Electrochemistry of Nanocrystalline La$_{0.5}$Sr$_{0.5}$MnO$_3$ Perovskite for the Oxygen Reduction Reaction in Alkaline Medium

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Abstract Nanocrystalline La$_{0.5}$Sr$_{0.5}$MnO$_3$ (LSMO) perovskite-sized powders were successfully synthesized at heating low temperature and time using the Pechini method based on polyesterification between citric acid and ethylene glycol. The electroactivity of carbon-supported perovskite (LSMO/C) for the oxygen reduction reaction (ORR) was evaluated. The results showed that the electroactivity of LSMO/C largely depends on the calcination temperature, type of electrolyte, and mass loading. It increased remarkably with the calcination temperature in 0.1 M KOH. The ORR performance was examined in KOH, NaOH, and K$_2$SO$_4$. Higher electroactivity was recorded in KOH electrolyte, as compared to NaOH and K$_2$SO$_4$.

Keywords Nanoparticles · Pechini method · Perovskite · Oxygen reduction · Alkaline medium

Introduction

The oxygen reduction reaction (ORR) has been studied at length due to its great importance in energy conversion and storage devices and extended to alkaline fuel cells [1], electrolyzers [2], and metal-air batteries [3]. In alkaline media, the ORR proceeds as in the acid counter-part, either via two-electron $\text{O}_2 + \text{H}_2\text{O} + 2\text{e}^- \rightarrow \text{HO}_2^- + \text{OH}^-$ or four-electron $\text{O}_2 + 2\text{H}_2\text{O} + 4\text{e}^- \rightarrow 4\text{OH}^-$ pathways. Many researchers are doing their best to find suitable electrocatalysts that could support the desired four-electron reduction process leading to an efficient energy conversion [4]. Although noble metals are suitable ones and have proved to be effective oxygen electrodes, the high costs and scarcity prevent their large-scale application. In order to reduce these costs, alternative non-precious metal oxides with comparable ORR activity to Pt-based catalysts are highly desirable for the development of fuel cells. In this regard, oxides with perovskite structure (ABO$_3$) have received wide attention due to their high activity and stability when used in alkaline media [5]. For instance, it has been reported that LaMnO$_3$ showed the best catalytic activity for both ORR and oxygen evolution reaction (OER) in 6 M KOH, whereas the LaCoO$_3$ showed the best stability [6]. Recently, considerable attention has been paid to strontium doped-perovskite materials, e.g., La$_{1-x}$Sr$_x$MnO$_3$ (LSMO) as cathode catalysts. These materials are broadly established as ORR/OER bifunctional electrode materials. The particle size and microstructure are significant for the electrodes’ performance [5, 7]. As one of the most frequently used oxygen electrodes of alkaline [7] and oxide [8] fuel cells, La$_{0.5}$Sr$_{0.5}$MnO$_3$ (LSMO) perovskite oxide has proved promising catalytic properties [8]. Although this perovskite oxide, mentioned above, has provided interesting electrocatalytic properties in fuel cells, the literature however reports, to the best of our knowledge, few studies about the effect of the preparation method using low-temperature synthesis during a short heating time (1 h) on the properties of LSMO perovskite as an oxygen electrode.

