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Nanoparticle-Decorated Ultrathin La₂O₃ Nanosheets as an Efficient Electrocatalysis for Oxygen Evolution Reactions

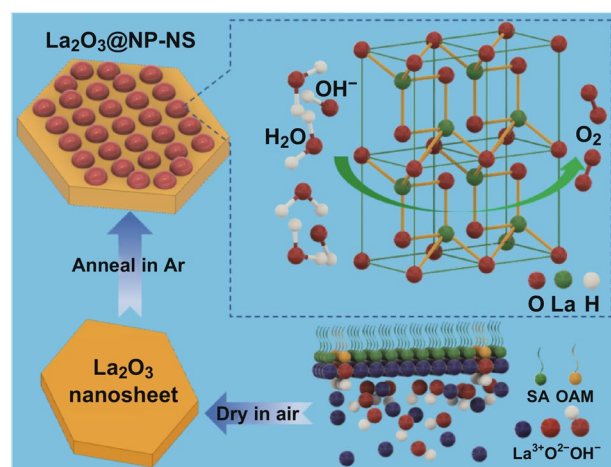
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HIGHLIGHTS

- The 2.27-nm-thick hybridized quasi-2D structure of La₂O₃ crystalline nanoparticles embedded in La₂O₃ amorphous nanosheets (La₂O₃@NP-NS) exhibited a low overpotential of 310 mV at 10 mA cm⁻².
- The mass activity of La₂O₃@NP-NS reached as high as 6666.7 A g⁻¹ at overpotential of 310 mV. Such a high mass activity was more than three orders of magnitude higher than that of benchmark IrO₂ (4.4 A g⁻¹) and RuO₂ (2.05 A g⁻¹) and five orders of magnitude higher than that of commercial La₂O₃ (0.048 A g⁻¹).

ABSTRACT Electrochemical catalysts for oxygen evolution reaction are a critical component for many renewable energy applications. To improve their catalytic kinetics and mass activity are essential for sustainable industrial applications. Here, we report a rare-earth metal-based oxide electrocatalyst comprised of ultrathin amorphous La₂O₃ nanosheets hybridized with uniform La₂O₃ nanoparticles (La₂O₃@NP-NS). Significantly improved OER performance is observed from the nanosheets with a nanometer-scale thickness. The as-synthesized 2.27-nm La₂O₃@NP-NS exhibits excellent catalytic kinetics with an overpotential of 310 mV at 10 mA cm⁻², a small Tafel slope of 43.1 mV dec⁻¹, and electrochemical impedance of 38 Ω. More importantly, due to the ultrasmall thickness, its mass activity, and turnover frequency reach as high as 6666.7 A g⁻¹ and 5.79 s⁻¹, respectively, at an overpotential of 310 mV. Such a high mass activity is more than three orders of magnitude higher than benchmark OER electrocatalysts, such as IrO₂ and RuO₂. This work presents a sustainable approach toward the development of highly efficient electrocatalysts with largely reduced mass loading of precious elements.

KEYWORDS Oxygen evolution reaction; Multiphase hybrid; Two-dimensional nanomaterials; Rare-earth oxides; Ionic layer epitaxy



1 Introduction

Today, oxygen evolution reaction (OER) is becoming an increasingly important process in many clean, renewable, and scalable electrochemical energy conversion and storage systems, such as fuel cells, electrochemical water splitting, solar fuel generation, and metal–air batteries [1–10]. To lead the exciting experimental developments of these energy systems to industrial applications, materials that catalyze OER with a high mass activity, a low overpotential, and a robust kinetic are highly desired [11–13]. Recent breakthroughs to lower the overpotential have revealed a large number of promising OER catalysts including carbon-based materials (e.g., graphene, CNT, and g-C₃N₄), and alternatives of transition metals (e.g., Mn, Co, Ni, and Fe) [13–25]. However, the low mass activity, high cost, and complicated fabrication procedure are still hindering scalable implementations of these materials in replacing the benchmark IrO₂ and RuO₂ that have high cost and limited supply [5, 13, 15, 26]. Therefore, development of high mass activity and high-efficient OER electrocatalysts based on earth-abundant elements is crucial and urgently needed to drive today's advanced energy technologies forward [3, 13, 17].

Recently, rare-earth metal-based materials, especially rare-earth metal oxides, are found promising for a wide range of catalytic applications, including steam methane reforming, photocatalysis, water–gas shift reactions, thermochemical water splitting, and organocatalytic reactions [27]. Among them, La₂O₃ is showing a great potential as an alternative electrocatalytic candidate owing to its multiple oxidation states, excellent chemical stability, and low toxicity [27–29]. However, the electrochemical catalytic performance of commercial La₂O₃ powders is far from satisfaction mainly because of their low ratio of active catalytic sites and poor conductivity [5, 30]. Morphological alterations, particularly the two-dimensional (2D) geometry with just one or a few atomic layers, are a promising solution for improving the catalytic performance due to the abundant active sites, delocalized spin states, high electrical conductivity, and low mass loading [31, 32]. Furthermore, hybridization of 2D materials with nanoparticles (NPs) could offer an even greater boost to the electrochemical properties by combining the structural and electronic advantages of different morphologies [30, 33–35], such

as lower the over potential of hydrogen evolution reaction (HER), and even raise the reversible capacity of lithium-ion batteries [36, 37]. Inspired by these previous advancements, reducing the dimension of La₂O₃ to a 2D structure and hybridizing with electrochemical active NPs may be a promising route leading La₂O₃ toward a high-performance electrochemical catalytic material in many energy conversion and storage systems. Based on this rationale, here, we report an ultrathin La₂O₃ nanosheets–NP hybrid structure (La₂O₃@NP-NS). The La₂O₃@NP-NS exhibited excellent OER performance with a low overpotential of 310 mV at 10 mA cm⁻² and a very small Tafel slope of 43.1 mV dec⁻¹ when the thickness of the La₂O₃ nanosheets was reduced to 2.27 nm. As a result of its uniform and ultrasmall thickness, they achieved a high mass activity, which was more than three orders of magnitude higher than benchmark IrO₂ and RuO₂, and five orders of magnitude higher than commercial La₂O₃ powder at the overpotential of 310 mV. This development presents an effective and scalable approach toward high-performance OER catalysts with a minimal use of precious rare-earth elements.

2 Experimental

2.1 Synthesis of 2.27-nm La₂O₃@NP-NS

Firstly, La₂O₃ nanosheets were synthesized by ionic layer epitaxy (ILE). In a typical process, 15 mL aqueous solution containing 0.2 mM La(NO₃)₃ and 2 mM hexamethylenetetramine (HMTA) was prepared in a glass vial with a 4.5 cm² opening area. After standing the aqueous solution in air for 40 min, 10 μL chloroform solution of mixed surfactants containing (~0.9 vol %) stearic acid (SA) solution and (~0.1 vol %) oleylamine (OAM) solution was spread on the water surface. The SA solution had a concentration of 1.8 mol L⁻¹. The OAM solution had a concentration of 1.8 mol L⁻¹ and mixed with hydrochloric acid to a concentration of 0.01 mol L⁻¹. This two-layer solution was exposed in atmosphere for 10 min to allow the chloroform to evaporate. Subsequently, the glass vial was screw-capped and placed in a 45 °C convection oven for 5 h for La₂O₃ nanosheets to grow. The La₂O₃ nanosheets were directly scooped onto a substrate and dried naturally in air. Secondly, the La₂O₃ nanosheets on the substrate

were annealed in Ar atmosphere at 400 °C for 1 h, yielding 2.27-nm La₂O₃@NP-NS for further characterization and properties measurements. Si substrates were used for SEM and AFM characterization. Fifty-nanometer Au-coated Si substrates were used for XPS characterization. Holy carbon grids were used for TEM. The ILE-synthesized La₂O₃ samples were firstly transferred on holy carbon grids for TEM measurement, and these ILE-synthesized La₂O₃ samples on TEM grid were then annealed in Ar atmosphere for TEM analysis of La₂O₃@NP-NS samples. Fluorine-doped tin oxide (FTO) glass substrates were used for electrocatalytic OER measurement.

