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Tellurium doped Lanthanum Manganite as Catalysts for the Oxygen Reduction Reaction

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Abstract

The effect of Te doping on the electrocatalytic activity of $\text{La}_{1-x}\text{Te}_x\text{MnO}_3$ towards the oxygen reduction reaction is investigated for the first time. $\text{La}_{1-x}\text{Te}_x\text{MnO}_3$ with x values up to 23% were synthesized from a single ionic liquid based precursor, yielding nanoparticles with mean diameter in the range of 40 to 68 nm and rhombohedral unit cell. Electrochemical studies were performed on carbon supported particles in alkaline environment. The composition dependence activity is discussed in terms of surface density of Mn sites and changes in the effective Mn oxidation state.

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Introduction

Oxygen electrocatalysis is one of the key process limiting the efficiency of energy conversion devices such as fuel cells, electrolyzers and metal-air batteries.¹⁻² In particular, the oxygen reduction reaction (ORR) is commonly associated with slow kinetics, requiring high overpotentials and high catalyst loadings. Current research activities are also focused on the development of non-noble metal electrocatalysis such as transition metal oxides.^{9,10} A number of studies have shown that Mn-based transition metal oxides are among the most catalytically active for the ORR in alkaline solutions.³⁻⁶ However, the parameters determining the reactivity of Mn sites in this complex systems remains a subject of discussion.

In the case of perovskites oxides (ABO_3), the key active site is determined by the B-cation. A multiplicity of parameters have been linked to electrocatalytic activity such as the nature of the A-site, d -orbital occupancy in the B-site, oxidation state, A-site surface segregation, oxygen vacancies, particle size and morphology.⁷⁻¹⁰ Suntivich *et.al.* proposed that single occupancy of e_g orbital gives the highest ORR activity,³ which appears to be supported by DFT studies reported by Calle-Vallejo and co-workers.⁷ However, recent studies by Celorrio *et.al.* showed that the B-site orbital occupancy may change under operational conditions, concluding that Mn-sites are uniquely active as they undergo changes in oxidation state in the region close to the formal ORR potential.¹¹ This point has also been recognized by other groups.¹²⁻¹³ The orbital occupancy of Mn can be influenced by the cations occupying the A-site, which might lead to tuning the activity of the catalysts.⁵ Celorrio *et al.* also demonstrated that increasing the amount of Ca^{2+} in $La_{1-x}Ca_xMnO_3$ not only increases the Mn oxidation state, but also decreases the average activity.⁵

In this study, we investigate tellurium-doped lanthanum manganite as an ORR catalyst for the first time. The rationale of Te doping is based on the studies by Yang *et.al.*, showing that $\text{La}_{1-x}\text{Te}_x\text{MnO}_{3+\delta}$ lead to high ratio $\text{Mn}^{2+}/\text{Mn}^{3+}$.¹⁴ $\text{La}_{1-x}\text{Te}_x\text{MnO}_3$ is synthesized from a versatile ionic liquid precursor, yielding nanoparticles with size ranging between 40 and 68 nm. At a fixed catalyst loading, the overall performance appears to decrease upon Te doping with respect to LaMnO_3 . However, normalization of the activity by the number density of electroactive Mn sites reveals a significant improvement of the B-site activity upon 10% Te doping of the A-site.

Experimental

$\text{La}_{1-x}\text{Te}_x\text{MnO}_3$ were synthesized employing an ionic liquid precursor based on the methodology described in previous studies.^{5, 11, 15} Firstly, ~9 mL 69% HNO_3 was added to a glass vial containing TeO_2 (0.001 mol) and heated to 80 °C under stirring for 3 hours, until the TeO_2 had completely dissolved. The colourless solution was diluted to 10 mL with deionized water to give a 0.1 M solution of Te^{4+} in aqueous HNO_3 . Later, 0.029 g ethylenediaminetetraacetic acid (EDTA) (0.01 mmol, 1:1 molar ratio of metal precursors:EDTA) was added to a 1 mL 1-ethyl-3-methylimidazolium acetate stirred at 80 °C. Then, 0.1 M aqueous $\text{Mn}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$, $\text{La}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, and the 0.1 M Te^{4+} solution in aqueous HNO_3 were added to the ionic liquid in the appropriate stoichiometric ratio. The total volume of aqueous solution added was 1 mL. Solutions were dehydrated by heating at 80 °C under stirring for 3 hours. Following dehydration, the mixture was transferred to an alumina crucible containing 100 mg microcrystalline cellulose. After 10 minutes of stirring at ~80 °C, a slightly viscous homogeneous gel was obtained. The crucible was transferred to a furnace,

and calcined at 850 °C for 4 hours with a temperature ramp rate of 5 °C/min. This temperature enabled the formation of phase pure $\text{La}_{1-x}\text{Te}_x\text{MnO}_3$ with x values below 25%