Herein, a Pechini route has been successfully used to synthesize homogeneous nanometric La$_{0.5}$Sr$_{0.5}$MnO$_3$ powder with high purity. The influence of anions and cations of electrolytes such as KOH, NaOH, and K$_2$SO$_4$ and mass loading on the ORR kinetics of this perovskite was examined.
The strontium-doped lanthanum manganite perovskite La$_{0.5}$Sr$_{0.5}$MnO$_3$ (LSMO) was synthesized following a procedure similar to that used for other metal oxide powders such as LaMnO$_3$ perovskite [9] and LiMn$_2$O$_4$ spinel [10]. The process involves the dissolution of nitrate salts in a mixture of a carboxylic acid, such as citric acid, and a polyol (ethylene glycol). Previous works on this process have suggested that three basic reactions, i.e., formation of metal chelates, esterification, and polymerization, occur during the formation of the polymeric precursor [11]. A clear solution was obtained by heating a mixture of citric acid and ethylene glycol (1:4 M ratio) at 90 °C for 30 min. The nitrates such as La(NO$_3$)$_3$.6H$_2$O, Sr(NO$_3$)$_2$, and CH$_3$MnO$_4$.4H$_2$O (Sigma-Aldrich) were added in the appropriate amounts based on the stoichiometric ratios of La$_{0.5}$Sr$_{0.5}$MnO$_3$ (La:Sr:Mn = 0.5:0.5:1 M ratio). The resulting solution was further heated at 140 °C for 30 min to induce esterification and remove excess of ethylene glycol. The viscous solution was vacuum dried at 180 °C to yield an organic polymer precursor [12]. Black powders were obtained by heating in air the precursor, at temperatures ranging from 500 to 800 °C with an increment of 100 °C. A single-phase perovskite structure was obtained by heating samples at 700 °C during 1 h. A single crystal of La$_{0.5}$Sr$_{0.5}$MnO$_3$ was grown by the floating zone method (FZM) [13] to confirm the stoichiometry of LSMO by energy-dispersive X-ray spectroscopy (EDX). Thermogravimetric analysis and differential scanning calorimetry (TGA-DSC) were taken with an apparatus SDT Q600 (MT INSTRUMENTS), under a controlled atmosphere of argon.

**Electrochemical Measurements**

All electrochemical experiments were performed at room temperature (25 °C) using an Autolab Potentiostat PGSTAT 30 and a thermostated three-electrode cell. The counter electrode was made of a glassy carbon with a large surface area. A reversible hydrogen electrode (RHE, 0.1 M KOH (pH 13.3), and 0.1 M NaOH (pH 13.2)) was used as the reference electrode. In 0.1 M K$_2$SO$_4$ electrolyte (pH 7.2), a Mercury-mercurous sulfate electrode (MSE: Hg/Hg$_2$SO$_4$, K$_2$SO$_4$ (0.1 M) 0.64 V vs. SHE) was employed. The applied electrode potential in this medium was also quoted with respect to the RHE using the equation \( E (\text{RHE})/V = E (\text{MSE}) + 0.059 \text{pH} + 0.64 \).

Before electrochemical measurements, the electrolyte was purged with nitrogen gas for 10 min. The electrode was subjected to 20 cycles between 0.05 and 1.2 V vs. RHE potential range to clean and activate the electrode surface. Linear sweep voltammetry (LSV) was performed from 1 to 0.2 V vs. RHE over a range of rotation rates of 400, 900, 1600, and 2500 rpm, at a scan rate of 5 mV s$^{-1}$, using a rotating disk electrode in an oxygen-saturated electrolyte solution. The electrolyte solutions were 0.1 M KOH, 0.1 M NaOH, and 0.1 M K$_2$SO$_4$.

**Results and Discussion**

**TGA/DSC Study**

Thermogravimetric analysis and differential scanning calorimetry (TGA-DSC) curves of the polymeric precursor are shown in Fig. 1. The total weight loss was 6 mg from room temperature to 900 °C. The first weight loss estimated to 7% in the temperature range 22–200 °C can be attributed to evaporation of water from LSMO. The second weight loss of 65% in the temperature range 200–350 °C is assigned to evaporation of ethylene glycol, decomposition of citric acid, and evaporation of structurally bounded water. The third step in the
process, ranging from 350 to 700 °C, with 5% weight loss can be explained by decomposition of the metal citrate complex and combustion of the residual organics. This behavior was found to be typical for the pyrolysis of certain citrate precursors [15]. The peaks, located at about 350 °C, are exothermic on the DSC curve due to decomposition of nitrates and organic matter and correspond to the second weight loss as detected in the TGA curve. The decomposition of all nitrates and citrates occurred below 500 °C, leading to the formation of the oxide powder. It can be inferred from these data that the final product can only be obtained by annealing at temperature equal to or higher than 700 °C.

