Sr$_2$Fe$_{1.4}$Mn$_{0.1}$Mo$_{0.5}$O$_{6-\delta}$ perovskite cathode for highly efficient CO$_2$ electrolysis

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High-temperature solid oxide cells afford chemical storage of renewable electricity. In particular, the electrochemical conversion of the greenhouse gas CO$_2$ is attracting increasing interest to facilitate a sustainable energy technology. In this work, the effectiveness of perovskite-structured Sr$_2$Fe$_{1.4}$Mn$_{0.1}$Mo$_{0.5}$O$_{6-\delta}$ (SFMM0.1) for use as cathode material for CO$_2$ electrolysis has been investigated. Both parent Sr$_2$Fe$_{1.5}$Mo$_{0.5}$O$_{6-\delta}$ (SFM) and SFMM0.1 are found to be redox stable in air and 5% H$_2$/Ar at 850 °C. Electrical conductivity relaxation experiments and first-principle calculations reveal that oxygen transport, CO$_2$ adsorption and reduction kinetics are enhanced upon doping of SFM with Mn. The faster CO$_2$ reduction kinetics observed for SFMM0.1 relative to SFM is reflected in a lower polarization resistance when both materials are used as single-phase electrodes in symmetrical cells. The polarization resistance in 50% CO/CO$_2$ at 800 °C decreases from 1.15 Ω cm$^2$ for SFM to 0.60 Ω cm$^2$ for SFMM0.1. Under similar conditions, the polarization resistance decreases further to 0.50 Ω cm$^2$ for a symmetrical cell with dual-phase SFMM0.1-SDC (samarium-doped ceria) electrodes. Unprecedented performance is demonstrated when SFMM0.1-SDC is integrated as the cathode in a solid oxide cell for electrolysis of pure CO$_2$, achieving a current density of 1.35 A cm$^2$ at 800 °C at 1.5 V.

Introduction

There is a worldwide interest to convert surplus electricity generated from renewables to storable fuels such as hydrogen, methane and liquid fuels.$^{1-3}$ In particular, solid oxide electrolysis of the greenhouse gas CO$_2$, with subsequent conversion of the products to useful fuels, attracts great interest.$^4$ Currently, the nickel-yttria-stabilized zirconia (Ni-YSZ) cermet electrode is the most commonly used cathode in high-temperature solid oxide electrolysis cells (SOECs) due to its high electrocatalytic activity toward CO$_2$ reduction. Its inherent drawback, however, is the limited redox stability of Ni. Under typical operating conditions of an SOEC device, Ni is easily oxidized to NiO, Ni(OH)$_2$ or highly volatile Ni(OH)$_2$, when the concentration of the reducing gas is too low and, hence, the oxygen partial pressure gets too high.$^5$ Furthermore, degradation and/or delamination may occur when the current density exceeds values of the order of $\sim$1 A cm$^{-2}$. Consequently, researchers still endeavor to find alternative cathode materials, thereby focusing on mixed ionic-electronic conducting oxides capable of redox changes, while retaining (long-term) stability under the harsh conditions of SOEC operation. Compared to the Ni-YSZ cermet electrode, where the electrochemical reduction is confined to the vicinity of the triple-phase boundary (TPB) lines, ceramic oxides with mixed ionic-electronic conductivity have the advantage in that they extend the active region over the entire gas/solid phase interface. Many mixed-conducting perovskite-type oxides have been tested for use as cathode for CO$_2$ electrolysis such as La$_{0.7}$Sr$_{0.25}$Cr$_{0.5}$Mn$_{0.5}$O$_3$ (LSCM)$^{6-8}$, La$_{0.8}$Sr$_{0.2}$FeO$_3$ (LST)$^{9-10}$, La$_{0.8}$Sr$_{0.2}$FeO$_3$ (LSF)$^{11}$ and Sr$_2$Fe$_{1.5}$Mo$_{0.5}$O$_{6-\delta}$ (SFM)$^{12-15}$ Amongst these, the iron-based materials LSF and SFM are reported to exhibit the highest electrocatalytic activity for CO$_2$ reduction in the intermediate to high temperature range (600–850 °C).

Isomorphic substitution of A- or B-sites by other types of cations is a known strategy to optimize the performance of perovskite oxide electrodes.$^{16-25}$ In previous research,$^{13-15}$ we showed that the double perovskite SFM is a promising electrode material for pure CO$_2$ electrolysis. Aiming at further promoting the rate of CO$_2$ reduction at the SFM surface, we have investigated in this work the influence of partial substitution of Fe in SFM by Mn (to produce Sr$_2$Fe$_{1.4}$Mn$_{0.1}$Mo$_{0.5}$O$_{6-\delta}$ (SFMM0.1)) on its effectiveness as a cathode material for CO$_2$ electrolysis. Density functional theory (DFT) computations are performed to evaluate oxygen migration and the interaction between adsorbed CO$_2$ and the Mn-doped SFM surface.

