Double-Perovskite Anode Materials Sr$_2$MMoO$_6$ (M = Co, Ni) for Solid Oxide Fuel Cells

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Double-perovskites Sr$_2$MMoO$_6$ (M = Co, Ni) have been investigated as anode materials for a solid oxide fuel cell. At room temperature, both Sr$_2$CoMoO$_6$ and Sr$_2$NiMoO$_6$ are tetragonal (I4/m). X-ray absorption spectroscopy confirmed the presence of Co$^{3+}$/Mo$^{6+}$ and Ni$^{3+}$/Mo$^{6+}$ pairs in the oxygen stoichiometric compounds. The samples contain a limited concentration of oxygen vacancies in the reducing atmospheres at an anode. Reoxidation is facile below 600 °C; they become antiferromagnetic at low temperatures $T_N = 37$ and 80 K for M = Co and Ni, respectively. As an anode with a 300 μm thick La$_{0.8}$Sr$_{0.2}$Ga$_{0.83}$Mg$_{0.17}$O$_{2.815}$ electrolyte and SrFe$_{0.2}$Co$_{0.8}$O$_{3-δ}$ as a cathode, Sr$_2$CoMoO$_6$ exhibited maximum power densities of 735 mW/cm$^2$ in H$_2$ and 527 mW/cm$^2$ in wet CH$_4$ at 800 °C; Sr$_2$NiMoO$_6$ shows a notable power output only in dry CH$_4$. The high performance of Sr$_2$CoMoO$_6$ in wet CH$_4$ may be due to its catalytic effect on steam reforming of methane, but some degradation of the structure that occurred in CH$_4$ obscures identification of the catalytic reaction processes at the surface. However, the stronger octahedral-site preference of Ni$^{3+}$ versus Co$^{3+}$ can account for the lower performance of the M = Ni anode.

Introduction

The solid oxide fuel cell (SOFC) is an electrochemical device that can be used for either stationary or mobile generation of electrical energy from a gaseous fuel. The conventional SOFC, which uses Y$_2$Zr$_{1-x}$Ti$_x$O$_{2−δ}$ (YSZ) as the electrolyte and a porous Ni–YSZ cermet anode, is commercially viable with pure H$_2$ or syngas as the fuel; but this anode is fouled by carbon deposition and sulfur poisoning when operated on natural gas. Development of an anode material that can operate on natural gas would provide a cheaper, more convenient SOFC. For this application, oxides that are mixed oxide-ion/electron conductors (MIECs) in the reducing atmosphere at the anode have been under investigation. The double-perovskite Sr$_2$MgMoO$_6$ is a promising MIEC with an excellent tolerance to sulfur that gives direct electrochemical oxidation in dry methane at 800 °C. Moreover, La-doped Sr$_2$MgMoO$_6$ performs somewhat better on natural gas. These results have prompted a study of other members of the Sr$_2$MMoO$_6$ family containing a 3d-block transition-metal M to investigate the role of the M cation.

The ordered double-perovskites A$_2$BB′O$_6$ have alternating BO$_{6/2}$ and B′O$_{6/2}$ corner-shared octahedra. Substitution at A or B sites can alter the cation valence and oxygen-vacancy concentration. As is well-known, cation valence and oxygen-vacancy concentration play important roles in the physical and electrochemical properties of the double perovskites. In Sr$_2$MgMoO$_6$, Mg ions show unchanged divalence; only the valence of Mo ions changes from $3t$ to $2t$ in the transition-metal M to investigate the role of the M cation.

The high performance of Sr$_2$CoMoO$_6$ in wet CH$_4$ may be due to its catalytic effect on steam reforming of methane, but some degradation of the structure that occurred in CH$_4$ obscures identification of the catalytic reaction processes at the surface. However, the stronger octahedral-site preference of Ni$^{3+}$ versus Co$^{3+}$ can account for the lower performance of the M = Ni anode.

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electrochemical performances of the double-perovskites Sr$_2$MoMoO$_6$ (M = Co, Ni) as anodes of a SOFC operating on dry and wet methane and on H$_2$.