XRD Characterization

Figure 2 shows some selected XRD patterns at 500, 600, and 700 °C obtained after in situ heat treating the gel precursor from room temperature to 700 °C. Scans were recorded at 0.06°/s for 2θ values between 12° and 80° at a heating rate of 10 °C/min. The results indicate that the formation of a perovskite phase started from 600 °C. No shift in the diffraction peaks can be observed for the sample, and all the peaks can be well indexed to a pure perovskite with a tetragonal crystal structure (space group I4/mcm, a = 5.4425 Å, b = 5.4425 Å, and c = 7.7661 Å) according to the PDF ICSD N° 01-089-0786. The average crystallite size (D) was determined using the Debye-Scherrer equation [16]:

\[ D = \frac{0.9 \lambda}{H K \cos \theta} \]  

where \( D \) is the average crystallite size (Å) in the direction hkl, \( H \) the full-width at half-maximum (FWHM) in radian, \( \theta \) the Bragg angle in degrees, and \( \lambda \) the incident X-ray wave length (\( \lambda_{Cu} = 1.5418 \) Å).

The average crystallite sizes (D) of the LSMO sample calcined for 1 h at 600, 700, and 800 °C were 12, 15.8, and 16.4 nm, respectively. This result confirms that the primary particles of the LSMO powders prepared by the Pechini method in the present work are nanosized perovskite phase materials and indicates that the grains’ size increases with the increasing heat-treatment temperature. The two peaks appearing at 44° and 64° on all the diffractograms are attributed to the support Kanthal (FeCrAl alloy) according to the PDF N° 00-042-1486.

Morphology

TEM images of LSMO powders heat-treated at different temperatures are shown in Fig. 3. The images show that all samples are nanoparticulated (sphere-like) practically identical, including quality, shape, and size, with regions showing a roughly agglomeration. The sizes of these nanoparticles (between 12 and 15 nm) are in good agreement with those obtained by the XRD diffraction patterns. The elemental energy-dispersive X-ray (EDX) spectrum in Fig. 3d shows the presence of La, Mn, Sr, and O elements of the sample heat treated at 700 °C for 1 h and their relative abundance in the \( \text{La}_{0.5}\text{Sr}_{0.5}\text{MnO}_3 \) perovskite. The EDX of the particles indicated a homogenous distribution of La, Mn, Sr, and O elements.

Electrochemical Characterization of LSMO/C Electrocatalyst

The ORR catalytic activity of (LSMO/C) heat-treated at different temperatures was measured by RDE depositing the samples with the same loading (68.5 μg\text{LSMO/cm}^2) on a glassy carbon (GC) electrode disk in O2-saturated 0.1 M
KOH electrolyte. Regardless of the heat treatment temperature, the LSV curves display a mixed kinetic diffusion-controlled region at high potentials and a diffusion-limiting current ($j_{\text{lim}}$) region at low potentials, see Fig. 4. The comparison between LSV curves indicates that the LSMO/C electrode heat-treated at 700 °C showed the highest catalytic activity with a concomitant half-wave potential ($E_{1/2}$) shifted to more positive potential and higher limiting current density ($j_{\text{lim}}$) than those heat treated at 500 and 600 °C. This phenomenon, apparently, puts in evidence the crystalline state of the sample [17]. Therefore, heat treatment at this temperature was selected for further experiments. The LSV curve of LSMO heat treated at 500 °C provided the highest surface area, but exhibited a poor catalytic activity with the lowest half-wave potential.