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Experimental

Synthesis

The citric acid-glycine combustion method was used to synthesize $\text{Sr}_x\text{Fe}_{1-x}\text{Mn}_x\text{Mo}_{2}O_{6}$ (SFMM$x$; $x = 0, 0.1, 0.2, 0.3, 0.4, 0.5$) powders. $\text{Sr(NO}_3)_2$, $\text{Fe(NO}_3)_3 \cdot 9\text{H}_2\text{O}$, $(\text{CH}_3\text{COO})_2\text{Mn} \cdot 4\text{H}_2\text{O}$ and $(\text{NH}_4)_2\text{Mo}_7\text{O}_{24} \cdot 4\text{H}_2\text{O}$ (purity of 99.0%) in the appropriate stoichiometric ratios were dissolved in distilled water. To this solution glycine and citric acid were added in a molar ratio of citric acid : glycine : total metal cations of 1.5 : 2 : 1. All the above chemicals with high purity (>99.0%) were purchased from Sinopharm Chemical Reagent Co., Ltd. After stirring with a magnetic mixer for 2 h, the solution was put into an induction cooker and heated to self-combustion, producing dark brown ashes. The ashes were collected and annealed at 1100 °C for 5 h in air, using a heating rate of 3 °C min$^{-1}$, and subsequently furnace-cooled to room temperature. $\text{Ce}_0.8\text{Sm}_0.2\text{O}_1.9$ (SDC) was used as electrode additive and was prepared using a similar sol–gel method. The combustion product was annealed at 600 °C for 2 h to form the phase-pure fluorite phase as checked by X-ray powder diffraction (see below). Powders of $\text{La}_0.9\text{Sr}_{0.1}\text{Ga}_{0.8}\text{Mg}_{0.2}\text{O}_3$ (LSGM) and $\text{La}_{0.6}\text{Sr}_{0.4}\text{Co}_{0.8}\text{Fe}_{0.2}\text{O}_3$ (LSCF), both from Fuel Cell Materials, USA, were used without any further treatment.

Characterization methods

The phase purity of powders was checked by X-ray diffraction using a Rigaku D/Max 2100 diffractometer with Cu-Kα radiation ($\lambda = 1.5418$ Å). XRD patterns were recorded in the range 20–80° with a step size of 0.02° and a scan speed of 1° min$^{-1}$ at 40 kV and 200 mA. The GSAS software package with the EXPGUI interface was used for Rietveld analysis. The JSM-6700F (JEOL GmbH, Germany) scanning electron microscope (SEM) was used for microstructural investigations. X-ray photoelectron spectroscopy (XPS) analysis on powder samples was conducted using a Kratos Analytical AXIS 165 with a monochromatic Al-Kα source for determination of the oxidation states. Measurements were performed on an as-synthesized SFMM0.1 powder that obtained after ex situ reduction at 850 °C in flowing 5% H$_2$/Ar (30 ml min$^{-1}$) for 12 h. The reduced powder was furnace-cooled to room temperature after which the gas flow was stopped. Care was taken to avoid re-oxidation of the sample (enclosed in a plastic sealed bag) during transfer to the XPS apparatus. Temperature-programmed desorption (TPD) was examined from 25 °C to 900 °C at 10 °C min$^{-1}$ in Ar (30 ml min$^{-1}$), using a Thermo-Finnigan TPDRO 1100 (Thermo Scientific, USA). Powder samples were pre-treated in pure CO$_2$ at 200 °C for 30 min. The TPD signal was normalized to the surface area of the sample. Brunauer–Emmett–Teller (BET) surface area measurements were performed on a 3H-3000A system (Beishide instrument S&T Co., Ltd, China) using nitrogen as probe gas at 77 K. Prior to measurements the samples were degassed under vacuum at 150 °C. Raman measurements were performed on a Renishaw inVia reflex Raman spectrometer (Renishaw plc, Wotton-under-Edge, UK).

Electrical conductivity and electrical conductivity relaxation measurements

Dense SFM and SFMM0.1 bars were prepared for electrical conductivity and electrical conductivity relaxation (ECR) measurements. To this end, the corresponding powder was mixed with 4 drops of a dispersed PVA solution per gram of powder and milled for 2 h. The infiltrated powder was uniaxially pressed into a rectangular bar under 300 MPa, and subsequently sintered at 1400 °C in air for 5 h to dimensions of $\sim 20 \times 4.8 \times 0.5$ mm$^3$. The relative density was in excess of 99.6% as measured by Archimedes’ method. Conductivity measurements were performed in the temperature range 700–850 °C, using a dc four-probe technique (Keithley, 2001-785D). Silver wires were used as probes, and silver gel (DAD87, Shanghai Institute of Synthetic Resin) was used to improve the contacts between the wires and the sample. The temperature was changed step-wise with intervals of 50 °C, using heating and cooling rates of 5 °C min$^{-1}$. The sample was held at each temperature for 10 h to ensure full equilibration with the ambient before data acquisition. For ECR measurements, the transient conductivity was measured following switching of the gas stream from 66.7% CO/CO$_2$ to 50% CO/CO$_2$ at 200 ml min$^{-1}$ (STP). The gas switching was achieved in less than 1 s. All gases were purchased from Nanjing special gas factory Co., Ltd. The transient conductivity was normalized and fit to the appropriate solution of Fick’s second law to obtain the surface exchange coefficient ($k_{\text{chem}}$) and the chemical diffusion coefficient ($D_{\text{chem}}$). Detailed descriptions of the ECR technique and the model used for data fitting are given elsewhere.27