**Experimental Section**

Sr$_2$CoMoO$_6$ and Sr$_2$NiMoO$_6$ samples were synthesized via a sol–gel method with stoichiometric SrCO$_3$ (Arlitest, 99%), (NH$_4$)$_2$Mo$_3$O$_9$·4H$_2$O (Fisher Scientific, assay MoO$_3$, 81.5%), and Co$_2$O$_4$ (Alfa, 99%) or NiO (Alfa, 99%) as the starting materials. SrCO$_3$ and Co$_2$O$_4$, NiO were first dissolved with diluted nitric acid and then mixed with (NH$_4$)$_2$Mo$_3$O$_9$·4H$_2$O. Ethylene diaminetetraacetic acid was added as a complexant to achieve a clear aqueous solution. The pH of the solution was adjusted to 9 with acetic acid to achieve a clear aqueous solution. The pH of the solution was adjusted to 9 with ammonia. The solution was evaporated on a hot plate to become a gel. The gel was first decomposed at 400 °C in air for 6 h and then calcined at 800 °C in air for 10 h. The calcined powder was pelletized and finally sintered at 1250 °C in air for 24 h to achieve a pure phase. Synthesis of other compounds, L$_{40}$Sr$_{5}$Sr$_{3}$Ga$_{0.5}$Sb$_{0.5}$Mo$_{0.17}$O$_{2.815}$, SrCo$_3$Fe$_2$O$_{3-δ}$, and L$_{40}$Ce$_{0.5}$O$_{2-δ}$, has been described elsewhere.

The phase purity and the lattice parameters of the samples were checked by X-ray powder diffraction (XRD, Philips X-pert, Cu Kα radiation). The diffraction profiles were analyzed with a Rietveld refinement program, RIETAN 2000. Redox behaviors of the samples were investigated by thermogravimetric analysis (TGA, Perkin-Elmer Pyris 1 and Netzsch STA 449 C) in air and in 5% H$_2$/Ar gas flows in the temperature range from room temperature to 1000 °C. In these TGA experiments, the amount of sample powder was ~15 mg and the heating/cooling rate was either 2 or 10 °C/min. Micrographs were taken by a scanning electron microscope (SEM, Hitachi: S4500). Magnetization measurements were made with a superconducting quantum interference device (Quantum Design: MPMS-XL5). The conductivity was measured by a standard dc four-probe method with our own setup. The samples for conductivity measurement were polished into rectangular bars; Pt wire and Pt paste were used to make the four probes. Before measurement, the samples were reduced in 5% H$_2$/Ar in our own setup at 800 °C for 20 h to ensure formation of oxygen vacancies. Oxygen partial pressure was monitored with a Thermalox CG1000 oxygen analyzer (Ametek).

The Mo L$_3$-edge, Ni K-edge, and Co K-edge X-ray absorption spectroscopy (XAS) measurements were performed in the fluorescence mode and on powdered samples on beamline X-19A at the National Synchrotron Light Source, Brookhaven National Laboratory. A double-crystal Si(111) monochromator was used. The XAS samples were prepared by dusting a fine powder of the samples onto scotch tape. In the case of the Ni and Co K-edge measurements, the X-ray beam transmitted through the samples allowed transmission-mode measurements along with edges of simultaneously run standards located on the down-beam side of the sample.

The absolute energy calibration was set to the elemental edge (first inflection point). The relative energy scale was maintained to better than ±0.05 eV with the simultaneously run standards. In the case of the Mo L$_3$-edges, standards were run periodically in the sample sequence and the energy scale is better than ±0.1 eV.

