of China (2017YFA0402800), the Natural Science Foundation of China (51472228, 51627901), the External Cooperation Program of BIC, Chinese Academy of Sciences (211134KYSB20130017), Hefei Science Center CAS (2016HSC-IU004), and the Fundamental Research Funds for the Central Universities (WK3430000004).

Notes and references

- 1 S. Choi, S. Yoo, J. Kim, S. Park, A. Jun, S. Sengodan, J. Kim, J. Shin, Y. J. Hu and Y. M. Choi, Highly efficient and robust cathode materials for low-temperature solid oxide fuel cells: PrBa_{0.5}Sr_{0.5}Co_{2-x}Fe_xO_{5+δ}, *Sci. Rep.*, 2013, **3**, 2426.
- 2 T. Wei, Y. H. Huang, R. Zeng, L. X. Yuan, X. L. Hu, W. X. Zhang, L. Jiang, J. Y. Yang and Z. L. Zhang, Evaluation of Ca₃Co₂O₆ as cathode material for high-performance solid-oxide fuel cell, *Sci. Rep.*, 2013, **3**, 1125.
- 3 T. Tsai and S. A. Barnett, Effect of LSM-YSZ cathode on thin-electrolyte solid oxide fuel cell performance, *Solid State Ionics*, 1997, **93**(3–4), 207–217.
- 4 J. Harris, C. Metcalfe, M. Marr, J. Kuhn and O. Kesler, Fabrication and characterization of solid oxide fuel cell cathodes made from nano-structured LSCF-SDC composite feedstock, *J. Power Sources*, 2013, **239**(239), 234–243.
- 5 B. Wei, L. Zhe, X. Huang, S. Li, G. Ai, Z. Liu and W. Su, Electrochemical characteristics of Ba_{0.5}Sr_{0.5}Co_{0.8}Fe_{0.2}O_{3-δ}-Sm_{0.2}Ce_{0.8}O_{1.9} composite materials for low-temperature solid oxide fuel cell cathodes, *Mater. Lett.*, 2006, **60**(29), 3642–3646.
- 6 L. Yang, Z. Liu, S. Wang, Y. M. Choi, C. Zuo and M. Liu, Mixed Proton-Oxide Ion-Electron Conducting Cathode for SOFCs Based on Oxide Proton Conductors, *J. Power Sources*, 2010, **195**(2), 471–474.
- 7 L. Yang, C. Zuo, S. Wang, Z. Cheng and M. Liu, A Novel Composite Cathode for Low-Temperature SOFCs Based on Oxide Proton Conductors, *Adv. Mater.*, 2010, **20**(17), 3280–3283.
- 8 C. Duan, J. Tong, M. Shang, S. Nikodemski, M. Sanders, S. Ricote, A. Almansoori and R. O'Hayre, Readily processed protonic ceramic fuel cells with high performance at low temperatures, *Science*, 2015, **349**(6254), 1321–1326.
- 9 M. Shang, J. Tong and R. O'Hayre, A promising cathode for intermediate temperature protonic ceramic fuel cells: BaCo_{0.4}Fe_{0.4}Zr_{0.2}O_{3-δ}, *RSC Adv.*, 2013, **3**(36), 15769–15775.
- 10 Z. Tao, L. Bi, L. Yan, W. Sun, Z. Zhu, R. Peng and W. Liu, A novel single phase cathode material for a proton-conducting SOFC, *Electrochem. Commun.*, 2009, **11**(3), 688–690.
- 11 Y. Rao, S. Zhong, H. Fei, Z. Wang, R. Peng and Y. Lu, Cobalt-doped BaZrO₃: A single phase air electrode material for



- reversible solid oxide cells, *Int. J. Hydrogen Energy*, 2012, **37**(17), 12522–12527.
- 12 Z. Wang, W. Yang, S. P. Shafi, L. Bi, Z. Wang, R. Peng, C. Xia, W. Liu and Y. Lu, A high performance cathode for proton conducting solid oxide fuel cells, *J. Mater. Chem. A*, 2015, **3**(16), 8405–8412.
- 13 Y. Choi, M. C. Lin and M. Liu, Computational study on the catalytic mechanism of oxygen reduction on $\text{La}_{(0.5)}\text{Sr}_{(0.5)}\text{MnO}_{(3)}$ in solid oxide fuel cells, *ChemInform*, 2007, **46**(38), 7214–7219.
- 14 Y. L. Lee, J. Kleis, J. Rossmesl, Y. Shao Horn and D. Morgan, Prediction of solid oxide fuel cell cathode activity with first-principles descriptors, *Energy Environ. Sci.*, 2011, **4**(10), 3966–3970.
- 15 Z. Wang, W. Yang, Z. Zhu, R. Peng, X. Wu, C. Xia and Y. Lu, First-principles study of O_2 reduction on $\text{BaZr}_{1-x}\text{Co}_x\text{O}_3$ cathodes in protonic-solid oxide fuel cells, *J. Mater. Chem. A*, 2014, **2**(39), 16707–16714.