2.2 Synthesis of Thicker La₂O₃ Nanosheets

La₂O₃@NP-NS (8.68 nm) and La₂O₃ (28.26 nm) nanosheets were synthesized through the same ILE and subsequent Ar annealing process as described above, where the reaction temperature during ILE process was increased to 60 and 80 °C for 5 h, respectively.

2.3 La₂O₃ Nanoparticles–FTO Electrode

La₂O₃ NPs were synthesized under the same ILE process at a temperature of 80 °C for 5 h. The NPs were collected from the bulk solution by centrifuge. As-received La₂O₃ NPs (2 mg) was ultrasonically dispersed into an ethanol solution to form a catalytic ink. This ethanol solution contained 0.5 mL of ethanol and 50 μL of a Nafion[®] solution (5 wt% in aliphatic alcohol from Sigma-Aldrich). La₂O₃ NPs–FTO electrode was prepared by depositing 3 μL catalytic ink onto surface of FTO, which corresponded to a loading of 0.02 mg of catalyst per cm². The ink was dried at room temperature in air.

2.4 IrO₂–FTO, RuO₂–FTO, and Commercial La₂O₃–FTO Electrode

Commercial IrO₂, RuO₂, or La₂O₃ powder (10 mg, from Sigma-Aldrich) was ultrasonically dispersed into an ethanol solution to form a catalytic ink. This ethanol solution contained 0.5 mL of ethanol and 50 μL of a Nafion[®] solution (5 wt% in aliphatic alcohol from Sigma-Aldrich). IrO₂–FTO or RuO₂–FTO electrode was prepared by depositing 10 μL catalytic ink onto surface of FTO, which corresponded to a

loading of 0.926 mg of catalyst per cm². The ink was dried at room temperature in air.

2.5 Material Characterizations

A Zeiss LEO 1530 field-emission scanning electron microscope was used to characterize morphologies of La₂O₃ nanosheets. The atomic force microscopy (AFM) tomography images were obtained using an XE-70 Park System. X-ray photoelectron spectroscopy (XPS) spectrum was obtained by a Thermo Scientific K-alpha XPS instrument with a 100 μm spot size, with the flood gun turned on during the measurements. The crystal structure was investigated by A FEI TF30 transmission electron microscopy (TEM) operated at 300 kV.

2.6 Electrochemical Testing

The OER performance was done on an Autolab PGSTAT302N workstation with a standard three-electrode system, which was used to record the catalytic activity of samples in N₂-saturated 1 M NaOH solution (PH= 13.6). The as-prepared La₂O₃@NP-NS on FTO substrate directly acted as the working electrode (mass loading: 0.0014 mg cm⁻² for 2.27-nm La₂O₃@NP-NC, 0.0055 mg cm⁻² for 8.68 nm La₂O₃@NP-NS, and 0.0186 mg cm⁻² for 28.3 nm La₂O₃ nanosheets), the saturated Ag/AgCl was used as the reference electrode, and a platinum wire was used as the counter electrode. The OER polarization curves were recorded at a scan rate of 5 mV s⁻¹ 1 M NaOH solution after purging with nitrogen for 30 min. All the potentials in this study are reported versus RHE. The electrochemical impedance spectroscopy (EIS) measurements were collected with frequencies ranging from 10 kHz to 0.1 Hz under an AC potential of 5 mV. For the stability test, the electrode was firstly activated in N₂-saturated 1 M NaOH solution for 25 cycles and then recorded the first CV polarization curves at a sweep rate of 5 mV s⁻¹. After continuous tests for 11 or 27 h, the final CV polarization curves were recorded at a sweep rate of 5 mV s⁻¹ again. The overpotential (η) was calculated by the following relationship: $\eta = E$ (vs. Ag/AgCl) + 0.21 V + 0.0592 × pH – 1.23 V – iR_u , where i is the current, and R_u equals to the R_{ct} value resolved from Nyquist plots. TOF can be calculated as $TOF = (j \times A) / (4 \times F \times n)$, where j (mA cm⁻²) is the current density at a particular overpotential, A is the

area of the working electrode, F is the Faraday constant ($96,500 \text{ C mol}^{-1}$), and n is the number of moles of the active materials. The mass activity can be calculated as: mass activity = j/m , where m is the mass loading of the working electrode (mg cm^{-2}) and j is the measured current density (mA cm^{-2}) at given potential. Electrochemical active surface area (ECSA) was calculated by the equation $\text{ECSA} = C_{\text{dl}}/C_s$, where C_s is the specific capacitance of the samples. Herein, C_s of 0.04 mF cm^{-2} was used according to the previously reported value of metal oxide/hydroxides in NaOH solution [38]. C_{dl} was calculated from the slope of the line in the plot of capacitive current density (j_{dl}) versus scan rates ν (V/s). A is surface area of electrode. $j_{\text{dl}} = (C_{\text{dl}} \times \nu)/A$, and cyclic voltammograms (CVs) were recorded in the double-layer regime (1 M NaOH, 0 to 0.1 V vs. Ag|AgCl) by varying scan rates ranging from 10 to 60 mV s^{-1} .

3 Results and Discussion

The La_2O_3 @NP-NS was synthesized by a solution-based ILE process at the water–air interface under a monolayer of mixed surfactants followed by Ar annealing as shown in Fig. S1. Firstly, the La_2O_3 nanosheets were synthesized by ILE process. This aqueous solution contained 0.2 mM $\text{La}(\text{NO}_3)_3$ and 2 mM HMTA as precursors and had a weak alkaline environment because HMTA decomposed into HCHO and ammonium hydroxide at 45 °C. Under weak alkaline conditions, La^{3+} could easily combine with OH^- to form $\text{Ce}(\text{OH})_3$, which would then be dehydrated to La_2O_3 after subsequent drying in the air. The as-synthesized La_2O_3 nanosheets exhibited a hexagonal shape with a bi-model size distribution (Fig. 1a). All the nanosheets appeared to be hexagonal, and their size exhibited a bi-model distribution. Larger nanosheets had a size of $15.7 \pm 2.4 \mu\text{m}$; while small ones were mostly around $1.8 \pm 0.9 \mu\text{m}$. This bi-model distribution was possibly due to the ILE mechanism where small nuclei or nanosheets tend to merge into big nanosheets driven by the reduction in surface energy [39]. Directly transferred La_2O_3 nanosheets covered almost the entire Si substrate surface, and no overlap was observed. AFM topography image (Fig. 1b) revealed that all the nanosheets had a very flat surface (surface roughness of 0.5 nm) with a uniform thickness of 2.31 nm (Fig. 1c). XPS analysis (Fig. S2a) showed that the spin–orbit splitting between $\text{La } 3d_{3/2}$ and $\text{La } 3d_{5/2}$ peaks was 16.8 eV, which was the typical characteristic

of La_2O_3 phase [40–42]. TEM image (Fig. S3) revealed the amorphous structure of as-synthesized La_2O_3 nanosheets. In addition, the thickness of the La_2O_3 nanosheets could be tuned by the synthesis temperatures (Fig. S4). As the synthesis temperature rose from 45 to 60, and 90 °C, the nanosheet thickness increased from 2.31 to 8.73, and 28.30 nm, respectively. All the nanosheets still exhibited similar bi-model hexagonal geometry with excellent surface flatness.

The La_2O_3 @NP-NS was obtained by annealing the as-synthesized La_2O_3 nanosheets in Ar at 400 °C for 1 h (Fig. S1). As shown by a low-magnification SEM image (Fig. 1d), the La_2O_3 @NP-NS exhibited the same morphology and size distribution after the annealing. No cracks or dissociations could be observed. AFM image discovered that numerous fine NPs appeared on the nanosheet surface with a fairly uniform distribution (Fig. 1e). The surface density of the NPs was about 20 per μm^2 . Enlarged AFM image (Fig. 1f) showed that these NPs mostly had a spherical shape, anchored on the surface of nanosheets, and their sizes were in the range from 29 to 46 nm. The nanosheets still had a thickness of 2.27 nm (Fig. 1g), while the NPs were extruded above surface of nanosheet surface from 1 to 3 nm.