Single SOFC test cells were fabricated by an electrolyte-supported technique with 300 μm thick L$_{40}$Sr$_{5}$Sr$_{3}$Ga$_{0.5}$Sb$_{0.5}$Mo$_{0.17}$O$_{2.815}$ (LSGM) as the electrolyte and SrCo$_3$Fe$_2$O$_{3-δ}$ (SCF) as the cathode. The fabrication method of the single cell has been described in detail in our previous work. A thin buffer layer of L$_{40}$Ce$_{0.5}$O$_{2-δ}$ (LDC) between the anode and the electrolyte was used to prevent interdiffusion of ionic species between anode and electrolyte.

| Table 1. Room-Temperature Lattice Parameters, Bond Lengths, Bond Angles, Density (d), Atom Occupancy (q$_{M}$), and Degree of Cationic Ordering ($ξ$) Obtained by Rietveld Refinement on XRD Patterns for Sr$_2$MoMoO$_6$ (M = Co, Ni) |
|------------------|------------------|------------------|
| M    | Co   | Ni   |
| space group | $a$ (Å) | $b$ (Å) | $c$ (Å) | $d$ (g/cm$^3$) | M–O (Å) | M–O (Å) | M–O (Å) | $\beta$ (°) | $\gamma$ (°) | $\alpha$ (°) | $R_p$ (%) | $R_{wp}$ (%) |
|---------|-------|-------|-------|--------|---------|---------|---------|--------|----------|-----------|-----------|----------|-----------|
| Co      | 5.5726(3) | 5.5726(3) | 7.9575(3) | 5.723 | 2.044(×2) | 2.039(×4) | 2.041 | 1.935(×2) | 1.935(×4) | 1.935 | 1.935(×2) | 1.935(×4) | 0.942 | 0.958 |
| Ni      | 5.5463(2) | 5.5463(2) | 7.8933(3) | 5.825 | 2.003(×2) | 1.987(×4) | 1.993 | 1.953(×2) | 1.951(×4) | 1.952 | 1.952(×2) | 1.952(×4) | 0.958 | 0.958 |

The XRD patterns show that the samples were double-perovskite phases. Sr$_2$CoMoO$_6$ and Sr$_2$NiMoO$_6$ both have a tetragonal structure with space group $I4/m$. The lattice parameters, bond lengths, bond angles, and site occupancies obtained by Rietveld refinement are displayed in Table 1. The lattice cell volume of M = Co is larger than that of M = Ni; the mean bond length (Co–O) is longer than that of (Ni–O). The order is consistent with their ionic radii (sixfold coordination), i.e., Co$^{2+}$ 0.745 Å (HS), Ni$^{2+}$ 0.69 Å. The bond length (Mo–O) is shorter than that of (M–O) because Mo$^{6+}$ (0.59 Å) and Mo$^{5+}$ (0.61 Å) are both smaller than the ionic radii of the M cations. Reduction from 180° of the M–O–Mo bond angles is caused by a cooperative rotation of the MO$_6$/2 and MoO$_6$/2 octahedra; these rotations increase as the geometric tolerance factor $r = (r_A + r_O)/\sqrt{2(r_B + r_O)}$ decreases. The $r_A$, $r_B$, and $r_O$ are respectively the room-temperature ionic radii of the A-site cation (ninefold coordination), the mean B-site ionic radius (sixfold coordination), and the oxide-ion radius (twofold coordination) with the Shannon$^{13}$ ionic radii. The $r$ factors are 0.932 and 0.945 for M = Co and M = Ni, respectively. This assignment for the valence states places the Mo$^{6+}$/Mo$^{5+}$ reduction potential above the M$^{3+}$/M$^{2+}$ reduction potentials of M = Co and Ni.

The double-perovskite B-site cations generally exhibit some antistructure disorder and antiphase boundaries. The value of the order parameter $ξ$ can be calculated as $ξ = 2(\delta m –


of the end point. The whole reaction can be expressed as
solution of sodium thiosulfate; starch was used as an indicator
situ was immediately titrated with a standard volumetric
then excess KI was added. The purple iodine produced in
powder was dissolved in diluted HCl aqueous solution and
vacancies exist in the reduced Sr2MgMoO6.5,6 Bernuy-Lopez
6.7% and 4.2%, respectively.