Powder X-ray diffraction was carried out on a Bruker D8 Advance using $\text{Cu K}\alpha$ radiation. Experiments were run between 10 and 80 degrees, using a step size of 0.02 degrees. The XRD patterns were refined by the Rietveld method using the FULLPROF program.¹⁶⁻¹⁷ The following parameters were refined: scale factor, background coefficients, zero-point error and positional coordinates. The isotropic thermal factors for all the atoms were fixed to a standard value of 1 \AA^2 . No cation deficiency was detected in the case of LaMnO_3 , thus the occupancy factors were fixed to 1. For $\text{La}_{1-x}\text{Te}_x\text{MnO}_3$, the occupancy factors for La and Te were also refined. Scanning electron microscopy was carried out on a JEOL SEM 5600LV scanning electron microscope. Transmission electron microscopy was carried out on a JEOL JEM-1400Plus instrument.

XPS was carried out at the Bristol NanoESCA Facility equipped with Argus spectrometer working at a base pressure of 4.0×10^{-11} mbar. Core level photoemission spectra were acquired in grazing incidence, i.e. 45° between the sample surface and the normal of the electron analyzer, with a monochromatic $\text{Al K}\alpha$ (1486.7 eV), pass energy of 20 eV at room temperature. The binding energy scale was referenced on the carbon C 1s photoemission line.

Cyclic voltammetry was carried out in argon saturated 0.1 M KOH solution using a glassy carbon working electrode, graphite counter electrode, and Hg/HgO reference electrode. Linear sweep voltammetry employing a rotating ring-disc electrode were carried out in O_2 saturated 0.1 M KOH solution using an ALS rotation controller, and an Ivium compactstat potentiostat. A glassy carbon disk (0.126 cm^2) and a Pt ring were used in all experiments. Working electrodes were prepared via a two-step drop casting process. First, an aqueous

Vulcan/Nafion suspension was dropped onto the surface of the electrode, followed by an aqueous suspension of the metal oxide nanoparticles. The sample loading on the electrode surface was $50 \mu\text{g}_{\text{vulcan}} \text{cm}^{-2}$, $50 \mu\text{g}_{\text{nafion}} \text{cm}^{-2}$ and $250 \mu\text{g}_{\text{oxide}} \text{cm}^{-2}$.

Results and discussion

Figure 1a shows the XRD patterns of LaMnO_3 , $\text{La}_{0.90}\text{Te}_{0.10}\text{MnO}_3$ and $\text{La}_{0.77}\text{Te}_{0.23}\text{MnO}_3$ as well as the result of the Rietveld refinement employing a rhombohedral lattice with the space group $R\bar{3}c$. LaMnO_3 and $\text{La}_{0.90}\text{Te}_{0.10}\text{MnO}_3$ are characterized by high degree of phase purity, whereas $\text{La}_{0.77}\text{Te}_{0.23}\text{MnO}_3$ exhibit a small peaks (marked as *) associated with an unidentified secondary phase. Ionic liquid precursors incorporating higher Te content yield multiple phases, in agreement with previous studies.¹⁸ Structural parameters were obtained by the Rietveld method and can be found in Table S1 of the Supporting information. The results show a minimum in the unit cell volume with increasing Te content (changes below 0.2%), while the Mn-O bond length increases by approximately 0.1% in $\text{La}_{0.77}\text{Te}_{0.23}\text{MnO}_3$. These changes in structural parameters appear consistent with previous studies.¹⁹

Figure 1b shows a representative SEM image of LaMnO_3 , illustrating the typical porous microstructure of the synthesized materials, while the TEM image in Figure 1c shows the nanoscale dimensions of the particles (SEM and TEM images of the other oxides can be found in Figure S1 and S2 of the Supporting Information, respectively). Particle size distributions obtained from measuring at least 80 particles can be found in Figure S3, revealing a slight increase in mean particle size with increasing Te content, from 39.9 ± 11.3 nm for LaMnO_3 to 67.9 ± 38.3 nm for $\text{La}_{0.77}\text{Te}_{0.23}\text{MnO}_3$ (see table S2).