Electrochemical measurements

Electrolyte-supported symmetrical cells and full solid oxide electrolysis cells were prepared to test the performance of electrodes. Pure SFMM0.1 powder or SFMM0.1-SDC mixed powders (weight ratio SFMM0.1 : SDC = 6 : 4) were grounded for 1 h and dispersed in a solution of ethyl cellulose in x-terpineol. The obtained electrode ink was screen-printed 3 times with intermediate drying at 90 °C for 2 h onto a 400 μm thick LSGM electrolyte disc (of 9 mm in diameter), followed by calcination at 1000 °C for 2 h under stagnant air to form a 10–12 μm thick porous electrode (with effective area of 0.91 cm$^2$).

For symmetrical cells, two identical electrodes were screen-printed on both sides of the electrolyte disc. The symmetrical cells were employed to measure the polarization resistance of the SFMM0.1 and SFMM0.1-SDC electrodes in 50% CO/CO$_2$. For the solid oxide electrolysis cells, LSCF-SDC and SFMM0.1-SDC composites were screen printed on opposite sides of the electrolyte disc, as oxygen and fuel electrode, respectively. The ink for the oxygen electrode was prepared in a similar way as described above. The cell was sealed to one end of an alumina tube, using silver paste, calcined at 800 °C for 2 h in stagnant air, and then the cathode exposed immediately to flowing CO$_2$ (30 ml min$^{-1}$). Silver mesh current collectors were attached to each electrode using silver paste for both the symmetrical-cell and the solid-oxide-electrolysis-cell tests. The schematics of
the electrochemical cell and experimental set-up used for CO₂
electrolysis tests is shown in Fig. S1.†

The electrochemical tests were performed in the temperature
range 700 °C–850 °C. The impedance of the symmetrical
cells was measured using a Solartron 1260 impedance/gain
phase analyzer in the range of frequencies from 0.01 Hz to 1
MHz under open circuit conditions and at an excitation voltage
of 10 mV. Polarization curves (linear sweep rate of 0.1 V s⁻¹ from
0 to 1.6 V) and impedance of the solid oxide electrolysis cells
were measured using a Princeton electrochemical workstation
(VMC-4 multichannel). The impedance spectra were fitted to an
equivalent circuit using ZView® software (Scribner, Associates
Inc., USA). The spectra obtained for the symmetrical cells were
further analyzed by the distribution of relaxation times (DRT)
method using a home-made software program. The program
employs Tikhonov regularization to enable the DRT
deconvolution.

Computational methods

Density functional theory (DFT) calculations were carried out
using the Vienna ab initio simulation package (VASP).²⁸,²⁹ The
generalized gradient approximation (GGA) with the Perdew-
Burke–Ernzerhof (PBE) functional was adopted to approximate
the exchange-correlation term.³⁰ The projector augmented wave
(PAW) method with C (1s²2s²2p²), O (1s²2s²2p⁴), Mn ([Ar]
3d⁵4s²), Fe ([Ar]3d⁶4s²), Sr ([Kr]5s²), and Mo ([Kr]4d⁵5s¹) was
used to describe the interaction between the ionic core and
valence electrons. Following previous DFT studies on SFM,³¹,³²
the exchange correlation energy of the 3d electrons was calcu-
lated within the generalized gradient approximation (GGA) with
an Uₜₜ parameter of 4 eV for Mn and Fe, and with no Uₜₜ
parameter for Mo. The kinetic energy cutoff was 520 eV, and the
energy convergence criterium was set to 10⁻⁵ eV. Structure
optimization was carried out until the forces on each atom were
less than 0.03 eV Å⁻¹. All the calculations were spin-polarized
and ferromagnetic states were used.