The iodometric titration method can be used to determine
directly the valence of some cations in the double perovs-
kites.14 According to their standard oxidation—reduction
potential, Co3+ and Ni3+ can efficiently oxidize I− to I2.
Therefore, we used this method to check the contents of Co5+
and Ni3+ ions. A weighed amount of double-perovskite
powder was dissolved in diluted HCl aqueous solution and
then excess KI was added. The purple iodine produced in
situation was immediately titrated with a standard volumetric
solution of sodium thiosulfate; starch was used as an indicator
of the end point. The whole reaction can be expressed as
follows:

\[
2M^{3+} + 2I^- = I_2 + 2M^{2+} \quad (M = \text{Co, Ni}) \tag{1}
\]

\[
2S_2O_3^{2−} + I_2 = 2I^- + S_4O_6^{2−} \tag{2}
\]

The percentages of Co5+/Co and Ni5+/Ni in the as-prepared
Sr2CoMoO6 and Sr2NiMoO6 samples sintered in air were
6.7% and 4.2%, respectively.

We have shown in our previous work that some oxygen
vacancies exist in the reduced Sr2MgMoO₆.5,6 Bernuy-Lopez
and Marrero-López et al.15,16 separately noted that reduction
of Sr₂MgMoO₆ can give rise to a limited number of oxygen
vacancies and hence to some lower Mo oxidation states than
Mo⁶⁺. Here, we focus on the temperature dependence of the
oxygen vacancies in the as-prepared Sr₂CoMoO₆ and
Sr₂NiMoO₆ samples in air and a 5% H₂/Ar atmosphere.

The perovskite structure does not accept interstitial oxygen.
Therefore, the presence of M³⁺ species in the air-sintered
samples requires the presence of some cation vacancies.
Partial evaporation of some of the constituent metals during
sintering at 1250 °C in air led to a cationic nonstoichiometry in
the as-prepared samples.17,18 We investigate how oxygen is
lost and reincorporated on raising and lowering of the
temperature with TGA. Since the Mo⁶⁺/Mo⁵⁺ redox couple is
at a higher energy than the M⁴⁺/M³⁺ couple, reduction of
the samples will, first, reduce the M³⁺ to M²⁺. (Note:
throughout we indicate formal valences on the cations, not
the actual charges they carry.)

On heating in air, Figure 1A, both samples show the onset
of weight loss at 300 °C and a step loss at 650 °C,
corresponding to ca. 0.05 O atoms/f.u., that is shown to be
irreversible in the case of M = Co. The M = Co sample
also shows a second reversible weight-loss step at 900 °C;
the M = Ni sample does not. This difference and the smaller
oxidation of Ni²⁺ versus Co²⁺ reflects, we believe, a greater
resistance to oxygen and Mo loss by the stronger preference

of Ni²⁺ for octahedral sixfold oxygen coordination. The step
weight losses signal oxygen loss in a finite volume fraction,
i.e., at defects. Oxygen vacancies trapped by formation of
molybdyl (Mo⁵⁺O) species at Mo-rich antiphase boundaries,
for example, would provide an irreversible, finite oxygen
loss. We postulate antisite Mo or Mo-rich antiphase bound-
aries are the operative defect since these would be common
to both samples. The second step in the M = Co sample at
900 °C reflects, according to this reasoning, reversible
trapping of an oxygen vacancy at some other defect associ-
ated with the M²⁺ ion since it does not occur in the M = Ni
sample. Given the presence of Mo vacancies, we tentatively
assign this step to trapping of oxygen vacancies at Co
neighboring a Mo vacancy. A Co²⁺ ion at a Mo vacancy is
stable in a lower oxygen coordination whereas the Ni²⁺ ion
is much less stable in lower coordination.

Next, the TGA curves of Figure 1B were recorded on
heating the as-prepared M = Co and M = Ni samples under
a 5% H₂/Ar atmosphere, which resembles that at the anode
of an operating SOFC. In this atmosphere, both samples
begin to decompose above 800 °C where we have confirmed
initiation of the reduction of MoO₃ with TGA measurements
on MoO₃ in the same atmosphere (not shown), which
contains Mo⁵⁺O units. In 5% H₂/Ar, the step found at 650
°C in air does not appear with either sample. On the other
hand, a significant reversible weight loss occurs in a step
above 300 °C in the M = Co sample, but not in the M = Ni
sample. Figure 2 shows that this weight loss in the M = Co
sample is due to a reversible loss of oxygen. On heating in