To understand whether this activity is attributed to the nanocrystalline state of the sample, the electrochemically active area was measured, as described in reference [18], for the three samples by evaluating the oxide roughness factor (RF), i.e., the ratio of the real area ($S_{\text{LMSO}}$) of a rough electrode to its geometrical one. The capacitive current density, $j_{\text{cap}}$, was determined as a function of the scan rate (30 to 80 mV s$^{-1}$) by recording cyclic voltammograms for all three heat-treated samples (500, 600, 700 °C) in a small potential region of 100 mV (i.e., 0.35 to 0.45 V). A typical example is depicted in the inset of Fig. 5. The double-layer capacitance ($C_{dl}$) was estimated from the slope of Fig. 5. The roughness factor (RF) was then calculated by assuming a double-layer capacitance of 60 μF cm$^{-2}$ for a smooth oxide surface [19]. Values of $C_{dl}$ and RF for the three samples are summarized in Table 1.

From the table, RF specifies that LSMO oxide calcined at 500 °C has the highest electrochemically active area than those calcined at 600 and 700 °C. Although the surface area value of LSMO/C/500 °C appears to be the highest, it revealed
the poorest electrocatalytic activity, suggesting that ORR process on LSMO/C is, therefore, assigned to the degree of crystallinity rather than to the surface area. To assess the effect of carbon Vulcan on the electroactivity of the LSMO perovskite, LSV voltammograms of the carbon-supported perovskite were compared to those non-supported perovskite, Fig. 6a, CVs at 50 mV/s for carbon Vulcan, LSMO/C and LSMO, see Fig. 6b. It can be observed, according to cyclic voltammograms for carbon in O₂ and N₂ saturated 0.1 M KOH, respectively, that carbon actively contributes in the ORR on LSMO/C composites. Composite electrodes as LSMO/C activate the ORR via four-electron pathway whereas carbon alone is only able to catalyze the two-electron one [14]. It comes up clearly that using Vulcan carbon as a support for LSMO improves the charge transfer of nanoparticulated LSMO at overpotential beyond the activation region, as compared to non-supported LSMO, which shows kinetic hindrance in agreement with the literature [20, 21]. Within the ORR-recorded potential interval, we can qualitatively evaluate the half-wave potential, $E_{1/2}$, and the kinetic current density, $j_k$, increase following this order $K_2SO_4 < NaOH < KOH$. Taking into consideration the roughness factor (RF) of each sample, the values of the real ORR kinetic current density obtained at 0.7 V (see insert in Fig. 7) in KOH, NaOH, and $K_2SO_4$ electrolytes were found to be 96.7, 49.3, and 14.3 $\mu$A/cm², respectively, indicating that KOH solution is a superior medium for ORR under both thermodynamic and kinetic considerations in agreement with the literature [22]. However, the high improvement of ORR activity on Pt electrode was assigned to the high oxygen solubility and diffusivity and lower viscosity of the solution by Jin et al. [23]. The oxygen diffusivity is related to the viscosity, $\eta$, of the electrolyte, and the diffusion coefficient $D_{O_2}$ is given by the Stokes-Einstein equation [24]:

$$D_{O_2} = \frac{k_BT}{6\pi\eta r}$$

where $r$ is the effective hydrodynamic radius of the oxygen molecule, $k_B$ is the Boltzmann constant, and $T$ is the absolute temperature. As the viscosity of NaOH electrolyte is higher than that of KOH [23], the $D_{O_2}$ in NaOH is lower than that in KOH; thus, a smaller limiting current density $j_{lim}$ is expected, according to Levich equation [25]. Furthermore, because Na⁺ ion is smaller in size and higher in charge density than the K⁺ ion, the salting-out effect, based on the electrolyte-oxygen interaction, is stronger in NaOH than in KOH [26], leading to the inhibition of oxygen dissolution and, consequently, to a smaller limiting current density in NaOH solution as seen in Fig. 7. Suntivich et al. [27] and Strmcnik et al. [28] proposed that the non-covalent interaction between cations (K⁺, Na⁺) and the oxide does not lead to the blockage of ORR active sites, but rather to the modification of the reaction intermediate energy. In the case of K⁺ which has the weakest interaction