- 16 D. Huan, Z. Wang, Z. Wang, R. Peng, C. Xia and Y. Lu, High-Performance Cathode with a Two-Layered R-P Structure for Intermediate Temperature Solid Oxide Fuel Cells, *ACS Appl. Mater. Interfaces*, 2016, **8**(7), 4592–4599.
- 17 B. Pe, Projector augmented-wave method, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1994, **50**(24), 17953–17979.
- 18 B. Morgan, D. Scanlon and G. Watson, Small polarons in Nb- and Ta-doped rutile and anatase TiO_2 , *J. Mater. Chem.*, 2009, **19**(29), 5175–5178.
- 19 G. Kresse and J. Furthmüller, Efficiency of *ab initio* total energy calculations for metals and semiconductors using a plane-wave basis set, *Comput. Mater. Sci.*, 1996, **6**(1), 15–50.
- 20 J. P. Perdew, K. Burke and M. Ernzerhof, Generalized Gradient Approximation Made Simple, *Phys. Rev. Lett.*, 1996, **77**(18), 3865–3868.
- 21 Y. A. Mastrikov, M. M. Kuklja, E. A. Kotomin and J. Maier, First-principles modelling of complex perovskite $(\text{Ba}_{1-x}\text{Sr}_x)(\text{Co}_{1-y}\text{Fe}_y)\text{O}_{3-\delta}$ for solid oxide fuel cell and gas separation membrane applications, *Energy Environ. Sci.*, 2010, **3**(10), 1544–1550.
- 22 Y. A. Mastrikov, R. Merkle, E. A. Kotomin, M. M. Kuklja and J. Maier, Formation and migration of oxygen vacancies in La Sr Co Fe O perovskites: insight from calculations and comparison with Ba Sr Co Fe O, *Phys. Chem. Chem. Phys.*, 2012, **15**(3), 911–918.
- 23 T. Ota, H. Kizaki and Y. Morikawa, Mechanistic Analysis of Oxygen Vacancy Formation and Ionic Transport in $\text{Sr}_3\text{Fe}_2\text{O}_{7-\delta}$, *J. Phys. Chem. C*, 2018, **122**(8), 4172–4181.
- 24 S. L. Dudarev, G. A. Botton, S. Y. Savrasov, C. J. Humphreys and A. P. Sutton, Electron-energy-loss spectra and the structural stability of nickel oxide: An LSDA + U study, *Phys. Rev. B*, 1998, **57**(3), 1505–1509.
- 25 G. Sürücü and A. Erkişi, The investigation of electronic and thermo-elastic properties of lanthanum orthoferrite (LaFeO_3), *Perovskite*, 2017, **25**, 1–19.
- 26 A. A. Markov, M. V. Patrakeev, V. V. Kharton, Y. V. Pivak, I. A. Leonidov and V. L. Kozhevnikov, Oxygen Nonstoichiometry and Ionic Conductivity of $\text{Sr}_3\text{Fe}_{2-x}\text{Sc}_x\text{O}_{7-\delta}$, *ChemInform*, 2007, **19**(16), 3980–3987.
- 27 H. J. Monkhorst, Special points for Brillouin-zone integrations, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1976, **16**(4), 1748–1749.
- 28 G. Henkelman, B. P. Uberuaga and H. Jónsson, A climbing image nudged elastic band method for finding saddle points and minimum energy paths, *J. Chem. Phys.*, 2000, **113**(22), 9901–9904.
- 29 Y. Ling, T. Guo, X. Zhang, R. A. Budiman, Y. Fujimaki, T. Nakamura, B. Lin, T. Kawada and K. Amezawa, Evaluation of electrical conductivity and oxygen diffusivity of the typical Ruddlesden-Popper oxide $\text{Sr}_3\text{Fe}_2\text{O}_{7-\delta}$, *Ceram. Int.*, 2017, **43**(18), 16264–16269.
- 30 P. Adler, Electronic State, Magnetism, and Electrical Transport Behavior of $\text{Sr}_{3-x}\text{A}_x\text{Fe}_2\text{O}_7$ ($x \leq 0.4$, A = Ba, La), *J. Solid State Chem.*, 1997, **130**(1), 129–139.
- 31 A. M. Ritzmann, J. M. Dieterich and E. A. Carter, Density functional theory + U analysis of the electronic structure and defect chemistry of LSCF ($\text{La}_{0.5}\text{Sr}_{0.5}\text{Co}_{0.25}\text{Fe}_{0.75}\text{O}_{3-\delta}$), *Phys. Chem. Chem. Phys.*, 2016, **18**(17), 12260.