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\(18\) Ivanov, S. A.; Eriksson, S. G.; Tellgren, R.; Rundlöf, H.; Tseggai,
air, a nearly complete regain of the weight lost occurs below 650 °C, and the weight-loss steps at 650 and 950 °C reappear. We therefore conclude that the small, irreversible weight loss in air above 350 °C in both samples, Figure 1, reflects reduction of M$^{3+}$ to M$^{2+}$; it occurs more abruptly in the M = Ni sample because of the stronger octahedral-site preference of the Ni$^{3+}$ ion. In the 5% H$_2$/Ar atmosphere, the M$^{3+}$ ions would be reduced to M$^{2+}$ already at room temperature and the vacancies introduced by this reduction are apparently trapped at the defects postulated to be Mo-rich antiphase boundaries. In the M = Co sample, the weight loss above 300 °C in 5% H$_2$/Ar would then represent a reduction of Mo$^{6+}$ to Mo$^{5+}$ by a reversible loss of oxygen that introduces mobile bulk oxygen vacancies. The stronger octahedral-site preference of Ni$^{3+}$ inhibits further loss of oxygen in the 5% H$_2$/Ar atmosphere once the Ni$^{3+}$ have been reduced to Ni$^{2+}$ and the oxygen vacancies so introduced have been trapped. Heating the M = Co sample in air after reduction in 5% H$_2$/Ar reintroduces oxide ions that move, below 650 °C, to annihilate the trapped vacancies, vacancies that are again created on further heating above 650 °C.

In ref 19, a TGA curve was presented for H$_2$-reduced Sr$_2$CoMoO$_6$ to demonstrate its reoxygenation in air up to 800 °C. Even though the authors do not pay attention to the slight lowering of the curve after it reaches its maximum around 650 °C, it is clear that their TGA curve is highly consistent with the present data and conclusions.

The Mo L$_3$-edge, Ni K-edge, and Co K-edge XAS measurements were performed to confirm the cation valences. Since a simultaneous standard was not possible at the low energies of the Mo L$_3$ edges, possible standards were run periodicaly in the sequence of samples and the relative energy was ±0.1 eV or better. In view of the low energy and strong “white line” p-to-d transition at the Mo L$_3$ edge self-absorption, degradation/rounding of the absorption edge peaks in the fluorescence mode was unavoidable. Here the “white line” (WL) terminology is conventional and refers to an atomic-like dipole transition into empty states that typically manifests a sharply peaked near-edge structure; in prior years it left a white-line streak on photographic film. As is routinely done, the spectra presented in this study had a linear background subtracted (determined over a ca. 80 eV interval below the edge) and were normalized to unity absorption step height across the edge. Here an average of the data in the 50–200 eV range above the edge was used to set the normalization value.

The L$_3$ near-edge WL features of 4d transition-metal compounds are due to transitions from the 2p to 4d states of the transition metal (such as Mo). In an octahedral ligand field, the d states of Mo$^{6+}$ are split into a sixfold-degenerate $t_{2g}$ ground state below a fourfold-degenerate $e_g$ state. In Figure 3a the L$_3$-edge WL features of the octahedrally coordinated compounds Sr$_2$CoMoO$_6$ and Sr$_2$NiMoO$_6$ are shown in comparison with the spectra of another octahedrally coordinated compound, MoO$_3$. The A and B features are associated, respectively, with the $t_{2g}$ and $e_g$ final states of the octahedrally coordinated compounds. It should also be noted that the relative intensity of the A feature decreases as the number of empty $t_{2g}$ states decreases (i.e., as the


electron count increases or valence decreases). Thus, the relative intensity ratio of the A to B features, \( I_A/I_B \), can be used to track the d-electron/hole count in such Mo compounds; i.e., a higher \( I_A/I_B \) usually corresponds to a higher Mo valence.