XPS spectra of LaMnO_3 , $\text{La}_{0.9}\text{Te}_{0.10}\text{MnO}_3$, $\text{La}_{0.77}\text{Te}_{0.23}\text{MnO}_3$ are displayed in Figure 2, showing the La $3d$ (left), Mn $2p$ (middle) and Te $3d$ (right) regions. The characteristic double peak associated with La $3d_{5/2}$ (834.0-834.9 eV) and La $3d_{3/2}$ (851.0-851.9 eV) are observed, with the La $3d_{5/2}$ binding energy consistent with a La^{3+} oxidation state.^{5, 11, 20-22} The peak at 874 eV, visible for the Te-containing samples has been assigned to the Te Auger MNN. The Te $3d$ photoemission lines involves the Te $3d_{5/2}$ at 576 eV and Te $3d_{3/2}$ at 586.6 eV. These values are consistent with a Te^{4+} oxidation state.²³⁻²⁴ Finally, the photoemission line associated with Mn $2p_{3/2}$ peak centered at 642 eV shows an interesting trend as function of Te content. This line contains contribution from Mn^{2+} (640.9 eV) and Mn^{3+} (641.9 eV) oxidation state.²⁵ Focusing on this narrow emission range (Figure S4), we can see a slight shift of the Mn $2p_{3/2}$ peak towards lower binding energy as the Te content increases. However, given the small differences in binding energies for the two oxidation states, we shall refrain from attempting to estimate $\text{Mn}^{2+}/\text{Mn}^{3+}$ ratio from this data.

Analysis of the relative composition of La, Te and Mn, after normalization of the individual photoemission lines by their sensitivity factor and inelastic mean free path, show a significant A-site surface enrichment. We obtained an A:B ratio of 62:38 for all three materials, broadly in agreement with data reported for other lanthanides.^{5, 26} We have also estimated La:Te ratios of 82:18 and 71:29 for $\text{La}_{0.90}\text{Te}_{0.10}\text{MnO}_3$ and $\text{La}_{0.77}\text{Te}_{0.23}\text{MnO}_3$, respectively. This analysis show both La and Te substantially segregate to the surface, with a slight increase in Te content with respect to the bulk values.

Figure 3 shows cyclic voltammograms recorded in an argon-saturated 0.1 M KOH solution of the carbon-oxide composite electrodes, revealing a two-step reduction from the initial Mn oxidation state to Mn^{2+} .^{5, 11-12} The position of the peak centered at 0.90 V remains almost unaffected, while the second reduction peak is slightly shifted towards higher potentials

upon increasing Te content. The most striking difference is in the current densities, with both Te doped materials showing significantly lower responses than LaMnO_3 . Considering that the catalyst loading is identical (i.e. $250 \mu\text{g}\times\text{cm}^{-2}$), this behavior indicates a decrease in the number density of electrochemically active Mn-sites at the electrode surface. The faradaic charges associated with the anodic and cathodic peaks are very similar, and the responses remain stable in the studied potential range, suggesting that the stability is not significantly compromised within the timescale of these experiments. Integration of the cathodic current peaks in the range of 0.2 to 1.2 V enables to estimate the effective number density of electroactive Mn sites (Γ_{Mn}).⁵ In order to calculate Γ_{Mn} , two limiting cases can be assumed with regards to the initial Mn oxidation state: the oxidation state is determined by the mean oxidation state of the A-site taking La^{3+} and Te^{4+} (Case 1), or fixed Mn oxidation state of +3 independently of the A-site composition (Case 2). Case 1 implicitly ignores changes in the oxygen content of the oxide, while Case 2 assumes that variations in the mean A-site charge is compensated by changes in oxygen stoichiometry. Our previous studies investigating $\text{La}_{1-x}\text{Ca}_x\text{MnO}_3$,⁵ in which the mean Mn oxidation state is directly estimated by XANES, confirms Case 1 as a valid approximation. In any case, Table S3 shows that both limiting cases provide values within the same order of magnitude.

The data in Table S3 shows the Γ_{Mn} is 40 to 60% smaller in the Te doped samples. In view of the similar extent of A-site surface segregation, the smaller Γ_{Mn} values in the case of the Te containing particles is mainly linked to smaller specific surface area. Although the mean particle size increases with increasing Te content, we see statistically similar Γ_{Mn} in for both Te containing oxides. This is due to the fact that size dispersion also increases in substantial fashion as shown in figure S3. It should also be mentioned that Γ_{Mn} in LaMnO_3 prepared at

850°C (this report) is lower than those prepared at 700°C from the same ionic liquid precursor,^{5, 11} confirming that calcination temperature is an important parameter to determining A-site segregation.