The elemental composition and chemical state of La_2O_3 @NP-NS were then investigated by XPS. As shown in Fig. 1h, the $\text{La } 3d$ spectrum was split into $\text{La } 3d_{3/2}$ and $\text{La } 3d_{5/2}$, located at 852.7 and 835.9 eV, respectively, with two satellite peaks at 856.3 and 839.4 eV, indicating the existence of La^{3+} . Typical characteristic 16.8 eV between $\text{La } 3d_{3/2}$ and $\text{La } 3d_{5/2}$ peaks of La_2O_3 phase was still shown. In addition, the $\text{O } 1s$ peaks at 532.5 and 530.5 eV could be assigned to the La – O bond in the La_2O_3 phase (Fig. 1i) [28, 43], further evidencing that the composition of La_2O_3 @NP-NS was La_2O_3 . By comparing the XPS results before and after Ar annealing, no obvious change was found for both La peaks and O peaks, suggesting both nanosheet and NPs had the same La_2O_3 chemistry. Thicker La_2O_3 nanosheets (8.73 and 28.30 nm) also showed the same morphology change after annealing (Fig. S5) with the same chemistry (Fig. S6). There was also a large quantity of NPs appeared uniformly on the surface of 8.68 nm nanosheets (Fig. S5c and inset). However, no NPs could be observed from the 28.26 nm nanosheets, possibly because the large thickness completely buried the NPs inside (Fig. S5d).

The crystal structure of La_2O_3 is hexagonal structure. There are two types of oxygen atoms in each hexagonal unit cell: One (denoted O4) is coordinated with four La atoms

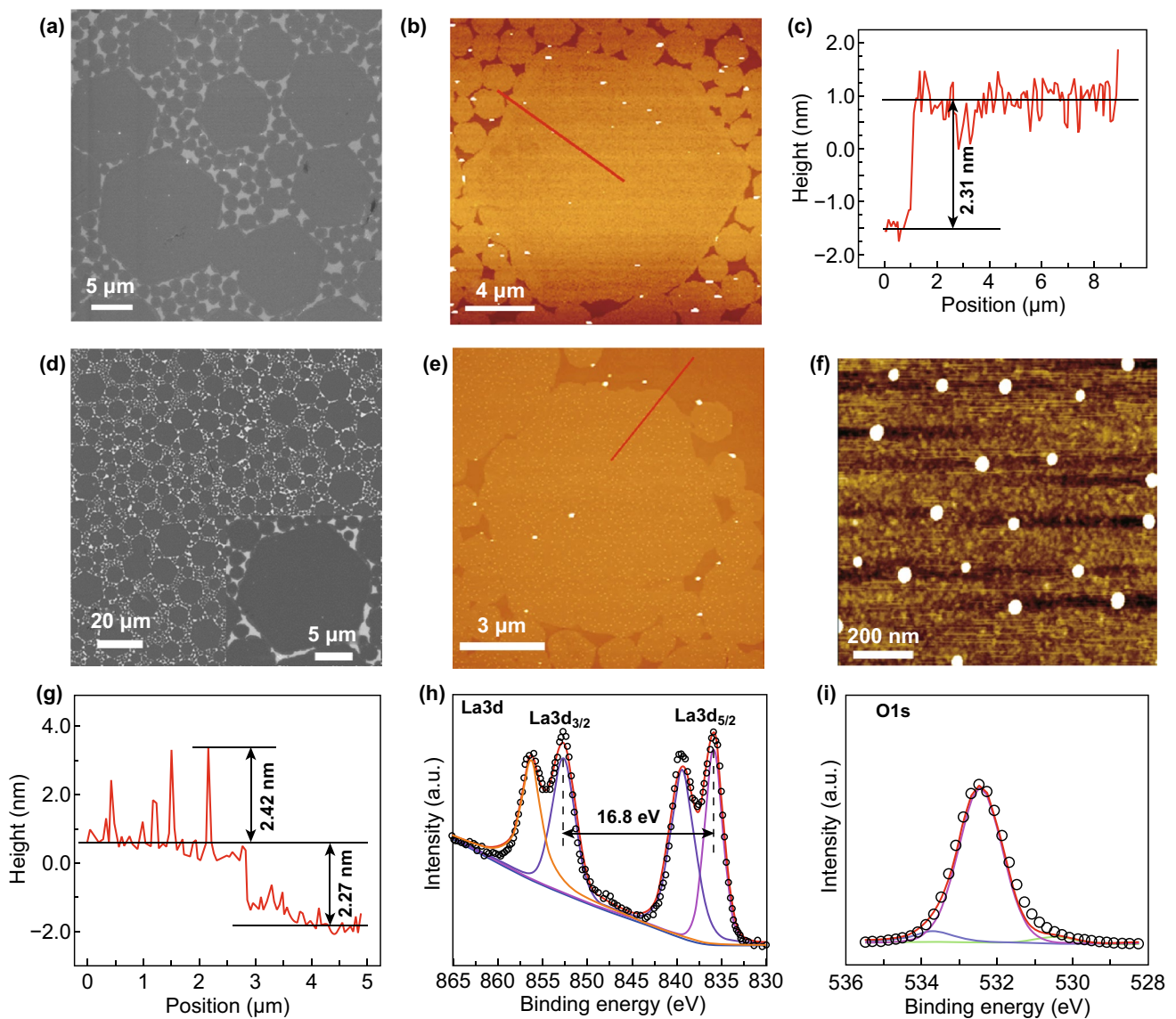


Figure 1 Morphology and elemental information of La_2O_3 nanosheets and $\text{La}_2\text{O}_3@NP\text{-NS}$. **a** SEM image of La_2O_3 nanosheets on a silicon substrate. **b** AFM topography scan of La_2O_3 nanosheets. **c** Height profiles derived from the red line in **b**. **d** SEM image of $\text{La}_2\text{O}_3@NP\text{-NS}$ on a silicon substrate at low magnification. Inset is high-magnification SEM image of $\text{La}_2\text{O}_3@NP\text{-NS}$. **e** AFM topography scan of $\text{La}_2\text{O}_3@NP\text{-NS}$. **f** High-magnification AFM topography scan of $\text{La}_2\text{O}_3@NP\text{-NS}$. **g** Height profiles derived from the red line in **e**. **h** La 3d and **i** O 1s XPS spectrum of $\text{La}_2\text{O}_3@NP\text{-NS}$

and the other (denoted O6) with six La atoms, while each La atom is coordinated with seven oxygen atoms: Four of them are O4, and the other three are O6 [44]. The crystal structure of $\text{La}_2\text{O}_3@NP\text{-NS}$ was then studied by TEM. Low-magnification TEM image showed a uniform contrast of all hexagonal $\text{La}_2\text{O}_3@NP\text{-NS}$ (Fig. 2a). A zoom-in image (Fig. 2b) at the corner area revealed that the angle between two adjacent edges of nanosheets was about 120° as marked by the dotted lines. The overall nanosheet showed a uniform

contrast but no clearly observable lattices. The very front of all the edges was mostly wavy. Both features indicate amorphous or polycrystallinity structure of the nanosheet. Selected area electron diffraction (SAED) pattern collected from multiple nanosheets surfaces (Fig. 2c) showed wide and diffusive diffraction rings, suggesting that the $\text{La}_2\text{O}_3@NP\text{-NS}$ indeed had a significant amount of an amorphous phase with small crystallites. The spread of interplanar spacing measured from the ring diameters was associated with