The double-peak features A and B in Figure 3A are seen in all spectra and can be attributed to the \( 2p \rightarrow 4d(t_{2g}) \) and \( 2p \rightarrow 4d(e_g) \) transitions, respectively. Since such double-peak features could have different widths for different compounds, it is more reliable to use the centrum or mean energy \( E_m \) (or characteristic energy\(^{22,23} \)) instead of the edge energy (defined as the inflection point of the rising part of the edge) to estimate the Mo valence. The values of \( E_m \) estimated for these three spectra are almost identical (2525.5 eV), supporting the belief that the values of the Mo valence are nearly the same for these three compounds. The very close Mo valence in these three compounds indicates that Ni and Co are predominantly in the divalent state like that for Mg. To confirm this deduction, we show the Co and Ni K-edges of the Sr\(_2\)MMoO\(_6\) compounds with M = Co and Ni, respectively, in Figure 3B,C, along with selected spectra of other compounds (see circles) standard.

\[ \chi(\text{emu/mol} \cdot \text{Oe}) \]

\[ T = 80 \text{ K} \]

\[ T = 37 \text{ K} \]

\[ \chi(\text{emu/mol} \cdot \text{Oe}) \]

\[ \chi^{-1}(\text{mOe/emu}) \]

\[ \chi^{-1}(\text{mOe/emu}) \]

\[ \text{M} = \text{Ni} \]

\[ \text{M} = \text{Co} \]

\[ \chi^{-1}(\text{emu/mol} \cdot \text{Oe}) \]

\[ \chi^{-1}(\text{mOe/emu}) \]

\[ \text{Sr}_2\text{MMoO}_6 \text{ with M = Co and Ni. They both show a long-range-ordered antiferromagnetic state at low temperature. The Neél temperature } T_N \text{ values are 37 and 80 K for M = Co and Ni, respectively. In the double-perovskite structure, the magnetically active M\(^{2+}\) ions are separated by magnetically neutral Mo\(^{6+}\) ions. The exchange interactions propagate through the intervening MoO\(_6\) octahedra; therefore, a relatively high level of covalent bonding is expected within the M–O–Mo–O–M pathways. The variation of the } T_N \text{ values depends primarily on the inverse of the energy separating the M\(^{3+}/\text{M}^{2+}\) and M\(^{2+}/\text{M}^{1+}\) redox energies; the correlation energy U for Co\(^{2+}\) is larger than the charge-transfer gap } \Delta \text{ for Ni. Interestingly, with reduction of the air-sintered } \text{Sr}_2\text{CoMoO}_6 \text{ sample in 5% H}_2/\text{Ar or H}_2 \text{ at 800 °C for 10 h, the reduced sample exhibits a spin-glass ferromagnetism in the measured temperature range from 5 to 380 K. A similar phenomenon was also observed by Viola et al.\(^{25} \) in oxygen-deficient } \text{Sr}_2\text{CoMoO}_6-\delta \text{ with a Curie temperature } T_C \text{ = 350–370 K. The observed spin-glass ferromagnetism in } \text{Sr}_2\text{CoMoO}_6-\delta \text{ comes from a ferromagnetic superexchange between Co\(^{2+}\) (3d\(^3\)) and Mo\(^{5+}\) (4d\(^1\)) moments in the cation-ordered regions with antiferromagnetic M\(^{2+}\)-O–M\(^{2+}\) interactions across antiphase boundaries and at antisite M\(^{2+}\). A good fitting with the Curie–Weiss law is observed above } T_N \text{ in the } \chi^{-1}–T \text{ curves for } M = \text{Co and Ni. The linear fitting above 200 K gives an effective paramagnetic moment } (\mu_{\text{eff}}) \text{ of 4.34(2) and 3.51(1} \mu_{\text{B}}/\text{f.u.} \text{ for } M = \text{Co and Ni, respectively. For } \text{Sr}_2\text{CoMoO}_6-\delta \text{, the measured } \mu_{\text{eff}} \text{ is higher than the spin-only 3.87} \mu_{\text{B}} \text{ for Co\(^{2+}\) ions. The measured } \mu_{\text{eff}} \text{ for Sr\(_2\)NiMoO\(_6\) is also high relative to the predicted spin-only 2.83} \mu_{\text{B}} \text{ for Ni\(^{2+}\) ions. Interpretation of the discrepancies between measured and spin-only } \mu_{\text{eff}} \text{ values is complicated by the existence of antiphase boundaries where short-range magnetic order may occur above the } T_N \text{ for the ordered regions.}