Figure 4 compares the current measured at the disk (i_{DISK} , bottom panel) and the ring (i_{RING} , top panel) of carbon-supported LaMnO_3 , $\text{La}_{0.90}\text{Te}_{0.10}\text{MnO}_3$ and $\text{La}_{0.77}\text{Te}_{0.23}\text{MnO}_3$ electrodes at 1600 rpm in O_2 -saturated 0.1 M KOH solution. The responses obtained from the Vulcan carbon support under the same experimental conditions are shown in Figure S5. All of the catalysts show significantly larger currents than the carbon support, while the ORR onset potential is over 100 mV more positive. LaMnO_3 exhibit larger cathodic current at the disc, with smaller ring current across the potential range. Current-potential curves recorded for $\text{La}_{0.77}\text{Te}_{0.23}\text{MnO}_3$ particles at various angular rotation rates are displayed in figure S6. Both Te doped oxides exhibit rather similar current-potential characteristics.

Figure 5 illustrates the dependence of i_{DISK} with the inverse of the angular rotation rate in the case of LaMnO_3 (figure 5a) and $\text{La}_{0.77}\text{Te}_{0.23}\text{MnO}_3$ (figure 5b), which follows the Koutecky-Levich relationship:

$$\frac{1}{i_{DISK}} = \frac{1}{i_k} + \frac{1}{i_L} = \frac{1}{i_k} + \frac{1}{0.62nAFcD^{2/3}\nu^{-1/6}\omega^{1/2}} \quad (1)$$

where n is the number of transferred electrons, A is the disk geometric area, F is the Faraday constant, c is the bulk oxygen concentration ($1.2 \cdot 10^{-6} \text{ mol cm}^{-3}$),²⁷ D is the oxygen diffusion coefficient ($1.9 \cdot 10^{-5} \text{ cm}^2 \text{ s}^{-1}$), ν is the kinematic viscosity ($0.01 \text{ cm}^2 \text{ s}^{-1}$), and ω is the angular rotation of the electrode. i_k and i_L are the kinetically and mass-transport limiting currents, respectively. It can be observed that LaMnO_3 (Figure 5a) shows a linear dependence, with a

slope consistent with $n = 4$.^{5, 11} In the case of $\text{La}_{0.77}\text{Te}_{0.23}\text{MnO}_3$ (Figure 5b), only a slight deviation from $n = 4$ is observed at high rotation rate.

The HO_2^- yield (2 e-electron process) as a function of potential was calculated from the ring and the disk current values at 1600 rpm (Figure S7a). The Vulcan support shows almost 100% HO_2^- yield between 0.2 and 0.6 V, while LaMnO_3 and both Te-doped catalysts exhibits less than 30%. The pathway selectivity can also be described in terms of the effective number of electrons shown in Figure S7b. The Vulcan support yields values close to 2, increasing to values above 3.4 upon introduction of the oxide catalysts.

These findings suggest that LaMnO_3 exhibits better performance than the Te doped under the specific catalyst formulation used, i.e. at equivalent mass loading. However, it should also be considered that mean particle sizes and size dispersion are composition dependent, which have an effect on the overall catalysts performance. Furthermore, XPS data show an increase in the A-site segregation in the presence of Te, although the surface composition remains dominated by La^{3+} . In order to account for these parameters, the kinetically limiting current can be normalized by Γ_{Mn} , i.e. $\hat{i}_k = i_k / (A \times \Gamma_{\text{Mn}})$.⁵

Figure 5c illustrates the variation of \hat{i}_k at 0.65 V as a function of the Te content, obtained from at least three different samples from various synthesis batches. The contribution of the carbon support to the overall current at this potential is negligible in comparison to the oxide loaded catalysts (see figure S4). Two sets of data are plotted based on the two limiting cases used for calculating Γ_{Mn} . It is interesting to see the performance of electrochemically active surface Mn sites increases on average in the presence of Te. Surface Mn sites in materials containing 10% Te appears twice more active than surface Mn-sites in LaMnO_3 . The performance seems to slightly degrade for larger Te content.