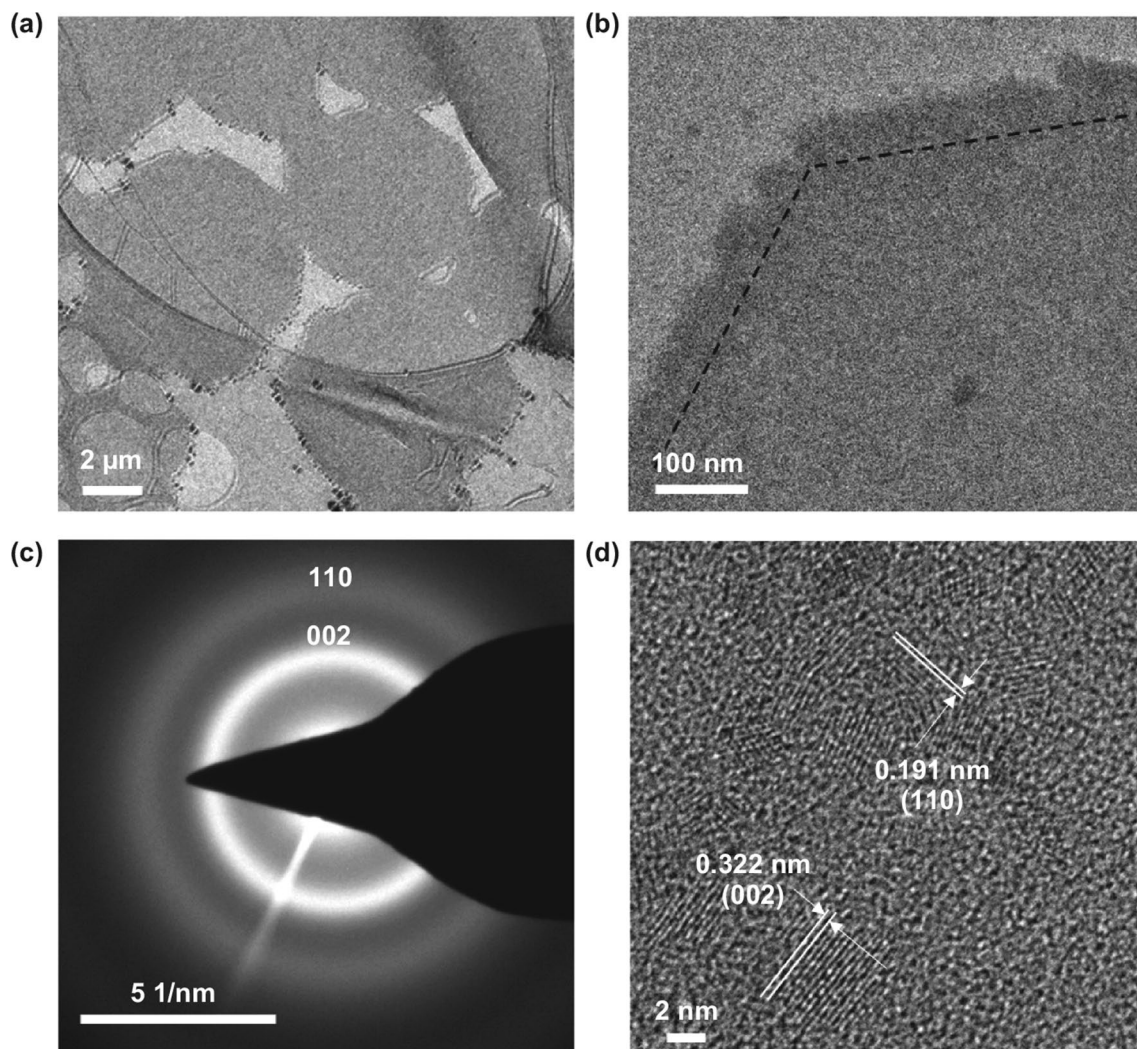


Figure 2 Structural characterization of 2.27-nm La_2O_3 @NP-NS. **a** TEM image of hexagonal La_2O_3 @NP-NS on a holey carbon TEM grid. **b** TEM image taken from a corner of a large La_2O_3 @NP-NS. **c** SAED pattern of La_2O_3 @NP-NS. **d** HRTEM image of La_2O_3 @NP-NS

the (002) and (110) planes of hexagonal-structured La_2O_3 , confirming its polycrystalline structure [45]. HRTEM image (Fig. 2d) revealed the NP-NS hybrid structure. All the NPs had a crystalline phase with a size ranging from 6 to 32 nm, in good agreement with AFM observations. The interplanar spacing from the NPs was identified to be 0.324 and 0.192 nm, corresponding to the (002) and (110) planes of hexagonal La_2O_3 [45]. No crystal lattices could be observed from the nanosheet region, suggesting the nanosheet was still remained as an amorphous phase providing a uniform matrix supporting the crystalline NPs. These structure analyses indicated that during annealing in the Ar, the amorphous La_2O_3 nanosheets could firstly begin to crystallize and

many tiny crystalline domains would thus form in the largely amorphous nanosheets; as the annealing time progressed, these tiny crystalline domains would then grow in both lateral and vertical size, gradually be extruded above surface of nanosheets and eventually grow into nanoparticles embedded in the nanosheets. Besides, Fig. S7 reveals the structure of crystallites distributed in the amorphous nanosheets region for 28.26 nm nanosheets, which confirmed that large thickness completely buried the NPs inside the 28.26 nm nanosheets after annealing.

To investigate the electrocatalytic performance of La_2O_3 @NP-NS for OER, the OER polarization curves of 2.27-nm La_2O_3 @NP-NS on FTO substrate were probed in

alkaline electrolyte (1 M NaOH). Through XPS analysis (Fig. S8), the La_2O_3 @NP-NS grown on FTO substrate has the same La_2O_3 chemistry with that on Si substrate. As a comparison, the OER activities of thicker La_2O_3 @NP-NS nanosheets, La_2O_3 NPs, commercial IrO_2 , RuO_2 , and La_2O_3 powders were also characterized under the same conditions. As shown in Fig. 3a, the 2.27-nm La_2O_3 @NP-NS exhibited the smallest overpotential of 310 mV at a current density of 10 mA cm^{-2} . Thicker nanosheets exhibited higher overpotential of 343 and 429 mV for 8.68 and 28.26 nm nanosheets, respectively. The very low overpotential from the 2.27-nm nanosheets outperformed the state-of-the-art noble metals oxides, e.g., IrO_2 (343 mV) and RuO_2 (408 mV) at the same current density. Besides, commercial La_2O_3 powders exhibited a very high overpotential of 1330 mV (Fig. S12a), confirming the significant morphological advantage of the 2D nanosheets.

Tafel plots from La_2O_3 @NP-NS with different thicknesses and other comparison samples were then characterized to evaluate and compare their reaction rate constants and electrocatalytic kinetics. As illustrated in Fig. 3b, Tafel slope of the 2.27-nm La_2O_3 @NP-NS was calculated to be only 43.1 mV dec^{-1} , significantly lower than that of thicker nanosheets (Table 1). In addition, the low Tafel slope of the 2.27-nm La_2O_3 @NP-NS was also superior to the benchmark IrO_2 , RuO_2 (Table 1), and some recently reported novel OER catalysts, such as $\text{CeO}_x/\text{CoO}_x$ (66 mV dec^{-1}) [46], NiO/CoN PINWs (44.5 mV dec^{-1}) [47], $\text{Co}_x\text{Ni}_{1-x}\text{Fe}_2\text{O}_4$ (46.4 mV dec^{-1}) [48], $\text{Ni}_{0.75}\text{V}_{0.25}\text{-LDH}$ (50 mV dec^{-1}) [49], and $\text{ZnCo}_2\text{O}_4/\text{N-CNT}$ (70.6 mV dec^{-1}) [50]. Such a low Tafel slope value from the 2.27-nm La_2O_3 @NP-NS suggests that the reaction rate constant and charge transfer rate were greatly improved as the nanosheets thickness reduced to the ultrathin nanometer scale. These improved reaction rate constant and charge transfer rate could be attributed to 2D electron gas, which was possibly presented at the interface of NP-NS in the 2D La_2O_3 nanohybrids and contributed to their improved electrocatalytic properties by enhancing electron mobility and conductivity of nanohybrids [51].

