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Published in:
International Journal of Electrochemistry

Link to article, DOI:
10.1155/2020/4013697

Publication date:
2020

Document Version
Publisher's PDF, also known as Version of record

Link back to DTU Orbit

Citation (APA):

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Electrochemical Reduction of Oxygen and Nitric Oxide on Mn-Based Perovskites with Different A-Site Cations

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Received 31 December 2019; Accepted 10 March 2020; Published 29 March 2020

Academic Editor: Adalgisa Rodrigues de Andrade

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Four LnMnO$_{3+\delta}$ (Ln = La, Pr, Sm, and Gd) perovskites were synthesized and characterized by powder XRD. It was shown that the perovskite lattice became more and more distorted when lowering the size of the A-site cation. The manganite-based perovskites were evaluated for the ability to electrochemically reduce oxygen and nitric oxide in the temperature range of 200 to 400 °C. At the lowest temperature, the electrodes were better at reducing nitric oxide than oxygen. At higher temperatures, the activity for the reduction of oxygen and nitric oxide became similar. The activation energies for the reduction of oxygen and nitric oxide were markedly different for LaMnO$_{3+\delta}$ and PrMnO$_{3+\delta}$ whereas it was similar for SmMnO$_{3+\delta}$ and GdMnO$_{3+\delta}$.

1. Introduction

Pollution from mobile and stationary sources is of still greater concern [1]. While the emissions of NOx from petrol-fired engines can be significantly lowered using the three-way catalytic converter [2], problems still remain for the cleanup of the exhaust from diesel-fired engines, due to excess oxygen in the exhaust gas [3]. The best known solution for the removal of nitrogen oxides from diesel engines is the selective catalytic reduction (SCR) with ammonia [4]. This solution requires, however, storage of the reducing agent on board the vehicle. There also exists the possibility of slip of the reducing agent.

An alternative solution to the SCR route is the reduction of the NOx in an all-solid-state electrochemical reactor, based on an oxide anion conductor, as suggested by Pancharatnam et al. [5]. In such a reactor, the NOx is reduced at the cathode to nitrogen and oxide anions. The oxide anions are then transported through the electrolyte to the anode where oxygen is formed. Obstacles still remaining for this solution are a low activity at the temperatures of the exhaust gas and a low selectivity due to simultaneous reduction of oxygen at the cathode, leading to high power consumption (see, e.g., [6] for a recent review).

Manganite-based perovskites have been investigated as cathodes for the electrochemical reduction of nitric oxide [7–9]. It has been shown that the manganite-based perovskites are active towards the reduction of nitric oxide even at the low temperature of a diesel exhaust gas.

In this study, manganite-based perovskites with different A-site cations are for the first time investigated for the reduction of oxygen and nitric oxide at the temperature interval of a real diesel exhaust gas. Manganite perovskite-based cathodes, with different A-site cations, have been studied before for the use as oxygen reduction cathodes [10, 11]. In some cases, the effect is quite large. Especially, it has been tried to use different A-site cations in Ln$_{0.6}$Sr$_{0.4}$MnO$_{3+\delta}$ and it was shown that the Pr-containing perovskite had the highest activity towards the reduction of oxygen at a temperature of 700°C [11]. No studies for the electrochemical reduction of nitric oxide on perovskites with different A-site cations have been reported in the open literature. The study is done on cone-shaped electrodes as suggested by Fabry and Kleitz [12]. The advantage of the cone-shaped electrodes is that the contact area is well defined and that the processing of the electrolyte and electrode is done separately, thereby avoiding reaction between the electrode and the electrolyte during sintering.
2. Experimental

Synthesis of the manganite-based perovskites was done using the glycine-nitrate process [13]. In short solutions of the metal nitrates were mixed in a beaker in the appropriate ratio. Glycine was then added, and the solutions were heated on a hot plate until ignition. Then, the resulting powders were calcined in a box furnace in air at 1100°C for 12 h. Phase purity of the as-synthesized powders was checked using powder X-ray diffraction. After this, pellets of the powders were made by pressing the powders in an appropriate die followed by sintering. Finally, diamond tools were used to shape the as-sintered pellets into cone-shaped electrodes. XRD was performed on the sintered pellets in order to be certain that the perovskite structure still was present after sintering at high temperatures.