Calculations were performed on the basis of the 40 atoms
containing [Sr₂Fe₆MoO₁₆] unit cell of the double perovskite
structure of Sr₂Fe₁₋ₓMnₓMo₀.₅O₆, represented in space group Fm₃m.
A 6 × 6 × 6 Monkhorst–Pack³⁴ k-point grid was used to sample
the Brillouin zone. For Mn-doped SFM, one Fe atom was
substituted by Mn in the optimized cell, leading to Sr₂Fe₁₋ₓ₋ₓMnₓMo₀.₅O₆ with x = 0.25 (SFMM0.25), as shown in Fig. 1.
Oxygen vacancies were modeled by removing a single neutral
atom from the unit cell, leading to Sr₂Fe₁₋ₓ₋ₓ₋ₓMnₓMo₀.₅O₆₋₋₋δ
with δ = 0.25. The oxygen formation energy was calculated according to

\[ E_o = E_{\text{defect}} - E_{\text{ideal}} + E\left(\frac{1}{2}\text{O}_2\right) \quad (1) \]

where \( E_{\text{defect}} \) and \( E_{\text{ideal}} \) are the total energies of the non-stoi-
chiometric and stoichiometric structures, respectively, and
\( E\left(\frac{1}{2}\text{O}_2\right) \) is the energy of a free O₂ molecule. The projected
densities of states (PDOS) were obtained as output of the VASP
code. Atomic charges were calculated using the Bader approach.

The minimum energy path (MEP) of oxygen migration and the
energy of the transition state (TS) were computed using the
climbing image nudged elastic band (CINEB) method.³³ A total
of 5 intermediate images was created with the forces minimized
to 0.03 eV Å⁻¹.

We used eight-layer SFM(001) and SFMM0.25(001) slabs,
separated by 15 Å of vacuum, to investigate the interaction
between CO₂ and their (001) surfaces with different surface
terminations. The bottom four layers were fixed at their bulk
position, while the top four layers were relaxed. A dipole
correction was applied to the direction normal to the surface,
while the Brillouin zone center was sampled at the gamma point
with a 3 × 3 × 1 k-point grid. The adsorption energy of a single
CO₂ molecule was computed according to

\[ E_{\text{ads}} = E_{\text{CO}_2(001)} - E_{(001)} - E(\text{CO}_2) \quad (2) \]

where \( E_{\text{CO}_2(001)} \) is the total energy of one CO₂ molecule adsorbed
on the (001) surface, \( E_{(001)} \) is the total energy of the free surface,
and \( E(\text{CO}_2) \) is the energy of a free CO₂ molecule.

Results and discussions

Phase analysis, structural and chemical stability

Fig. S2† shows room temperature X-ray diffraction patterns of
samples Sr₂Fe₁₋ₓ₋ₓ₋ₓMnₓMo₀.₅O₆₋₋₋δ (SFMMx, x = 0, 0.1, 0.2, 0.3,
0.4, 0.5). Consistent with previous work SFM (x = 0) is found to
adopt a cubic structure (space group Fm₃m). The XRD patterns
of the Mn-doped samples resemble those of undoped SFM (x = 0),
demonstrating that the former compositions adopt a struc-
ture similar to that of the parent phase. No shifts in the
diffraction peaks are observed upon substitution of Fe by Mn,
which may be accounted for by the very similar radii of Fe³⁺ and
Mn²⁺ ions.³⁶ However, some extra weak re-
dlections are found in
the patterns of samples with \( x = 0.3, 0.4 \) and 0.5. Their presence
is assigned to SrMoO₄ impurity phase. Previous research has
revealed that SFM forms a solid solution up to composition
Sr₂Fe₁₋ₓ₋ₓ₋ₓMn₀.₅₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋₋GraphNode 1 Unit cell of Sr₂Fe₁₋ₓ₋ₓ₋ₓMnₓMo₀.₅O₆ (SFM) in which one Fe atom is substituted by Mn to produce Sr₂Fe₁₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋ₓ₋xDCO₂ molecule was computed according to

\[ E_{\text{ads}} = E_{\text{CO}_2(001)} - E_{(001)} - E(\text{CO}_2) \quad (2) \]

where \( E_{\text{CO}_2(001)} \) is the total energy of one CO₂ molecule adsorbed
on the (001) surface, \( E_{(001)} \) is the total energy of the free surface,
and \( E(\text{CO}_2) \) is the energy of a free CO₂ molecule.
clarify this. To avoid any phase coexistence, SFMM0.1 was selected for further investigations in this study.

The structural stability of SFMM0.1 was tested by firing in 5% H₂/Ar at 850 °C for 12 h. Rietveld refinements of the XRD patterns (Fig. 2) show that the perovskite structure is retained after firing and no extra peaks are observed. Lattice parameters and reliability factors obtained from Rietveld refinements of the patterns obtained after firing in air or 5% H₂/Ar are shown in Table S1.† The cubic lattice parameter of as-synthesized SFMM0.1 slightly increases from 7.84176(4) Å to 7.86728(4) Å after firing under reducing conditions. This increase correlates with a lowering in the average oxidation state of the B-site cations (Fe, Mo, Mn) in SFMM0.1 and, hence, with an apparent increase in the concentration of oxygen vacancies.