- 32 L. Zhao, J. Drennan, C. Kong, S. Amarasinghe and S. P. Jiang, Insight into surface segregation and chromium deposition on $\text{La}_{0.6}\text{Sr}_{0.4}\text{Co}_{0.2}\text{Fe}_{0.8}\text{O}_{3-\delta}$ cathodes of solid oxide fuel cells, *J. Mater. Chem. A*, 2014, **2**(29), 11114–11123.
- 33 J. Druce, H. Téllez, M. Burriel, M. D. Sharp, L. J. Fawcett, S. N. Cook, D. S. Mcphail, T. Ishihara, H. H. Brongersma and J. A. Kilner, Surface termination and subsurface restructuring of perovskite-based solid oxide electrode materials, *Energy Environ. Sci.*, 2014, **7**(11), 3593–3599.
- 34 D. Oh, D. Gostovic and E. D. Wachsman, Mechanism of $\text{La}_{0.6}\text{Sr}_{0.4}\text{Co}_{0.2}\text{Fe}_{0.8}\text{O}_3$ cathode degradation, *J. Mater. Res.*, 2012, **27**(15), 1992–1999.
- 35 M. E. Lynch, L. Yang, W. Qin, J. J. Choi, M. Liu, K. Blinn and M. Liu, Enhancement of $\text{La}_{0.6}\text{Sr}_{0.4}\text{Co}_{0.2}\text{Fe}_{0.8}\text{O}_{3-\delta}$ durability and surface electrocatalytic activity by $\text{La}_{0.85}\text{Sr}_{0.15}\text{MnO}_{3\pm\delta}$ investigated using a new test electrode platform, *Energy Environ. Sci.*, 2011, **4**(6), 2249–2258.
- 36 S. P. Simner, M. D. Anderson, M. H. Engelhard and J. W. Stevenson, Degradation mechanisms of La-Sr-Co-Fe- O_3 SOFC cathodes, *Electrochem. Solid-State Lett.*, 2006, **9**(10), A478–A481.
- 37 M. Wei, H. Li, G. Guo, Y. Liu and D. Zhang, Effects of PdO modification on the performance of $\text{La}_{0.6}\text{Sr}_{0.4}\text{Co}_{0.2}\text{Fe}_{0.8}\text{O}_{3-\delta}$ cathodes for solid oxide fuel cells: A first principle study, *Int. J. Hydrogen Energy*, 2017, **42**(36), 23180–23188.
- 38 A. Longo, L. F. Liotta, D. Banerjee, V. L. Parola, F. Puleo, C. Cavallari, C. J. Sahle, M. M. Sala and A. Martorana, The Effect of Ni Doping on the Performance and Electronic Structure of LSCF Cathodes Used for IT-SOFCs, *J. Phys. Chem. C*, 2018, **122**(2), 1003–1013.
- 39 Y. Choi, D. S. Mebane, M. C. Lin and M. Liu, Oxygen Reduction on LaMnO_3 -Based Cathode Materials in Solid Oxide Fuel Cells, *Chem. Mater.*, 2007, **19**(7), 1690–1699.
- 40 G. Henkelman, A. Arnaldsson and H. Jónsson, A fast and robust algorithm for Bader decomposition of charge density, *Comput. Mater. Sci.*, 2006, **36**(3), 354–360.



- 41 Z. Wang, R. Peng, W. Zhang, X. Wu, C. Xia and Y. Lu, Oxygen reduction and transport on the $\text{La}_{1-x}\text{Sr}_x\text{Co}_{1-y}\text{Fe}_y\text{O}_{3-\delta}$ cathode in solid oxide fuel cells: a first-principles study, *J. Mater. Chem. A*, 2013, **1**(41), 12932–12940.
- 42 J. A. Dawson, J. A. Miller and I. Tanaka, First-Principles Insight into the Hydration Ability and Proton Conduction of the Solid State Proton Conductor, Y and Sn Co-Doped BaZrO_3 , *Chem. Mater.*, 2015, **27**(3), 901–908.
- 43 B. Merinov, Proton diffusion pathways and rates in Y-doped BaZrO_3 solid oxide electrolyte from quantum mechanics, *J. Chem. Phys.*, 2009, **130**(19), 1021.
- 44 M. S. Islam, P. R. Slater, J. R. Tolchard and T. Dinges, Doping and defect association in $\text{AZrO}_{(3)}$ ($A = \text{Ca}, \text{Ba}$) and $\text{LaMO}_{(3)}$ ($M = \text{Sc}, \text{Ga}$) perovskite-type ionic conductors, *Dalton Trans.*, 2004, **19**(19), 3061–3066.
- 45 S. J. Stokes and M. S. Islam, Defect chemistry and proton-dopant association in BaZrO_3 and BaPrO_3 , *J. Mater. Chem.*, 2010, **20**(30), 6258–6264.
- 46 M. E. Björketun, P. G. Sundell, G. Wahnström and D. Engberg, A kinetic Monte Carlo study of proton diffusion in disordered perovskite structured lattices based on first-principles calculations, *Solid State Ionics*, 2005, **176**(39), 3035–3040.