The cone-shaped electrodes were tested in a pseudo-3-electrode setup [14], with a Pt reference/counter electrode and a one end closed yttria stabilized zirconia tube (Vesuvius) (see Figure 1). The end, of the zirconia tube, exposed to the cone-shaped electrodes was polished down to 1 μm with diamond paste. Approximately 60 g of weight was applied to the cone-shaped electrodes. The electrochemical measurements were done using a Gamry Femtotstat potentiostat. The voltammograms were recorded between 0.4 V and -0.8 V vs. air, starting at open circuit voltage (OCV) and sweeping in the cathodic direction first. The voltammograms were recorded with sweep rates 1 or 10 mV/s. Electrochemical impedance spectroscopy (EIS) was also recorded at OCV. The EIS was recorded from 300 kHz to 50 mHz, with 10 points per decade. An amplitude of 25 mV rms was used. The EIS was used to find the contact area of the cone-shaped electrodes using Newman’s formula [15]. The electrochemical behavior of the manganite-based perovskites was tested in atmospheres containing either 10% oxygen in argon or 1% nitric oxide in argon. Brooks mass flow controllers were used to control the flow rate and gas composition. The flow rate was kept at 20 mL/min throughout. The electrochemical measurements were done at 200, 300, and 400°C.

3. Results

Results from the powder XRD diffraction show that the as-sintered perovskites are of single phase. The unit cell parameters can be found in Table 1. The lanthanum-containing manganite has a hexagonal structure, whereas the other manganites have an orthorhombic structure. The manganites become more and more distorted from the ideal cubic perovskite structure when lowering the ionic radii of the A-site cation.

Examples of voltammograms are given in Figures 2-7 (the results for the LaMnO₃⁺δ is taken from [9], but is plotted here again for clarity). In Figure 2, the results of the measurements in the oxygen-containing atmosphere at 200°C are shown. It is seen that the LaMnO₃⁺δ and the PrMnO₃⁺δ perovskites have an equal activity and that the SmMnO₃⁺δ and GdMnO₃⁺δ perovskites also have an equal activity, which is markedly higher than that for the LaMnO₃⁺δ and PrMnO₃⁺δ perovskites. Again at 300°C (see Figure 3), the Sm- and Gd-containing perovskites have an almost identical activity. At this temperature, the La-containing perovskites have a much higher activity than the other perovskites, and the PrMnO₃⁺δ perovskite has a much lower activity. At 400°C (Figure 4), the PrMnO₃⁺δ perovskite again performs the worst. At potentials between 0 and -0.7 V vs. air, the LaMnO₃⁺δ still performs the best, when it comes to the reduction of oxygen. However, at low potentials, the downward bend of the voltammograms is stronger on the Sm- and Gd-based manganites than on the LaMnO₃⁺δ perovskite, making them more active towards oxygen reduction at the very lowest applied potentials. The manganite-based perovskites are also able to oxidize the oxide ions to oxygen. The PrMnO₃⁺δ perovskite is the most active in this sense.

In the nitric oxide-containing atmosphere, a markedly different behavior of the perovskites is observed (see Figures 5–7). At the lowest temperature (200°C), the lanthanum-based manganite has a much higher activity than the other perovskites (see Figure 5). The magnitude of the current densities is markedly higher in the nitric oxide-containing atmosphere than in the oxygen-containing atmosphere, at this temperature. As it is also observed in the oxygen-containing atmosphere, the differences between the

![Figure 1: A sketch of a setup for measurements on cone-shaped electrodes.](image-url)
activities of the perovskites level out at the higher temperatures (see Figures 6 and 7). At 400°C, the strong downward bend in the voltammograms for the SmMnO$_{3+\delta}$ and GdMnO$_{3+\delta}$ perovskites, also observed in the oxygen-containing atmosphere, at low potentials is seen again. P'hemaximum cathodic current density obtained is given in Table 2. It is observed that the apparent selectivity is highest for the La compound.