Conductivity data taken above 350 °C while heating in different atmospheres are shown in Figures 5 and 6 for nominal Sr\(_2\)CoMoO\(_6-\delta\) and Sr\(_2\)NiMoO\(_6-\delta\), respectively. The conductivity above 350 °C is dominated by electrons on the Mo\(^{6+}/\text{Mo}^{5+}\) couple given the much lower mobility of the oxygen vacancies. The as-prepared M = Co sample gave positive thermoelectric power at room temperature showing that the conduction was by holes in the Co\(^{3+}/\text{Co}^{2+}\) couple (see Figure S4 in the Supporting Information), but the Co\(^{3+}\) are all reduced to Co\(^{2+}\) above 350 °C. In air, the log(\(\sigma(T)\)) vs \( T^{-1} \) curves for M = Co, Figure 5B, are linear below 650 °C, but the curve deviates to higher conductivity above 650 °C where the TGA curves of Figure 1A show a step. Electrons would be on Mo\(^{5+}\) coordinated by an oxygen vacancy, but trapping of oxygen vacancies at Mo\(^{6+}\) would release electrons to the matrix. The M = Ni sample in air had a larger resistivity with a nonlinear Arrhenius behavior below 650 °C, consistent with a small step in the TGA curve near 500 K and a saturation of the number of charge carriers.


After reduction in 5% H₂/Ar for 20 h at 800 °C, the resistivity of the M₀₀₀ sample in that atmosphere shows a broad maximum near 650 °C where the TGA curve in air shows a step that we have postulated to be associated with formation of Mo₃O species that trap oxygen vacancies. Since the system was carefully sealed so as to prevent leakage from air, the partial pressure of oxygen in the gas was extremely low (pO₂ = 10⁻¹⁹ atm). Mo⁵⁺ ions would be trapped at oxygen vacancies unless formation of Mo₃O species released the electrons. Nevertheless, at least half of the electrons introduced by the oxygen vacancies would be free to reduce the Co³⁺ to Co²⁺ or to move on the Mo in sixfold oxygen coordination. Although a small fraction of metallic cobalt or nickel was found in the XRD data of the samples after reduction in 5% H₂/Ar at 800 °C for 20 h (see Figure 7), this fraction was too small to percolate through the sample. Therefore, the grain conductivity can be assumed to be dominated by transfer of electrons from Mo⁵⁺ to Mo⁶⁺ octahedral sites. This assumption is supported by the increase in conductivity above 600 °C, which could be due to excitation of electrons from Co²⁺ to Mo⁶⁺ or, more likely, by release above 600 °C of electrons trapped at fivefold-coordinated Mo=O. The increase in resistivity with temperature below 600 °C is more problematic; but the motional enthalpy of polaronic conduction may increase with temperature or the grain-boundary contribution to the conductivity may be dominant at lower temperatures. An analogous increase in the conductivity of the M₀₀₀ sample above 600 °C is also apparent.

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The double-perovskite samples were tested as anode materials in SOFCs having a 300 µm thick LSGM electrolyte and SrFe₀.₂Co₀.₈O₆₋δ cathode. Figure 8 shows SEM images for the single cell with Sr₂CoMoO₆ (SCMO) as the anode after operating in H₂ and CH₄. A clear cross section of the cell configuration can be seen in Figure 8A. Estimated from the SEM images, the thicknesses of the LSGM, SCMO, LDC, and SCF layers are 310, 26, 8, and 20 µm, respectively. The LDC buffer layer obviously exists between LSGM and...
SCMO, which efficiently prevents a diffusion reaction between the anode and the electrolyte. Both anode and cathode layers are porous, which favors transfer of oxygen, fuel gas, and exhaust products.

Power density and cell voltage as functions of current density at 800 °C in H₂, dry CH₄, and wet CH₄ for (A) Sr₂CoMoO₆ and (B) Sr₂NiMoO₆.