<table>
<thead>
<tr>
<th>Heat-treated LSMO sample</th>
<th>$C_{dl}$ (μF/cm²)</th>
<th>RF</th>
<th>$j_{lim}$ mA/cm²</th>
<th>$j_k$ μA/cm²</th>
</tr>
</thead>
<tbody>
<tr>
<td>700 °C</td>
<td>2880</td>
<td>48</td>
<td>5</td>
<td>104</td>
</tr>
<tr>
<td>600 °C</td>
<td>3480</td>
<td>58</td>
<td>3</td>
<td>51</td>
</tr>
<tr>
<td>500 °C</td>
<td>3600</td>
<td>60</td>
<td>1.028</td>
<td>21</td>
</tr>
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</table>
with the $OH_{\text{surf}}$, the modification is minimal, and thus, the ORR proceeds as if the cation was not present. In the case of sulfate anions $SO_4^{2-}$, the ORR overpotential losses and the electroactivity increased. In general, the presence of strongly adsorbed anions in the electrolytes such as $H_2SO_4$, $H_3PO_4$ results in the appearance of high overpotentials for the ORR on Pt cathode. This effect is not observed, however, when the ORR is evaluated in non-adsorbing (negligible) electrolytes such as HClO$_4$ or alkaline electrolytes [29]. For instance, poorer performance of Pt cathodes for the ORR in concentrated $H_3PO_4$ than in KOH may be attributed to limitations arising from adsorption of phosphoric acid anions [30]. According to Dominguez et al. [31], the observed negative effect of sulfates for the ORR on both Fe/N/C and Fe/N/CNT composites in 0.5 $H_2SO_4$ is due to both the removal of acid-soluble Fe species and the presence of sulfates onto the active sites. These anions remain bonded to the active sites rather than simply blocking the access of $O_2$ into the active sites.

**Effect of the LSMO Loading**

In order to investigate the optimal amount of LSMO in the LSMO/C for best ORR activity, LSMO/C electrocatalysts with various mass loadings of the LSMO oxide, heat-treated at 700 °C, were studied and evaluated. Figure 8 represents LSVs obtained on the LSMO/C electrodes in 0.1 M KOH under the same conditions, that is, oxygen-saturated electrolyte, 25 °C, and an angular velocity $\omega$ of the RDE of 900 rpm. To compare the electrocatalytic activity of LSMO/C electrodes with that of benchmark materials, such as Pt/Vulcan [32], a commercial Pt/C (20 wt.% Pt on carbon, Johnson-Matthey) was also tested under the same conditions. In the studied interval of catalyst mass loadings, it is observed that the higher the mass loading added of LSMO/C, the higher the half-wave potential and the limiting and kinetic current densities. Within the experimental error, mass loadings of LSMO/C of 68.5 $\mu g_{LSMO}/cm^2$ and that of 96 $\mu g_{LSMO}/cm^2$ delivered the same results, demonstrating that this mass loading between 70 and 90 $\mu g_{LSMO}/cm^2$ can be considered as an...
The active surfaces of the various sample loadings were estimated by evaluating the LSMO roughness factors which were found to be 39, 45, and 48 for LSMO loadings 27.5, 41, and 68.5 μgLSMO/cm², respectively. As expected, the RF increases with increasing LSMO loading. Based on the results provided by the LSV and capacitive curves, it should be pointed out that the ORR activity depends intimately on the catalytic site density. The influence of catalytic loading on the ORR activity was recently reported on precious and non-precious metal centers, such as Pt/C [33], CoSe2/C in acid [34], and alkaline media [35]. It should be mentioned that an increase of catalyst depositing volume from 0.5 to 7 μL suggests the higher thickness of LSMO/C film which reflects the higher number of site density. As discussed previously in detail [36], the activity of the catalyst is directly related to the site density.