EIS was further investigated to reveal the charge transfer resistance (R_{ct}) of La_2O_3 @NP-NS and to understand its role in the enhanced OER performance. As shown in Fig. 3c, Nyquist plot of the 2.27-nm La_2O_3 @NP-NS had a much smaller semicircle compared to thicker nanosheets. From the Nyquist plots, R_{ct} of the 2.27-nm La_2O_3 @NP-NS was found only about 38Ω (Fig. 3d), which was lower than those of

thicker nanosheets, La_2O_3 nanoparticles, and RuO_2 powders, and only slightly larger than that of IrO_2 powder (Table 1). The largely reduced R_{ct} was possibly due to the greatly shortened charge transfer distance between the catalysis surface and substrate as a result of the ultrasmall thickness [30].

One major advantage of the ultrathin nanosheet for electrochemical catalysis is its ultrahigh ratio of active catalytic sites. Figure 3e demonstrates the calculated mass activity plots. A remarkable mass activity of 6666.7 A g^{-1} was obtained from the 2.27-nm La_2O_3 @NP-NS at overpotential 310 mV. It was one to more than two orders of magnitudes higher than other morphologies of La_2O_3 (Table 1) at the same overpotential. Such a large mass activity evidenced the unique merit of the ultrathin 2D geometry, which involved an extremely low mass and rendered considerably large exposed surface area allowing a very high ratio of catalytically active sites within the material. Besides, the NPs extruded above surface of nanosheet should also contribute to high ratio of active catalytic sites. Furthermore, this mass activity was three orders of magnitude higher than benchmark IrO_2 (4.4 A g^{-1}) and RuO_2 (2.05 A g^{-1}), and five orders of magnitude higher than commercial La_2O_3 at the same overpotential, suggesting that our 2.27-nm La_2O_3 @NP-NS could be very competitive electrochemical catalysts for industrial applications. The electrochemical active surface area (ECSA) was also calculated based on cyclic voltammograms curves (Fig. S13) to estimate the exposure of active sites. As shown in Fig. S14, the values of ECSA for 2.27 (0.025) and 8.68 nm (0.021 cm^2) La_2O_3 @NP-NS were close and about twice as much as 28.26 nm La_2O_3 nanosheets, which could be because the NPs extruded above surface of nanosheet also contributed to active catalytic sites. Moreover, the ECSA value of commercial La_2O_3 powder (0.31 cm^2) was much bigger than that of 2.27-nm La_2O_3 @NP-NS, but it exhibited poor OER performance, which could attribute to good conductivity caused by ultrathin thickness, but poor intrinsic OER catalytic activity of commercial La_2O_3 powder [32].

The turnover frequency (TOF) was another critical OER parameter that evaluates the number of O_2 molecules formed per active metal site per second. As shown in Fig. 3f, the TOF of 2.27-nm La_2O_3 @NP-NS reached up to 5.79 s^{-1} at the overpotential of 310 mV, which was about one to five orders of magnitudes higher than those comparison samples (Table 1). The extremely high TOF could be attributed to the largely facilitated electron transfer from the ultrathin nanosheet to the FTO substrate as well as the very large



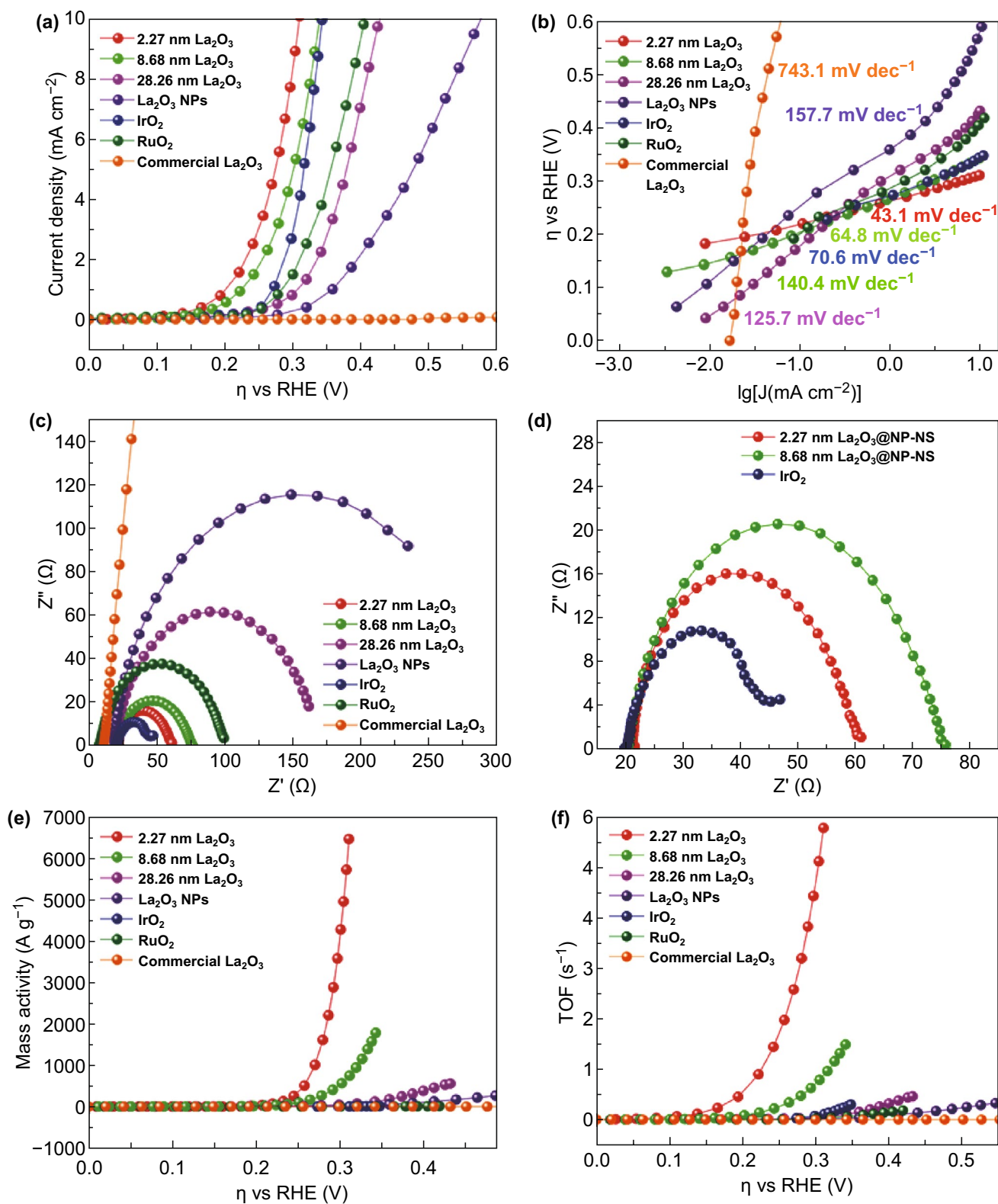


Figure 3 Electrocatalytic OER performance comparison of 2.27-nm La_2O_3 @NP-NS (red), 8.68 nm La_2O_3 @NP-NS (green), and 28.26 nm La_2O_3 nanosheets (magenta), ILE synthesized La_2O_3 NPs (violet), commercial IrO_2 (blue), RuO_2 (olive), and La_2O_3 powder (orange). **a** OER polarization curves measured in 1 M NaOH solution. **b** Tafel plots. **c** Nyquist plots measured in 1 M NaOH solution at a potential of 310 mV versus RHE. **d** Enlarged Nyquist plots. **e** Mass activity determined from current density as a function of η . **f** TOF determined from j as a function of η . (Color figure online)

Table 1 OER performance comparison between the samples synthesized in this work

Samples	Over potential (mV) @ 10 mA cm ⁻²	Tafel slope (mV dec ⁻¹)	R _{ct} (Ω)	Mass activity (A g ⁻¹) @ 310 mV	TOF (s ⁻¹) @ 310 mV
2.27 nm La ₂ O ₃ @NP-NS	310	43.1	38	6666.7	5.79
8.68 nm La ₂ O ₃ @NP-NS	343	64.8	57	713	0.88
28.26 nm La ₂ O ₃	429	125.7	143	51	0.055
ILE synthesized La ₂ O ₃ NPs	580	157.7	272	17.4	0.012
Commercial IrO ₂ powder	343	70.4	25.4	4.4	0.0026
Commercial RuO ₂ powder	408	140.6	89.6	2.05	7.23 × 10 ⁻⁴
Commercial La ₂ O ₃ powder	1330	743.1	47,598	0.048	4.87 × 10 ⁻⁵

mass activity. The significant enhancement in both mass activity and TOF confirmed the ultrathin La₂O₃@NP-NS could provide a highly efficient OER performance with a minimal use of rare-earth elements.