We rationalize this early trend in terms of changes in the electron density at Mn sites. Incorporating 10% Te in the A-site, most probably in a 4+ oxidation state, effectively increases the electron density at the B-site, leading to an enhancement in the catalytic activity for ORR. Indeed, the slight shift in the binding energy of the Mn 2p_{3/2} peak suggests a decrease in the oxidation state on Mn sites in the presence of Te (figures 2 and S4). A similar correlation have been recently reported in the case La_{1-x}Ca_xMnO₃.⁵

In conclusion, these studies reveal an increase in the mean activity of surface Mn sites upon Te⁴⁺ doping in the A-site. The most active surface Mn sites were obtained upon 10% replacement of La³⁺ by Te⁴⁺, leading to an increase of the electron density at the Mn site without compromising the coordination structure. To confirm this hypothesis, a number of parameters should be systematically explored such as the effect of low oxidation state cations with different sizes as well as the activity of different LaMnO₃ phases. In any case, our studies demonstrate the need of establishing universal descriptors, such as $\hat{\mu}_k$, which uncovers the mean activity of surface Mn sites, independently of the particle size, dispersion quality and complex phenomena such as A-site segregation.

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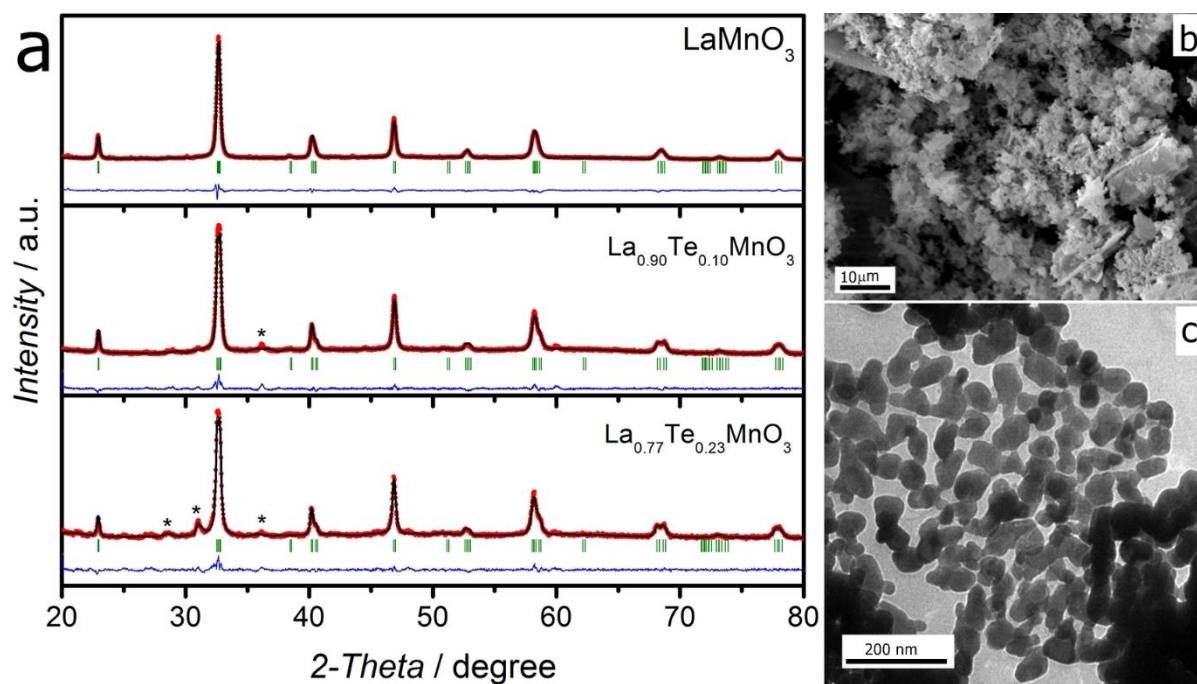


FIG. 1 (a) XRD patterns of LaMnO_3 , $\text{La}_{0.90}\text{Te}_{0.10}\text{MnO}_3$ and $\text{La}_{0.77}\text{Te}_{0.23}\text{MnO}_3$. Red dots correspond to the experimental values while the black thin line corresponds to the Rietveld refinement. The thin blue line shows the difference between experimental and refined patterns. The green bars correspond to the positions of the allowed Bragg reflections for the main phase. (b) SEM and (c) TEM images of as-prepared LaMnO_3 .