To examine the pathway for the ORR via two or four electrons, LSV curves at various rotation speeds \( \omega \) were plotted. Figure 10a, given as an example, shows for LSMO/C loaded with a mass of 68.5 μgLSMO/cm² calcined at 700 °C that the limiting current density increases with increasing \( \omega \) due to the improved mass transport of oxygen at the electrode surface. This series of experiments assesses the idea that the ORR process is essentially being done via a series process, whatever the nature of the catalytic center, as demonstrated herein with LSMO perovskites, on LaCoO3 [37], and other kinds of electrode materials [34–36].

The desired feature of a good catalyst towards the ORR would be to reduce molecular oxygen to \( \text{OH}^- \) through the direct four-electron process [38]. A two-electron reduction of \( \text{O}_2 \) via \( \text{HO}_2^- \) peroxide [39] leads to a lower energy conversion efficiency. In order to verify the ORR pathway, Koutecky-Levich (K-L) plots corresponding to the LSV curves of Fig. 10a are plotted in Fig. 10b using the Koutecky-Levich (K-L) equation [25]:

\[
\frac{1}{j} = \frac{1}{j_k} + \frac{1}{j_{\text{lim}}} = \frac{1}{j_k} + \frac{1}{B \omega^{1/2}} \quad (3)
\]

\[
B = 0.62 nFAC_O D_O^{2/3} \nu^{-1/6} \quad (4)
\]

where \( j, j_k, \) and \( j_{\text{lim}} \) are the measured disk current density, kinetic current density, and diffusion-limiting current density, respectively, and \( \omega \) is the electrode rotation rate (rad s\(^{-1}\)). Furthermore, \( n \) is the number of electrons in the ORR, \( F \) is the Faraday constant (96,500 C mol\(^{-1}\)), \( A \) is the area of the disk electrode (0.07 cm\(^2\)), \( C_O \) is the oxygen concentration in 0.1 M KOH (1.14 × 10\(^{-3}\) mol cm\(^{-3}\)), \( D_O \) is the oxygen diffusion coefficient in 0.1 M KOH (1.73 10\(^{-5}\) cm\(^2\) s\(^{-1}\)), and \( \nu \) is the kinematic viscosity of the 0.1 M KOH solution (0.01 cm\(^2\) s\(^{-1}\)). All plots in Fig. 10b determine linear and parallel relationship with \( j_{\text{lim}}^{-1} \) as function of \( \omega^{-1/2} \) at four
selected applied electrode potentials, implying therefore consistent diffusional characteristics of ORR kinetics on the LSMO/C electrocatalyst. From the slopes, B of the K-L plots (Fig. 10b), calculation of the number of electrons transferred per O₂ provides, as for as Pt/C composite, four electrons for LSMO/C.

Conclusion

This study was carried out to evaluate the activity of the perovskite La₀.₅Sr₀.₅MnO₃/C used as an electrocatalyst for the oxygen reduction reaction (ORR). The ORR on the LSMO/C electrocatalyst surface was studied using voltammetry techniques and the influence of heat treatment temperature. The mass loading and the nature of electrolyte were also examined. The results obtained allowed the following conclusions:

(i) The Pechini method was successfully used at low temperature to produce La₀.₅Sr₀.₅MnO₃ homogeneous nanometric-sized material with high purity.

(ii) The LSMO calcined at 700 °C for 1 h showed higher electroactivity in 0.1 KOH with a higher half-wave potential than its counterparts heat-treated at 500 and 600 °C.

(iii) The performance for ORR of the LSMO/C electrocatalyst in different electrolytes, e.g., KOH, NaOH, and K₂SO₄, was examined. Higher ORR electroactivity has been observed in KOH electrolyte as compared to NaOH and K₂SO₄ ones.

(iv) The higher the LSMO loading added to the LSMO/C composite, the higher is the ORR electroactivity. The LSMO/C with higher loadings exhibited an encouraging electrocatalytic activity, approaching that of the more expensive fuel cell Pt/C cathode examined under the same conditions.

(v) By examining Koutecky-Levich plots, the ORR materials’ activity is apparently due to four-electron charge transfer. This is the result of increasing the number of accessible active sites with the mass loading.

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