To further figure out the origin of the active sites for La₂O₃@NP-NS, electrocatalytic OER performances of 2.31 nm La₂O₃ nanosheets before Ar annealing were also investigated and compared to that of 2.27-nm La₂O₃@NP-NS, as shown in Fig. S15. It was found that the OER performance of 2.31 nm La₂O₃ nanosheets was greatly enhanced after Ar annealing at 400 °C for one hour. The overpotential was shifted toward a lower overpotential by 59 mV after annealing, with corresponding Tafel slope reducing from 95.4 to 43.1 mV dec⁻¹. Moreover, mass activity increased greatly from 3238 to 6666.7 A g⁻¹. All these results confirmed that the catalytic active sites of La₂O₃@NP-NS were attributed to precipitated La₂O₃ nanoparticles after Ar annealing, as well as ultrathin nanosheet region.

To better understand the OER performance of the ultrathin La₂O₃@NP-NS hybrid structure, the OER characteristic data (e.g., overpotential, mass loading, mass activity, and TOF) from other representative state-of-the-art OER catalysts were collected and compared (Table S1). While the overpotential from this work was generally superior or at least comparable to those Ni-, Co-, Mn-, or Fe-based OER catalysts, the biggest advantage of our nanosheets was obviously the extremely large mass activity. As a result of the ultrasmall thickness and uniform coverage over a large area, the mass loading of our La₂O₃@NP-NS was only 0.0014 mg cm⁻², which brought the mass activity to more than three orders of magnitude higher than typical OER catalysts such as CoFe-LDHs [52], Co₃O₄ [53], and Ni@NC [54].

For nanostructured catalysts, the stability is vital due to the largely increased atom mobility on nanoscale surfaces.

Impressively, the ultrathin La₂O₃@NP-NS exhibited superb stability with 90% retention of the initial current density after 11 h of continuous OER operation in 1 M NaOH at $\eta = 345$ mV (Fig. 4a). SEM image showed that before (Fig. S16a) and after (Fig. S16b) the 11-hour continuous OER test, the coverage of nanosheets was nearly the same and no obvious damage could be observed; XPS analysis (Fig. S17a) revealed that the La₂O₃@NP-NS almost retained their chemistry. In addition, the OER polarization curve showed

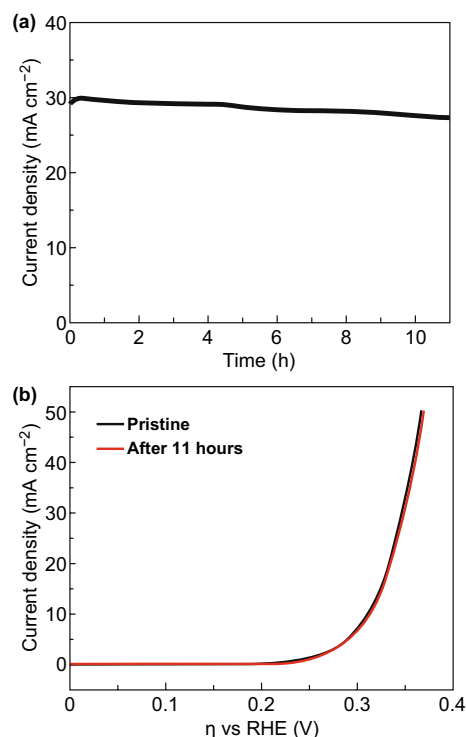


Figure 4 Electrochemical stability tests. **a** Current density measured at $\eta = 345$ mV (vs. RHE) as a function of time. **b** OER polarization curves of 2.27-nm La₂O₃@NP-NS before and after 11-h OER

only a slight decay after the OER test (Fig. 4b), which confirmed appropriate function of the $\text{La}_2\text{O}_3@\text{NP-NS}$ coating during the long-term operation. The high stability of the ultrathin $\text{La}_2\text{O}_3@\text{NP-NS}$ nanosheets might be attributed to the amorphous–crystallites hybrid structure. The amorphous La_2O_3 matrix could stabilize the small La_2O_3 NPs from aggregation by limiting the atomic diffusion, and on the other hand, the anchored La_2O_3 NPs could act as spacers to prevent the 2D La_2O_3 nanosheets from restacking during electrochemical cycling [37, 51]. Besides, the stability test of 2.27-nm $\text{La}_2\text{O}_3@\text{NP-NS}$ with longer time was also investigated. As shown in Fig. S18a, the stability test survived for 27 h because there was a sharp decrease in the current density at the time range from 25 to 27 h. SEM images (Fig. S18c, d) revealed that the majority of $\text{La}_2\text{O}_3@\text{NP-NS}$ fell off and only part of samples was adsorbed on the FTO substrate after 27 h stability test, which resulted in the decrease in current density during stability test and confirmed that OER performance truly came from the ultrathin $\text{La}_2\text{O}_3@\text{NP-NS}$. Long-hour immersion in alkaline solution may weaken the adsorption of $\text{La}_2\text{O}_3@\text{NP-NS}$ on FTO substrate, which could cause the $\text{La}_2\text{O}_3@\text{NP-NS}$ to fall off from the substrate during stability test.

4 Conclusions

In conclusion, $\text{La}_2\text{O}_3@\text{NP-NS}$ was synthesized by a facile solution-based ILE approach in a large area. Outstanding electrocatalytic OER performance was observed when the nanosheet thickness was reduced to 2.27 nm. The 2.27-nm $\text{La}_2\text{O}_3@\text{NP-NS}$ exhibited a low overpotential of 310 mV at 10 mA cm^{-2} , a small Tafel slope of 43.1 mVdec^{-1} , and charge transfer resistance of 38 Ω . A high mass activity of 6666.7 A g^{-1} and turnover frequency of 5.79 s^{-1} were also obtained from the 2.27-nm $\text{La}_2\text{O}_3@\text{NP-NS}$ at an overpotential of 310 mV. This very large mass activity was more than three orders of magnitude higher than benchmark IrO_2 (4.4 A g^{-1}) and RuO_2 (2.05 A g^{-1}), and five orders of magnitude higher than commercial La_2O_3 (0.048 A g^{-1}) at the same overpotential. The nanosheets also exhibited an impressive stability for over 11-h continuous operation with only 10% decay. The excellent electrocatalytic OER activity could be attributed to the ultrathin thickness that enables overall short out-of-plane charge diffusion length and facilitated electron transfer from the nanosheet surface to

the substrate. This development opens up a promising strategy for the development of highly efficient electrocatalysts with largely reduced loading of catalytic materials. Creating the hybrid ultrathin 2D morphologies could potentially become an ultimate solution to preserve precious elements in advanced and sustainable electrocatalyst design in many energy and environmental systems.