The activation energies are given in Table 3. The activation energy is highest for the LaMnO$_{3+\delta}$ perovskite in the oxygen-containing atmosphere, 0.87 eV. This compound also has the lowest activation energy, 0.37 eV, in the nitric oxide-containing atmosphere. For the SmMnO$_{3+\delta}$ and GdMnO$_{3+\delta}$ perovskites, the activation energy is almost the same, around 0.65 eV, in the oxygen- and nitric oxide-containing atmospheres.
Figure 6: Voltammograms recorded at 300°C in 1% NO in Ar on the four manganite-based perovskites with different A-site cations. The voltammograms are recorded with a sweep rate of 1 mV/s. Also at this temperature, the LaMnO$_{3+\delta}$ perovskite is the most active. The differences between the three other manganite-based perovskites are smaller at this temperature.

Table 2: Maximum cathodic current densities in oxygen- and nitric oxide-containing atmospheres for the four manganite-based perovskites, given in mA/cm$^2$. The current densities increase with increasing temperature. The apparent selectivity for the reduction of nitric oxide is highest for the LaMnO$_{3+\delta}$ perovskite at 200°C.

<table>
<thead>
<tr>
<th>Compound</th>
<th>LaMnO$_{3+\delta}$</th>
<th>PrMnO$_{3+\delta}$</th>
<th>SmMnO$_{3+\delta}$</th>
<th>GdMnO$_{3+\delta}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>T/gas</td>
<td>O$_2$</td>
<td>NO</td>
<td>O$_2$</td>
<td>NO</td>
</tr>
<tr>
<td>200°C</td>
<td>0.01</td>
<td>0.23</td>
<td>0.01</td>
<td>0.12</td>
</tr>
<tr>
<td>300°C</td>
<td>2.60</td>
<td>2.00</td>
<td>0.20</td>
<td>1.20</td>
</tr>
<tr>
<td>400°C</td>
<td>4.60</td>
<td>3.20</td>
<td>1.00</td>
<td>3.00</td>
</tr>
</tbody>
</table>

Table 3: The activation energies for the reduction of oxygen and nitric oxide in the temperature interval 200 to 400°C given in eV. The activation energy is highest for the end member LaMnO$_{3+\delta}$ in the oxygen-containing atmosphere. The activation energies in the oxygen-containing atmospheres are almost the same for the three other manganite-based perovskites.

<table>
<thead>
<tr>
<th>Compound</th>
<th>LaMnO$_{3+\delta}$</th>
<th>PrMnO$_{3+\delta}$</th>
<th>SmMnO$_{3+\delta}$</th>
<th>GdMnO$_{3+\delta}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gas</td>
<td>O$_2$</td>
<td>NO</td>
<td>O$_2$</td>
<td>NO</td>
</tr>
<tr>
<td>$E_A$</td>
<td>0.87</td>
<td>0.37</td>
<td>0.64</td>
<td>0.45</td>
</tr>
</tbody>
</table>
containing atmospheres. The activation energy in the oxygen-containing atmosphere for the PrMnO$_{3+δ}$ perovskite is similar to the activation energies of the SmMnO$_{3+δ}$ and GdMnO$_{3+δ}$ perovskites. However, the activation energy for the PrMnO$_{3+δ}$ perovskite, in the nitric oxide-containing atmosphere, is markedly lower than the activation energies for the SmMnO$_{3+δ}$ and GdMnO$_{3+δ}$ perovskites. The differences in activation energies make the LaMnO$_{3+δ}$ and PrMnO$_{3+δ}$ perovskites the ones with the highest apparent selectivity towards the reduction of nitric oxide at the lowest temperatures.

4. Discussion

The crystal structures obtained for the four different perovskites with different A-site cations are in accordance with the literature [16–18]. The distortion of the ideal cubic perovskite structure, seen in this study, will normally lead to a lowering of the electronic conductivity, when the size of the A-site cation is lowered. This is due to less overlap between the p-orbitals from the oxygen and the d-orbitals from the B-site cation, the so-called superexchange. However, probably the effect is so small and has no practical importance for the use of the manganese-based perovskites as electrodes.