Fig. 3a shows XPS spectra for the B site elements in as-synthesized SFM.0.1. The Fe 2p⁴/₂ peak can be deconvoluted into three constituents, corresponding to Fe²⁺, Fe³⁺ and Fe⁴⁺, while the Mo 3d⁵/₂ peak can be deconvoluted into Mo⁵⁺ and Mo⁶⁺ peaks.†† Compared with pure SFM,‡ Mn-doping reduces the average oxidation state of Fe from +3.16 to +3.04, while the value of 6.00 for Mo remains unaffected. The Mn 2p⁴/₂ peak can be deconvoluted into Mn²⁺, Mn³⁺ and Mn⁴⁺ peaks centered at 641.8 eV, 643.1 eV and 643.5 eV, respectively,‡‡ yielding an average oxidation state for Mn of +2.70. The XPS results thus confirm that the average oxidation state of the B site cations in as-synthesized SFM is reduced from +3.87 to +3.76 in SFMM0.1. Upon treatment of SFMM0.1 in 5% H₂/Ar (Fig. 3b), the average oxidation states of Mn, Fe and Mo reduce from +2.7 to +2.27, from +3.04 to +2.92 and from +6.0 to +5.9, respectively (see Table S2†), demonstrating the high redox activity of the Mn dopant. The average oxidation state of the B site cations in reduced SFMM0.1 is calculated to be +3.63. Experimental and deconvoluted O 1s XPS spectra for as-synthesized SFMM0.1 at 1100 °C in air and after reduction at 850 °C in 5% H₂/Ar are given in Fig. S3.†

To analyze the influence of vacancy formation on oxidation states, we computed effective Bader charges of the atoms in SFMM0.1 before and after creation of an oxygen vacancy, which gives a semi-quantitative picture of the redistribution of the electron density upon removal of oxygen. Corresponding results are listed in Table S3.† The Bader charges of the redox active ions are +1.69 e for Fe and +1.86 e for Mn, both of which are much lower than the corresponding values derived from data of XPS, suggesting a significant degree of covalency in the Fe–O and Mn–O bonds.†† After removal of neutral oxygen, the redistribution of the electrons leads to Bader charges of +1.62 e for Fe and +1.67 e for Mn. The obtained results demonstrate the high redox ability of the Mn dopant relative to that of atoms of the host lattice.

Finally, the chemical stability of SFMM0.1 powder in CO₂ atmosphere was checked. XRD analysis revealed no phase change after exposure to pure CO₂ at 850 °C for 12 h.

**Electrical conductivity**

Electronic transport in SFM occurs by a small polaron hopping mechanism and proceeds via percolation pathways along the Fe–O–Fe bonding network in the crystal. As shown in Fig. S4† the conductivity of SFM in air reaches a value of about 20 S cm⁻¹ at 800 °C. With the addition of Mn the conductivity at this temperature decreases to 16 S cm⁻¹, suggesting that under oxidizing conditions the concentration of p-type charge carriers is slightly reduced by the doping with Mn. Arrhenius plots of the electrical conductivities of SFM and SFMM0.1 in both 5% H₂/Ar and 50% CO/CO₂ are shown in Fig. 4, clarifying that under the prospective operating conditions of CO₂ electrolysis the conductivity of SFM is enhanced upon doping with Mn. The conductivity found for SFM in 5% H₂/Ar, 16 S cm⁻¹ at 800 °C, is in good agreement with previous results, increasing to a value of 25 S cm⁻¹ for SFMM0.1. The higher conductivity exhibited by both materials in the more reducing atmosphere, i.e. in 5% H₂/Ar relative to that in 50% CO/CO₂, occurs on account of the predominant n-type conductivity under reducing conditions. Introducing Mn and/or oxygen nonstoichiometry in SFM may trigger fluctuations in the electronic...
band structure. Previous PDOS calculations on SFM showed that the Fe 3d and O 2p states are strongly hybridized, accounting for the high electronic conductivity of SFM and other Fe-rich perovskites. Our calculations show that hybridization around the Fermi level is preserved in perfect ($d = 0$) and oxygen-defective ($d = 0.25$) SFM0.25.

Electrical conductivity relaxation

ECR experiments on SFM and SFM0.1 ceramics were conducted to determine the influence of partial substitution on oxygen diffusion and the material’s CO$_2$ reduction ability. Normalized conductivity curves for both samples acquired at different temperatures following switching of the gas stream from 66.7% CO/CO$_2$ to 50% CO/CO$_2$ are shown in Fig. 5. Relative to that for SFM, faster re-equilibration kinetics is found for SFM0.1. This difference in re-equilibration time scales is more pronounced at lower temperatures. The solid lines in Fig. 5 are from curve fitting of the data to model equations, revealing that under the conditions of the experiments the relaxation behavior
of both samples is under mixed control of bulk diffusion and surface exchange. Arrhenius plots of $D_{\text{chem}}$ and $k_{\text{chem}}$ are shown in Fig. 6, which makes evident that below about 850 °C partial substitution of Fe with Mn in SFM increases the magnitude of both parameters. At 700 °C, the increase of both parameters relative to the values observed for pure SFM is about 3–4 times. Further note from Fig. 6a and b that substitution leads to a profound decrease in the apparent activation energies of both parameters $D_{\text{chem}}$ and $k_{\text{chem}}$. Secondary electron SEM images of SFM and SFMM0.1 recorded after the ECR experiments are shown in Fig. S5.† Comparison with SEM images taken before the experiments (not shown) indicated no apparent changes in the morphology of both samples.