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References

1. S. Chu, A. Majumdar, Opportunities and challenges for a sustainable energy future. *Nature* **488**, 294–302 (2012). <https://doi.org/10.1038/nature11475>
2. M. Dresselhaus, I. Thomas, Alternative energy technologies. *Nature* **414**, 332–337 (2001). <https://doi.org/10.1201/b17893>
3. Y.Y. Wang, C. Xie, Z.Y. Zhang, D.D. Liu, R. Chen et al., In situ exfoliated, n-doped, and edge-rich ultrathin layered double hydroxides nanosheets for oxygen evolution reaction. *Adv. Funct. Mater.* **28**, 1703363 (2018). <https://doi.org/10.1002/adfm.201703363>
4. Y.S. Jin, H.T. Wang, J.J. Li, X. Yue, Y.J. Han et al., Porous MoO_2 nanosheets as non-noble bifunctional electrocatalysts for overall water splitting. *Adv. Mater.* **28**, 3785–3790 (2016). <https://doi.org/10.1002/adma.201506314>
5. M. Tahir, L. Pan, F. Idrees, X. Zhang, L. Wang et al., Electrocatalytic oxygen evolution reaction for energy conversion and storage: a comprehensive review. *Nano Energy* **37**, 136–157 (2017). <https://doi.org/10.1016/j.nanoen.2017.05.022>

6. Y.C. Mao, C. Ning, N. Zhang, Y. Hu, M.Y. Li et al., Enhancing photoelectrochemical performance of TiO₂ nanowires through a facile acid treatment method. *J. Electrochem. Soc.* **165**, 799–803 (2018). <https://doi.org/10.1149/2.0431813jes>
7. Y.C. Mao, Y.G. Cheng, J.Q. Wang, H. Yang, M.Y. Li et al., Amorphous NiO electrocatalyst overcoated ZnO nanorod photoanodes for enhanced photoelectrochemical performance. *New J. Chem.* **40**, 107–112 (2016). <https://doi.org/10.1039/c5nj01815c>
8. X.T. Wang, T. Ouyang, L. Wang, J.H. Zhong, T. Ma et al., Redox-inert Fe³⁺ in octahedral sites of Co–Fe spinel oxides with enhanced oxygen catalytic activity for rechargeable Zn–air batteries. *Angew. Chem. Int. Ed.* **58**, 13291–13296 (2019). <https://doi.org/10.1002/ange.201907595>
9. T. Ouyang, Y.Q. Ye, C.Y. Wu, K. Xiao, Z.Q. Liu et al., Heterostructures composed of N-doped carbon nanotubes encapsulating Co and β-Mo₂C nanoparticles as bifunctional electrodes for water splitting. *Angew. Chem. Int. Ed.* **58**, 4923–4928 (2019). <https://doi.org/10.1002/anie.201814262>
10. H. Su, X.T. Wang, J.X. Hu, T. Ouyang, K. Xiao et al., Co–Mn spinel supported self-catalysis induced N-doped carbon nanotubes with high efficiency electron transport channels for zinc–air batteries. *J. Mater. Chem. A* **7**, 22307–22313 (2019). <https://doi.org/10.1039/C9TA08064C>
11. D.H. Guo, R.K. Shibuya, C. Akiba, S. Saji, T. Kondo et al., Active sites of nitrogen-doped carbon materials for oxygen reduction reaction clarified using model catalysts. *Science* **351**, 361–365 (2016). <https://doi.org/10.1126/science.aad0832>
12. B.Y. Xia, Y. Yan, N. Li, H.B. Wu, X.W.D. Lou, X. Wang, A metal–organic framework-derived bifunctional oxygen electrocatalyst. *Nat. Energy* **1**, 15006 (2016). <https://doi.org/10.1038/energy.2015.6>
13. W.R. Cheng, X. Zhao, H. Su, F. Tang, W. Che et al., Lattice-strained metal–organic-framework arrays for bifunctional oxygen electrocatalysis. *Nat. Energy* **4**, 115–122 (2019). <https://doi.org/10.1038/s41560-018-0308-8>
14. S. Zheng, L. Zheng, Z. Zhu, J. Chen, J. Kang, Z. Huang, D. Yang, MoS₂ nanosheet arrays rooted on hollow rGO spheres as bifunctional hydrogen evolution catalyst and supercapacitor electrode. *Nano-Micro Lett.* **10**, 62 (2018). <https://doi.org/10.1007/s40820-018-0215-3>
15. T.Y. Ma, S. Dai, M. Jaroniec, S.Z. Qiao, Graphitic carbon nitride nanosheet–carbon nanotube three-dimensional porous composites as high-performance oxygen evolution electrocatalysts. *Angew. Chem. Int. Ed.* **53**, 7281–7285 (2014). <https://doi.org/10.1002/anie.201403946>
16. M.G. Walter, E.L. Warren, J.R. McKone, S.W. Boettcher, Q. Mi et al., Solar water splitting cells. *Chem. Rev.* **110**, 6446–6473 (2010). <https://doi.org/10.1021/cr1002326>
17. D. Chen, C. Chen, Z.M. Baiyee, Z. Shao, F. Ciucci et al., Non-stoichiometric oxides as low-cost and highly-efficient oxygen reduction/evolution catalysts for low-temperature electrochemical devices. *Chem. Rev.* **115**, 9869–9921 (2015). <https://doi.org/10.1021/acs.chemrev.5b00073>
18. B.W. Zhang, K. Jiang, H.T. Wang, S. Hu, Fluoride-induced dynamic surface self-reconstruction produces unexpectedly efficient oxygen-evolution catalyst. *Nano Lett.* **19**, 530–537 (2019). <https://doi.org/10.1021/acs.nanolett.8b04466>
19. R.R. Zhang, Y.C. Zhang, L. Pan, G.Q. Shen, N. Mahmood et al., Engineering cobalt defects in cobalt oxide for highly efficient electrocatalytic oxygen evolution. *ACS Catal.* **8**, 3803–3811 (2018). <https://doi.org/10.1021/acscatal.8b01046>
20. X.N. Li, H. Liu, Z.Z. Chen, Q.M. Wu, Z.Y. Yu et al., Enhancing oxygen evolution efficiency of multiferroic oxides by spintronic and ferroelectric polarization regulation. *Nat. Commun.* **10**, 1409 (2019). <https://doi.org/10.1038/s41467-019-09191-0>
21. Y. Hou, M. Qiu, M.G. Kim, P. Liu, G. Nam et al., Atomically dispersed nickel–nitrogen–sulfur species anchored on porous carbon nanosheets for efficient water oxidation. *Nat. Commun.* **10**, 1392 (2019). <https://doi.org/10.1038/s41467-019-09394-5>
22. Z.F. Huang, J.J. Song, Y.H. Du, S.B. Xi, S. Dou et al., Chemical and structural origin of lattice oxygen oxidation in Co–Zn oxyhydroxide oxygen evolution electrocatalysts. *Nat. Energy* **4**, 329–338 (2019). <https://doi.org/10.1038/s41560-019-0355-9>
23. L. Hu, T.Z. Xiong, R. Liu, Y.W. Hu, Y.C. Mao et al., Co₃O₄@Cu-based conductive metal–organic framework core-shell nanowire electrocatalysts enable efficient low-overall-potential water splitting. *Chem. Eur. J.* **25**, 6575–6583 (2019). <https://doi.org/10.1002/chem.201900045>
24. L. Hu, Y. Hu, R. Liu, Y. Mao, M.S.J.T. Balogun, Y. Tong, Co-based MOF-derived Co/CoN/Co₂P ternary composite embedded in N- and P-doped carbon as bifunctional nanocatalysts for efficient overall water splitting. *Int. J. Hydrogen Energy* **44**, 11402–11410 (2019). <https://doi.org/10.1016/j.ijhydene.2019.03.157>
25. F.X. Wang, L. Hu, R. Liu, H. Yang, T.Z. Xiong et al., Hybrid implanted hybrid hollow nanocube electrocatalyst facilitates efficient hydrogen evolution activity. *J. Mater. Chem. A* **7**, 11150–11159 (2019). <https://doi.org/10.1039/c9ta00931k>
26. J. Suntivich, K.J. May, H.A. Gasteiger, J.B. Goodenough, Y. Shao-Horn, A perovskite oxide optimized for oxygen evolution catalysis from molecular orbital principles. *Science* **334**, 1383–1385 (2011). <https://doi.org/10.1126/science.1212858>
27. T. Montini, M. Melchionna, M. Monai, P. Fornasiero, Fundamentals and catalytic applications of CeO₂-Based materials. *Chem. Rev.* **116**, 5987–6041 (2016). <https://doi.org/10.1021/acs.chemrev.5b00603>
28. J.G. Kang, Y.I. Kim, D.W. Cho, Y. Sohn, Synthesis and physicochemical properties of La(OH)₃ and La₂O₃ nanostructures. *Mat. Sci. Semicon. Proc.* **40**, 737–743 (2015). <https://doi.org/10.1016/j.mssp.2015.07.050>
29. A.A. Yadav, A.C. Lokhande, R.B. Pujaria, J.H. Kimb, C.D. Lokhande, The synthesis of multifunctional porous honey comb-like La₂O₃ thin film for supercapacitor and gas sensor applications. *J. Colloid Interface Sci.* **484**, 51–59 (2016). <https://doi.org/10.1016/j.jcis.2016.08.056>
30. T.A. Shifa, F. Wang, Y. Liu, J. He, Heterostructures based on 2d materials: a versatile platform for efficient catalysis. *Adv. Mater.* **31**(45), 1804828 (2018). <https://doi.org/10.1002/adma.201804828>