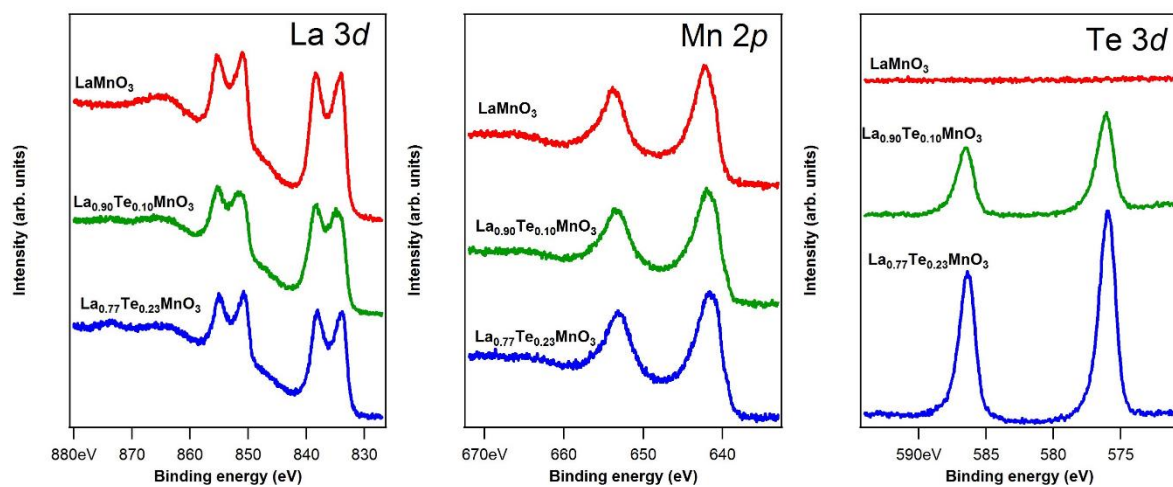


FIG. 2 Photoemission spectra of La 3d (left panel), Mn 2p (middle panel) Te 3d (right panel) of the various $\text{La}_{1-x}\text{Te}_x\text{MnO}_3$ oxides taken in grazing incidence using a monochromatic Al $K\alpha$ X-ray source.