**Oxygen migration**

A high concentration of oxygen vacancies in SFM and Mn-doped SFM facilitates transport of $O^{2-}$ ions via a vacancy diffusion mechanism. Another factor that dominates the $O^{2-}$ diffusivity is the migration barrier ($E_{\text{mig}}$) associated with hopping along the diffusion path to the nearest-neighbor vacancy. Our CINEB calculations predict the formation energy of oxygen vacancies, $E_v(\text{Fe–VO}–\text{Fe})$, corresponding with the removal of oxygen in Fe–O–Fe bonds, in both structures to be the lowest compared with values for formation of oxygen vacancies between dissimilar metal atoms. For SFMM0.25 it is found that $E_v(\text{Fe–VO}–\text{Fe})$ is 0.17 eV, while $E_v(\text{Fe–VO}–\text{Mo})$ and $E_v(\text{Fe–VO}–\text{Mn})$ are 1.32 eV and 0.36 eV, respectively. The higher values for the latter two may be expected on the basis of the weaker Fe–O bond compared to
Mn–O and Mo–O bonds. The value obtained for \( E_v(\text{Fe–}V\text{O–Fe}) \) in SFM is 0.13 eV, which is notably lower than that for SFMM0.25, indicates that beyond breaking of the two Fe–O bonds the formation of an oxygen vacancy is also affected by the local surroundings. On account of the low values of \( E_v(\text{Fe–}V\text{O–Fe}) \) in SFM and SFMM0.25, we envision that O\(_2\)/C\(_0\) transport in both structures is dominated by hopping between adjacent (Fe–O–Fe) sites and (Fe–V\text{O–Fe}) sites such migration involves passage over a saddle point on the potential energy surface and, consequently, the O\(^{2-}\) ion will follow a curved path as schematically depicted in Fig. 7a. The MEPs obtained by our CINEB calculations (Fig. 7b) show that \( E_m\text{igr} \) reduces from 0.36 eV in SFM to a value as low as 0.15 eV in SFMM0.25. For the activation energy of oxygen self-diffusion, which is given by the sum of \( E_v(\text{Fe–}V\text{O–Fe}) \) and \( E_m\text{igr} \), we thus find values of 0.49 eV and 0.32 eV for SFM and SFMM0.25, respectively. The calculations were carried out for \( \delta = 0.25 \). Though it cannot be excluded that higher migration enthalpies are obtained for lower, probably more realistic oxygen deficiencies, the results clearly indicate that the presence of Mn lowers the migration energy despite the migration path being near the Fe atoms, corroborating with the results from ECR experiments.

**CO\(_2\) adsorption**

Theoretical calculations were carried out to study the interaction of CO\(_2\) with the SFM and SFMM0.25 surfaces. Primary calculations on SFM showed the SrO (100) termination to be the most stable surface. To study the influence of Mn-doping on CO\(_2\) adsorption, however, we focused our calculations on the (Fe, Mn, Mo) O\(_2\)-terminated surface. We first calculate the CO\(_2\) adsorption energies on the perfect \( (i.e., \text{non-defective}) \) surfaces, and then proceed with considering the effect of an oxygen vacancy created next to the adsorption site. CO\(_2\) adsorption energies, C–O bond lengths and O–C–O angles calculated for different adsorbed CO\(_2\) configurations are listed in Table S4.

CO\(_2\) chemisorbs atop surface oxygen atoms. Chemisorption involves charge transfer to the adsorbed CO\(_2\) molecule (and bending of the initially linear CO\(_2\) molecule due to electron transfer to the antibonding \( \pi^* \) orbitals of CO\(_2\) with as a result more negative adsorption energies compared to physisorption. Different oxygen sites are available on the (100) surfaces of SFM and SFMM0.25. These are the Fe–O–Fe and Fe–O–Mo sites on the (Fe, Mo) O\(_2\)-terminated (100) surface of SFM, with respective adsorption energies of 0.98 eV and 0.91 eV. On the (Fe, Mn, Mo) O\(_2\)-terminated (100) surface of SFMM0.25 these are the Fe–O–Mn and Fe–O–Mo sites with respective adsorption energies of 1.00 eV and 1.11 eV. Hence, CO\(_2\) adsorbs preferentially atop the oxygen atom with the lowest oxygen vacancy formation energy. The calculations further show that the CO\(_2\) molecule adsorbs as a monodentate surface carbonate (CO\(_3^{2-}\)) species as illustrated for adsorption on preferential oxygen sites on both surfaces in Fig. 8a and c. For both SFM and SFMM0.1, it is found that the O–C–O angles are bent to about 131°, and that the C–O bond lengths are elongated to values close to 1.27 Å, relative to the gaseous values of 180° and 1.18 Å, respectively. The ideal carbonate ion has trigonal planar symmetry with C–O bond lengths of about 1.28 Å.
surface carbonate ion is considered to be an important intermediate for activation and reduction of CO$_2$.