In the following tentative explanations, the differences in the electrochemical behavior of the four perovskites will be given. For the reduction of oxygen, the PrMnO$_{3+δ}$ perovskite always shows the lowest activity, in the investigated temperature range. This is in contradiction with the literature [11], but it should be kept in mind that this study has been undertaken at much lower temperatures than that in the literature and that the manganites investigated in the literature were strontium doped. The reason for the low activity of the Pr-containing manganite could be because the Pr can exist also in the oxidation state Pr(IV), leading to a higher concentration of cation vacancies in the perovskite structure. The behavior of the LaMnO$_{3+δ}$ perovskites is very different than that of the other perovskites. This could be due to that the mobility of oxide ions is more thermally activated on the LaMnO$_{3+δ}$ perovskite than on the other perovskites. That the voltammograms for the SmMnO$_{3+δ}$ and GdMnO$_{3+δ}$ perovskites bend more strongly downwards at low potentials could be because these two perovskites are easier to reduce than the LaMnO$_{3+δ}$ and PrMnO$_{3+δ}$ perovskites (see [19]), leading to the creation of oxide ion vacancies in perovskite structure at low oxygen partial pressures (low potentials). This can be explained on the background of the inductive effect. When the size of the A-site cation is lowered, the A-site cation will attract more electron density from oxygen. This will again make oxygen attracting more electron density from the B-site cation, lowering the electron density on the B-site cation. This will make it easier to reduce the B-site cation, as observed in this study.

The differences of the manganite-based perovskite cathodes are larger at the lowest temperature also in the nitric oxide-containing atmosphere, and the differences level out at the highest temperature (see Table 2). The nitric oxide is perhaps bound more strongly on the LaMnO$_{3+δ}$ and PrMnO$_{3+δ}$ perovskites at the lowest temperatures. That the activity increases less with increasing temperature for the LaMnO$_{3+δ}$ and PrMnO$_{3+δ}$ perovskites could also be due to weak chemisorption of nitric oxide on these compounds at elevated temperature. It has been shown in the literature that oxygen is bound less strongly on these compounds than on GdMnO$_{3+δ}$ [19]. At 400°C, the same trends as in the oxygen-containing atmosphere are observed, that is, the reduction of the perovskites at low potentials is easier when the size of the A-site cation is lowered.

For the oxygen evolution, the PrMnO$_{3+δ}$ perovskite shows superior activity at 400°C. Pr-containing compounds have before been shown to be good catalysts for oxidation processes [20]. The activity of the PrMnO$_{3+δ}$ perovskite for oxidizing nitric oxide to nitrogen dioxide is also highest at 400°C, even though the effect is much less pronounced than in the oxygen-containing atmosphere.

That the activation energies are almost the same on PrMnO$_{3+δ}$, SmMnO$_{3+δ}$ and GdMnO$_{3+δ}$ perovskites in the oxygen-containing atmosphere indicates that the reaction mechanism is the same for these three manganite-based perovskites. However, for the LaMnO$_{3+δ}$ perovskite, the higher activation energy suggests that the reduction of oxygen takes place through a different mechanism than on the three other perovskites. In the nitric oxide-containing atmosphere, the activation energies suggest that the reduction of nitric oxide proceeds through different mechanisms for the LaMnO$_{3+δ}$ and PrMnO$_{3+δ}$ perovskites and for the SmMnO$_{3+δ}$ and GdMnO$_{3+δ}$ perovskites.

5. Conclusion

Four manganite-based perovskites, LaMnO$_{3+δ}$, PrMnO$_{3+δ}$, SmMnO$_{3+δ}$, and GdMnO$_{3+δ}$ perovskites, were synthesized and shown to be single phased by powder XRD. The manganites were shown to be better at reducing nitric oxide than oxygen at the lowest temperature. At higher temperatures, the differences leveled out. The work clearly revealed that the choice of the A-site cation is important when designing electrodes for electrochemical reduction of oxygen or nitric oxide at low temperatures. The differences in the activity of then manganites were tentatively explained on the background of the inductive effect, chemisorption, and the oxide ion mobility in the crystal lattice.

Data Availability

All data will be available on request.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

References