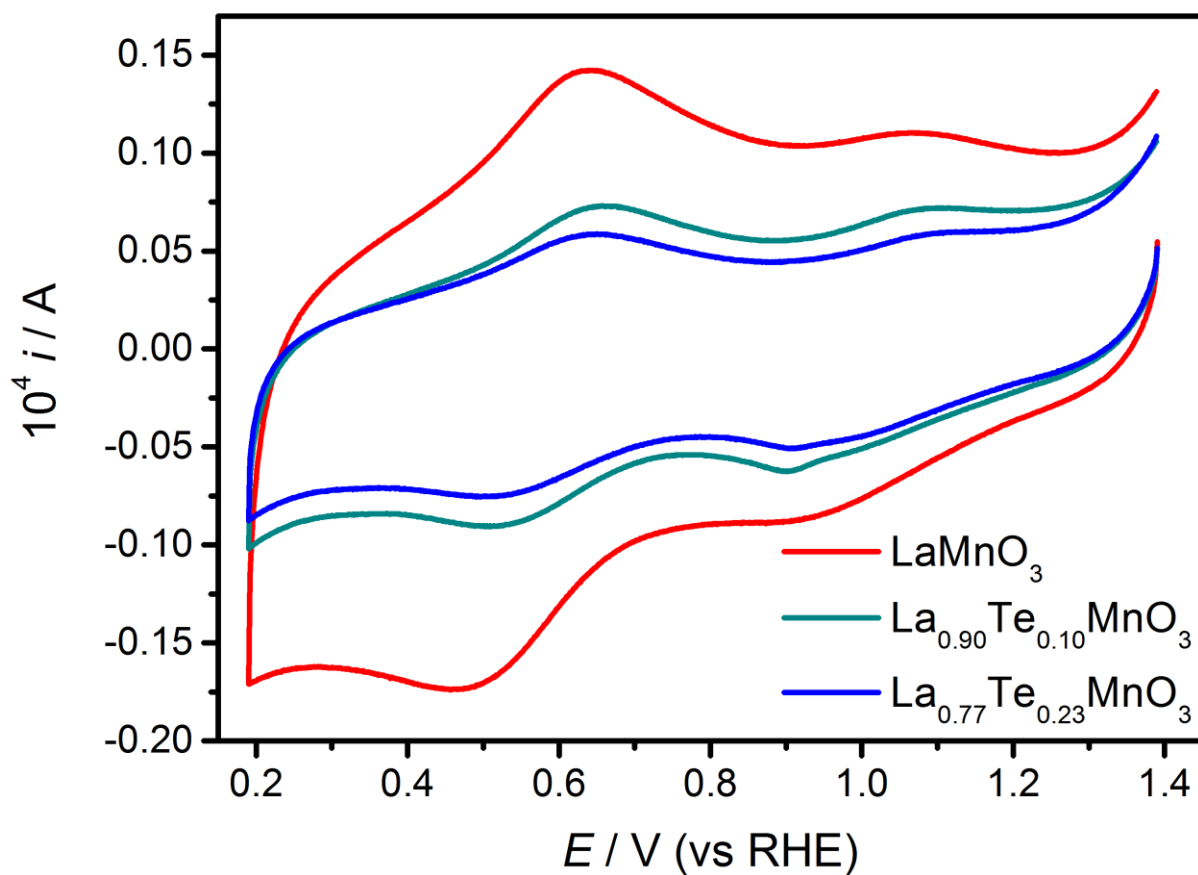


FIG. 3 Cyclic voltammograms of LaMnO_3 , $\text{La}_{0.90}\text{Te}_{0.10}\text{MnO}_3$ and $\text{La}_{0.77}\text{Te}_{0.23}\text{MnO}_3$ supported on Vulcan-carbon in Ar-saturated 0.1 M KOH electrolyte solution at 0.01 V s^{-1} . Electrode loaded with $50 \mu\text{g cm}^{-2}_{\text{nafion}}$, $50 \mu\text{g cm}^{-2}_{\text{vulcan}}$, and $250 \mu\text{g cm}^{-2}_{\text{oxide}}$.