Next, the effect of a nearest-neighbor oxygen vacancies on CO$_2$ adsorption is considered. A different configuration for CO$_2$ emerges compared with that found upon adsorption on the vacancy-free (100) surfaces as the CO$_2$ molecule now forms a bidentate carbonate configuration with one of the C–O bonds strongly elongated to about 1.31 Å. The bidentate configuration results in a more negative adsorption energy relative to that of the monodentate configuration found on the vacancy-free surface. For SFM the adsorption energy is lowered from $0.98$ eV to $1.23$ eV, while for SFMM0.1 it is lowered from $1.11$ eV to $1.37$ eV. Apart from the lower adsorption energies the distorted CO$_2$ molecule is presumed to be highly activated. The more exothermic values of the CO$_2$ adsorption energies found for SFMM0.25 than for SFM substantiated by the theoretical calculations indicate that partial substitution of Fe with Mn enhances CO$_2$ adsorption. This is further corroborated by the results from temperature-programmed desorption measurements on SFM and SFMM0.1 powder samples that have been treated in pure CO$_2$ at 200 °C for 30 min. Fig. S6† shows that much enhanced desorption is found for SFMM0.1 as compared to SFM which extends up to about 900 °C.

**Electrochemical performance**

Both LSGM electrolyte-supported symmetrical and full electrolysis cells were fabricated. Symmetric cells were employed to measure the polarization resistance ($R_p$) of single-phase SFMM0.1 and dual-phase SFMM0.1-SDC electrodes under reducing conditions. Fig. 9a shows corresponding impedance spectra recorded at 800 °C in 50% CO/CO$_2$. Also included for comparison is the impedance spectrum for a symmetrical cell with SFM electrodes recorded under identical conditions from our previous study. The latter cell was fabricated similarly to the symmetrical cells employed in the present study. Under the specified conditions, $R_p$ for SFM is $1.12$ Ω cm$^2$, while $0.60$ Ω cm$^2$ is found for the SFMM0.1 electrode. Corresponding values at 700 °C are $8.76$ Ω cm$^2$ and $4.36$ Ω cm$^2$ for the SFM and SFMM0.1 electrode, respectively, confirming that $R_p$ decreases significantly with partial substitution of Fe by Mn. The electrode performance further improves for the SFMM0.1-SDC composite electrode: $0.5$ Ω cm$^2$ at 800 °C, and $0.90$ Ω cm$^2$ at 700 °C. The results from symmetrical cells are compiled in Table S5.

A distribution of relaxation times (DRT) analysis was performed on the impedance spectra recorded at 800 °C. In a DRT plot, the location of a peak reflects a characteristic relaxation time, while the integral peak area is a measure of the...
polarization loss. In general, low frequency peaks are assigned to gas diffusion and surface processes such as adsorption/desorption, while high frequency peaks are linked to charge transfer and ion migration processes.

As seen from Fig. 9b multiple peaks can be identified in the DRT curves of the spectra. The high frequency peaks are probably caused by the transformation procedure and have no direct physical meaning. Comparison of the DRT curves shows that doping of SFM with Mn notably lowers the low frequency-peak in the range 0.1–1 Hz, which we tentatively ascribe to lowering of the polarization losses due to enhanced CO₂ adsorption and associated reduction kinetics consistent with the results from DFT calculations and ECR (see above sections Oxygen migration and Electrical conductivity relaxation). For the SFMM0.1-SDC composite electrode, the peak at 0.1–1 Hz has virtually vanished. Fig. S7† shows impedance spectra of the symmetrical cells recorded under open circuit conditions, at 800 °C, in 50% CO/CO₂ and (b) corresponding results from DRT analyses of the spectra. Impedance spectra are shown after subtraction of the apparent electrolyte resistance. Data for SFM in (a) is replotted from ref. 14.

Fig. 9 (a) Impedance spectra of electrolyte-supported symmetrical cells SFM|LSGM|SFMM0.1|LSGM|SFMM0.1 and SFMM0.1-SDC/LSGM/SFMM0.1-SDC recorded under open circuit conditions, at 800 °C, in 50% CO/CO₂, and (b) corresponding results from DRT analyses of the spectra. Impedance spectra are shown after subtraction of the apparent electrolyte resistance. Data for SFM in (a) is replotted from ref. 14.

Fig. 10 Electrochemical performance of a single electrolyte-supported electrolysis cell, LSCF-SDC/LSGM/SFMM0.1-SDC, incorporating composite SFMM0.1-SDC as fuel electrode: (a) I–V curves, and (b) impedance spectra at open circuit voltage. The drawn lines represent fitted data. Measurements were conducted at different temperatures, as indicated. Air was supplied to the anode and pure CO₂ to the cathode.