![Figure 9](image_url)

We used power cycles to test the stability of the anodes in different fuels. Figure 10 shows the maximum power density as a function of power cycle for various anodes Sr₂MMoO₆ (M = Co and Ni) at 800 °C in H₂, dry CH₄, and wet CH₄.

![Figure 10](image_url)

The higher power output with H₂ fuel for the M = Co sample is consistent with its higher concentration of oxygen vacancies and higher electronic conductivity, which we believe to be the result of the stronger octahedral-site preference of the Ni²⁺ ions. From Figure 10, we can see that both Sr₂CoMoO₆ and Sr₂NiMoO₆ anodes run stably over power cycling in H₂, which can be ascribed to their stable phases in H₂. The XRD pattern in Figure 7 has typically shown that Sr₂CoMoO₆ is almost pure after being reduced at 800 °C in H₂ for 20 h.

The performances are quite different for Sr₂CoMoO₆ and Sr₂NiMoO₆ in dry and wet methane. Sr₂NiMoO₆ shows a higher power in dry CH₄, demonstrating that M = Ni favors the main process for direct oxidation of methane. The reaction can be expressed as

\[
\text{CH}_4 + 4\text{O}^{2-} = \text{CO}_2 + 2\text{H}_2\text{O} + 8\text{e}^- \quad (3)
\]

The O²⁻ ions are created in the cathode and transported through the electrolyte from the cathode. Sr₂CoMoO₆ exhibits a much higher power in wet CH₄, indicating a preferential reaction pathway on the anode through steam reforming of methane:

\[
\text{CH}_4 + \text{H}_2\text{O} = \text{CO} + 3\text{H}_2 \quad (4)
\]

We should take into account the catalytic effect of Sr₂CoMoO₆ on the above reformer reaction. Metallic Co was observed in Sr₂CoMoO₆ exposed to CH₄ (see the XRD patterns in Figure 7). Moreover, Co has been found to have high catalytic activity for the reforming of methane. Therefore, it is reasonable to propose that the high performance for Sr₂CoMoO₆ in wet CH₄ fuel is due to the catalytic effect of surface Co⁰ on the reformer reaction. Metallic Ni can also act as a catalyst for steam reforming of CH₄, but the power density of Sr₂NiMoO₆ is lower than that of Sr₂CoMoO₆ in wet CH₄. We note that Ni easily induces formation of graphitic carbon on the catalyst, which causes

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catalytic deactivation and blockage of the reactor. In addition, after exposure of the sample to CH4, a tiny amount of SrCO3 and SrMoO4 also became visible in the XRD data (Figure 7). SrCO3 and SrMoO4, which would be on the surface of the anodes, may block the catalytic pathways for reforming of the fuel. These impurities would increase with operation time and may be the reason why the power density in CH4 drops rapidly over cycling in Figure 10.

Conclusions

Magnetic, structural, and XAS data show the existence of Co2+/Mo6+ and Ni2+/Mo6+ pairs in the double-perovskites Sr2MMoO6. The Mo6+/Mo5+ reduction potential is low enough to accept electrons from H2 and CH4, thus allowing dissociative chemisorption of the fuel. Moreover, displacement of a Mo6+ or a Mo5+ ion within an octahedral site to form a stable Mo=O species would trap oxygen vacancies in Mo-rich antiphase boundaries and release electrons trapped at oxygen vacancies; also, Mo=O species may favor reaction of the fuel with a surface oxide ion by lowering its acidity and energy for desorption of the oxidized reaction product. Polaronic conduction on the mixed-valent Mo6+/Mo5+ array allows transfer of the electrons from the surface reaction to the current collector, and bulk oxygen vacancies allow transfer of O2− ions from the oxide-ion electrolyte to the anode surface to replenish the surface oxygen that is lost. In methane, the rate-limiting processes are the chemical reactions occurring at the surface of the anode, but the dominant processes are obscured by the evolution onto the anode surface of Co or Ni and SrCO3 with SrMoO4 in the reducing atmosphere at the anode.

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Supporting Information Available: Rietveld refinement of XRD patterns, magnetic susceptibility, thermoelectric power, and the average activation energy of electronic conduction for the anode materials. This material is available free of charge via the Internet at http://pubs.acs.org.

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