31. H. Jin, C. Guo, X. Liu, J. Liu, A. Vasileff et al., Emerging two-dimensional nanomaterials for electrocatalysis. *Chem. Rev.* **118**, 6337–6408 (2018). <https://doi.org/10.1021/acs.chemrev.7b00689>
32. P. Tian, Y.H. Yu, X. Yin, X.D. Wang, A wafer-scale 1 nm Ni(OH)₂ nanosheet with superior electrocatalytic activity for the oxygen evolution reaction. *Nanoscale* **10**, 5054–5059 (2018). <https://doi.org/10.1039/c7nr09042k>
33. D. Deng, K.S. Novoselov, Q. Fu, N. Zheng, Z. Tian et al., Catalysis with two-dimensional materials and their heterostructures. *Nat. Nanotechnol.* **11**, 218–230 (2016). <https://doi.org/10.1038/nnano.2015.340>
34. M. Yu, S. Zhou, Z. Wang, J. Zhao, J. Qiu, Boosting electrocatalytic oxygen evolution by synergistically coupling layered double hydroxide with MXene. *Nano Energy* **44**, 181–190 (2018). <https://doi.org/10.1016/j.nanoen.2017.12.003>
35. C. Tang, H.S. Wang, H.F. Wang, Q. Zhang, G.L. Tian et al., Spatially confined hybridization of nanometer-sized NiFe hydroxides into nitrogen-doped graphene frameworks leading to superior oxygen evolution reactivity. *Adv. Mater.* **27**, 4516–4522 (2015). <https://doi.org/10.1002/adma.201501901>
36. Y. Li, H. Wang, L. Xie, Y. Liang, G. Hong et al., MoS₂ Nanoparticles grown on graphene: an advanced catalyst for the hydrogen evolution reaction. *J. Am. Chem. Soc.* **133**, 7296–7299 (2011). <https://doi.org/10.1021/ja201269b>
37. J. Mujtaba, H.Y. Sun, G.Y. Huang, K. Mølhave, Y.G. Liu et al., Nanoparticle decorated ultrathin porous nanosheets as hierarchical Co₃O₄ nanostructures for lithium ion battery anode materials. *Sci. Rep.* **6**, 20592 (2016). <https://doi.org/10.1038/srep20592>
38. C.L.M. Charles, J. Suho, C.P. Jonas, F.J. Thomas, Benchmarking heterogeneous electrocatalysts for the oxygen evolution reaction. *J. Am. Chem. Soc.* **135**, 16977 (2013). <https://doi.org/10.1021/ja407115p>
39. F. Wang, J.H. Seo, G. Luo, M.B. Starr, Z. Li et al., Nanometre-thick single-crystalline nanosheets grown at the water–air interface. *Nat. Commun.* **7**, 10444 (2016). <https://doi.org/10.1038/ncomms10444>
40. Y. Li, B. Guan, A. Maclennan, Y.F. Hu, D.D. Li et al., Porous waxberry-like MnO₂/La₂O₃ microspheres for high performance asymmetric supercapacitor. *Electrochim. Acta* **241**, 395–405 (2017). <https://doi.org/10.1016/j.electacta.2017.04.175>
41. T. Honma, Y. Benino, T. Fujiwara, T. Komatsu, Electronic polarizability, optical basicity, and interaction parameter of La₂O₃ and related glasses. *J. Appl. Phys.* **91**, 2942–2950 (2002). <https://doi.org/10.1063/1.1436292>
42. C. Yang, H. Fan, S. Qiu, Y. Xi, Y. Fu, Microstructure and dielectric properties of La₂O₃ films prepared by ion beam assistant electron-beam evaporation. *J. Non-Cryst. Solids* **355**, 33–37 (2009). <https://doi.org/10.1016/j.jnoncrysol.2008.09.029>
43. M.F. Sunding, K. Hadidi, S. Diplas, O.M. Løvrvik, T.E. Norby et al., XPS characterisation of in situ treated lanthanum oxide and hydroxide using tailored charge referencing and peak fitting procedures. *J. Electron. Spectrosc.* **184**, 399–409 (2011). <https://doi.org/10.1016/j.elspec.2011.04.002>
44. B. Li, H. Metiu, DFT studies of oxygen vacancies on undoped and doped La₂O₃ surfaces. *J. Phys. Chem. C* **114**, 12234–12244 (2010). <https://doi.org/10.1021/jp103604b>
45. P. Huang, Y. Zhao, J. Zhang, Y. Zhu, Y. Sun, Exploiting shape effects of La₂O₃ nanocatalysts for oxidative coupling of methane reaction. *Nanoscale* **5**, 10844–10848 (2013). <https://doi.org/10.1039/c3nr03617k>
46. J.H. Kim, K. Shin, K. Kawashima, D.H. Youn, J. Lin et al., Enhanced activity promoted by CeO_x on a CoO_x electrocatalyst for the oxygen evolution reaction. *ACS Catal.* **8**, 4257–4265 (2018). <https://doi.org/10.1021/acscatal.8b00820>
47. J. Yin, Y.X. Li, F. Lv, Q.H. Fan, Y.Q. Zhao et al., Porous nanowires as efficient bifunctional catalysts for Zn–air batteries. *ACS Nano* **11**, 2275–2283 (2017). <https://doi.org/10.1021/acsnano.7b00417>
48. V. Maruthapandian, O. Mathankumar, V. Saraswathy, B. Subramanian, O. Muralidharan, Study of the oxygen evolution reaction catalytic behavior of Co_xNi_{1-x}Fe₂O₄ in alkaline medium. *ACS Appl. Mater. Interfaces* **9**, 13132–13141 (2017). <https://doi.org/10.1021/acsami.6b16685>
49. K. Fan, H. Chen, Y. Ji, H. Huang, P.M. Claesson et al., Nickel–vanadium monolayer double hydroxide for efficient electrochemical water oxidation. *Nat. Commun.* **7**, 11981 (2016). <https://doi.org/10.1038/ncomms11981>
50. Z.Q. Liu, H. Cheng, N. Li, T.Y. Ma, Y.Z. Su, ZnCo₂O₄ quantum dots anchored on nitrogen-doped carbon nanotubes as reversible oxygen reduction/evolution electrocatalysts. *Adv. Mater.* **28**, 3777–3784 (2016). <https://doi.org/10.1002/adma.201506197>
51. Y. Liu, C. Ma, Q. Zhang, W. Wang, P. Pan et al., 2D electron gas and oxygen vacancy induced high oxygen evolution performances for advanced Co₃O₄/CeO₂ nanohybrids. *Adv. Mater.* **31**, 1900062 (2019). <https://doi.org/10.1002/adma.201900062>
52. P. Li, M.Y. Wang, X.X. Duan, L.R. Zheng, X.P. Cheng et al., Boosting oxygen evolution of single-atomic ruthenium through electronic coupling with cobalt-iron layered double hydroxides. *Nat. Commun.* **10**, 1711 (2019). <https://doi.org/10.1038/s41467-019-09666-0>
53. Y. Zhang, B. Ouyang, J. Xu, G. Jia, S. Chen et al., Rapid synthesis of cobalt nitride nanowires: highly efficient and low-cost catalysts for oxygen evolution. *Angew. Chem. Int. Ed.* **55**, 8670–8674 (2016). <https://doi.org/10.1002/anie.201604372>
54. J. Ren, M. Antonietti, T.P. Fellingner, Efficient water splitting using a simple Ni/N/C paper electrocatalyst. *Adv. Energy Mater.* **5**, 1401660 (2015). <https://doi.org/10.1002/aenm.201401660>