The performance of the SFMM0.1-SDC composite electrode for CO₂ electrolysis was further tested in a full electrolyte-supported electrolysis cell using LSCF-SDC as oxygen electrode and LSGM as the electrolyte. Corresponding results are shown in Fig. 10a. Using pure CO₂, without addition of reducing gas (e.g., H₂) or pre-reduction of the electrode, the current density at an applied voltage of 1.5 V is found to increase from 0.47 A cm⁻² at 700 °C to 1.80 A cm⁻² at 850 °C. The I–V curves exhibit distinct non-linear behavior. This phenomenon has also been observed in other studies where perovskite-type oxides are used as electrodes for CO₂ or H₂O electrolysis, without pre-mixing of the feed gases with CO or H₂,⁷,⁹,¹⁰,⁴³–⁴⁵ and which is in contrast to the linear I–V curves observed when Ni-YSZ based cathodes are used.

Nyquist plots of the single cell measured at OCV at different temperatures are given in Fig. 10b. The spectra can be fitted using the equivalent circuit $R_s((R_1Q_1)(R_2Q_2)(R_3Q_3))$, where $R_s$ is the apparent ohmic resistance of the electrolyte, $R_i$ ($i = 1, 2$ and 3) is a resistance, and $Q_i$ ($i = 1, 2$ and 3) is a constant phase element. At 800 °C, the total electrode polarization resistance ($R_p = R_1 + R_2 + R_3$) at OCV is 0.58 Ω cm², noting that the LSCF-SDC anode minorly contributes to the value of $R_p$. The excellent performance of the SFMM0.1-SDC electrode is immediately
Table 1 Comparison of literature values of current densities achieved in pure-CO2 electrolysis over different all-ceramic cathodes in electrolyte-supported cells operated at 1.5 V at 800 °C with the value obtained in this study

| Fuel electrode | Current density [A cm⁻²] | Reference |
|---------------|--------------------------|-----------|
| La₀.₇₅Sr₀.₂₅Ce₀.₅Mn₀.₅O₃₋ₓ-SDC | 0.075 | 7 |
| (La₀.₇₅Sr₀.₂₅)₀.₉₅(Cr₀.₅Mn₀.₅)₀.₀₅O₋ₓ-GDC | 0.26 | 8 |
| (La₀.₇₅Sr₀.₂₅)₀.₉₅(Cr₀.₅Mn₀.₅)₀.₀₅-GDC | 0.18 | 8 |
| La₀.₇₅Sr₀.₂₅Ce₀.₅Fe₀.₅O₋ₓ-SDC | 0.09 | 7 |
| La₀.₇₅Sr₀.₂₅Ce₀.₅Fe₀.₅Th₀.₁O₋ₓ | 0.075 | 10 |
| La₀.₇₅Sr₀.₂₅Fe₀.₅Mn₀.₁O₋ₓ-SDC | 0.15 | 10 |
| La₀.₇₅Sr₀.₂₅Fe₀.₅O₋ₓ-SDC | 0.06 | 10 |
| La₀.₂Sr₀.₈Ti₀.₅O₋ₓ-SDC | 0.105 | 11 |
| La₀.₂Sr₀.₈Ti₀.₅Mn₀.₁O₋ₓ-SDC | 0.26 | 11 |
| (La₀.₂Sr₀.₈)₀.₉₅Fe₀.₅Mn₀.₁O₋ₓ-SDC | 0.22 | 11 |
| (La₀.₂Sr₀.₈)₀.₉₅Fe₀.₅Cr₀.₁Mn₀.₁O₋ₓ-SDC | 0.45 | 11 |
| SFMM₀.₁-SDC | 1.35 | This work |

† SDC and GDC refer to Sm₀.₅Ce₀.₅O₂₋ₓ and Gd₀.₅Ce₀.₅O₂₋ₓ, respectively.

Conclusions

In conclusion, an all ceramic cathode has been developed which combines a high stability with a high performance in direct CO₂ electrolysis. When SFMM0.1-SDC is incorporated as cathode in an electrolyte-supported SOEC for electrolysis of pure CO₂, a current density of 1.35 A cm⁻² is achieved at 800 °C under an applied bias of 1.5 V, without the addition of a reducing gas such as H₂. The enhanced cell performance can be accounted for by data from electrical conductivity relaxation experiments and first-principle calculations, revealing that oxygen transport, CO₂ adsorption and reduction kinetics are enhanced upon doping of parent SFM with Mn. The SFMM0.1-SDC composite electrode further exhibits excellent coking resistance. This work shows that SFMM0.1-SDC is a promising alternative to replace metal-based electrodes for direct CO₂ electrolysis.

Conflicts of interest

There are no conflicts to declare.

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