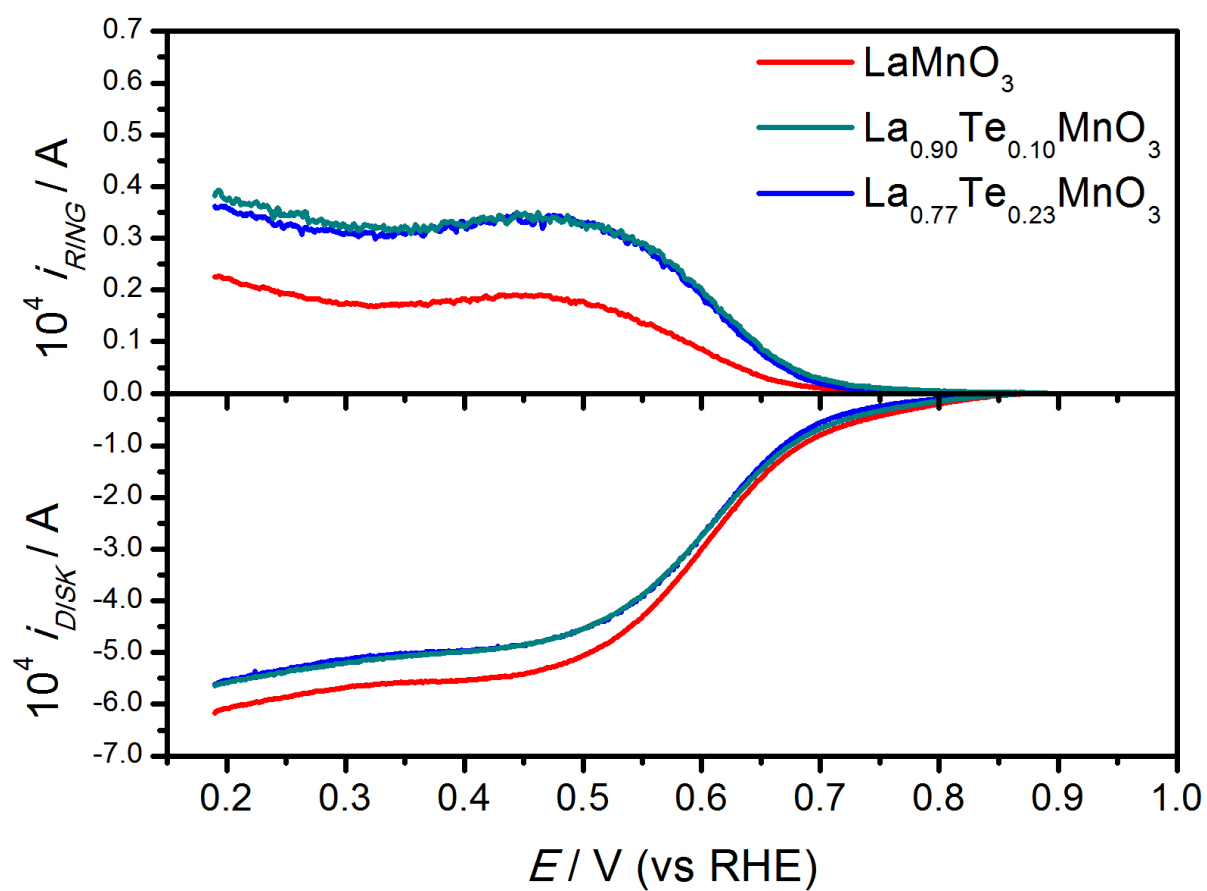


FIG. 4 RRDE responses of the various $La_xTe_{1-x}MnO_3$ nanoparticles supported at Vulcan layer at 1600 rpm in O_2 -saturated 0.1 M KOH at 0.010 V s^{-1} . The Pt ring was held at a constant potential of 1.10 V. The oxide content in each electrode was $250 \mu\text{g cm}^{-2}$.

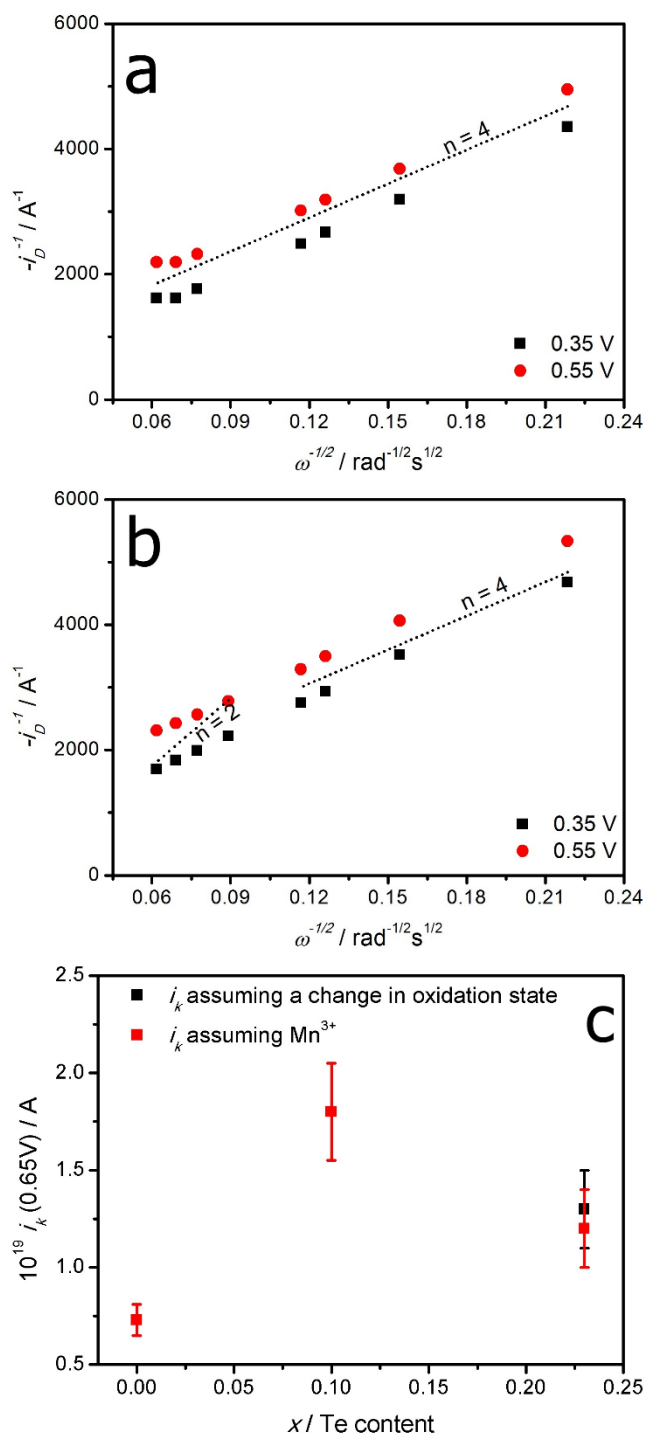


FIG. 5. Koutecky–Levich plots of (a) LaMnO₃ and (b) La_{0.77}Te_{0.23}MnO₃ electrodes in O₂-saturated 0.1 M KOH at 0.55 V and 0.35 V. Dotted lines represent the limiting slopes for the 2 and 4-electron processes. (c) Kinetically limited current at 0.65 V normalised by the effective number of electroactive Mn atoms (\hat{i}_k)