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## CeO<sub>2</sub> supported high-valence Fe oxide for highly active and stable water oxidation<sup>†</sup>

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Despite the high intrinsic electrocatalytic activity for the oxygen evolution reaction (OER), stable high valence metal ions such as Fe<sup>4+</sup> are very difficult to form in oxide catalysts. In this work, by elaborating a novel Fe<sup>IV</sup>/CeO<sub>2</sub>@NF (nickel foam) catalyst, we stabilized high-valence Fe ions on a CeO<sub>2</sub> support and achieved a record low overpotential of 219 mV to reach the current density of 50 mA cm<sup>-2</sup>. Theoretical calculations revealed that the fluent d-f electron transfer between ultra-small FeO<sub>x</sub> nanoparticles (US-FeO<sub>x</sub>) and CeO<sub>2</sub> guarantees the robust high valence of surface Fe sites, which enables the optimum adsorption and efficient conversions in the OER process. Meanwhile, the electronic modulations induced by the US-FeO<sub>x</sub> also improve the site-to-site electron transfer to lower the reaction energy barriers for excellent OER performance. Moreover, the Fe<sup>IV</sup>/CeO<sub>2</sub>@NF catalyst delivered excellent stability, sustaining a high current density (200 mA cm<sup>-2</sup>) for over 500 h, and the simple preparation method gave access to a large-area electrode (100 cm<sup>2</sup>), paving the way for large-scale hydrogen production by water splitting.

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### Broader context

Water electrolysis-based hydrogen production in a sustainable fashion is one of the most promising ways to meet the global energy demand of a green future. The development of highly active and stable water oxidation catalyst is the key for large-scale application of water electrolysis because of the kinetically sluggish four-electron transfer process of oxygen evolution reaction (OER). Despite the high intrinsic electrocatalytic activity for the OER, stable high valence metal ions such as Fe<sup>4+</sup> are very difficult to form in oxide catalysts. Herein we successfully stabilized high-valence Fe ions on Ce<sup>4+/3+</sup>O<sub>2</sub> support and achieved superior OER performance. The novel Fe<sup>IV</sup>/CeO<sub>2</sub>@NF (nickel foam) catalyst achieved a record low overpotential of 219 mV to reach the current density of 50 mA cm<sup>-2</sup> and delivered excellent stability of sustaining a high current density (200 mA cm<sup>-2</sup>) for over 500 h. Theoretical calculations revealed that the fluent d-f electron transfer between ultra-small FeO<sub>x</sub> nanoparticles (US-FeO<sub>x</sub>) and CeO<sub>2</sub> guarantees the robust high valence of surface Fe sites, which enables the optimum adsorption and efficient conversions in the OER process. Moreover, the simple preparation method gave access to a large-area catalyst electrode (100 cm<sup>2</sup>), paving the way for large-scale hydrogen production by water splitting. This work sets the stage for designing and preparing large-area high valence transition metal-based catalysts with high OER activity and high stability for industrial production of clean hydrogen from water splitting.

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### Introduction

In order to take on the challenges of imminent energy, environmental and climatic crises caused by burning traditional fossil fuels, the development of renewable clean energy sources such as hydrogen, wind and solar energy has become a top priority agenda. Among possible solutions, water electrolysis-based hydrogen production in a sustainable fashion is the most promising one.<sup>1-3</sup> Extensive efforts have been made to tackle the kinetically sluggish four-electron transfer process of the OER, which causes huge energy loss and limits the large-scale application of water electrolysis for hydrogen production.<sup>4-7</sup> Many non-noble metal-based electrocatalysts have been developed to reduce the overpotential for water oxidation.<sup>8-10</sup>



Transition metal-based oxides or hydroxides have displayed good OER performance in alkaline environments yet free of the shortcomings of the scarcity and expensiveness of precious metal-based catalysts,<sup>11–13</sup> but their catalytic activity and stability under high current operating conditions need to be further improved.

It is widely known that the valence state of transition metals, *e.g.*, Fe ions, is pivotal to the OER activity of transition-metal-based catalysts. The high-valence transition metals have been reported to be the actual OER active sites and shown high OER intrinsic activity.<sup>14–18</sup> The Fe<sup>3+</sup> ions in the widely applied catalysts, such as NiFe-LDH, need be oxidized to Fe<sup>4+</sup> first under applied potentials before they can contribute to the OER process.<sup>19–21</sup> It thus appears that direct synthesis of stable Fe<sup>4+</sup>-based catalysts should reduce the overpotential of the OER. However, given that the typical oxidation states of Fe are +2 and +3, it is very difficult to obtain such stable Fe<sup>4+</sup>-containing electrocatalysts. As far as we know, Fe<sup>4+</sup> has only been reported in ABO<sub>3</sub> perovskite structure catalysts; however, that synthesis process was complicated and needed a very high annealing temperature ( $\geq 800$  °C).<sup>22,23</sup>

Because of the easy switch between Ce<sup>4+</sup> and Ce<sup>3+</sup>, CeO<sub>2</sub> has been widely used as a metal catalyst support to modulate the electronic structure through the so-called strong metal–support interaction (SMSI).<sup>24,25</sup> This also applies to metal oxides/hydroxides on a CeO<sub>2</sub> support, but the electronic interactions are commonly weaker and could hardly change the valence states of surface metal ions.<sup>26,27</sup> It has been reported that the transferred charge between the surface metal and support is inversely related to the metal cluster size,<sup>25,28</sup> the electronic interaction between the CeO<sub>2</sub> support and metal ions could thus be enhanced by reducing the particle size, making possible the formation of stable Fe<sup>4+</sup> on the CeO<sub>2</sub> support.

Here, we report a stable FeO<sub>x</sub> catalyst with a stable high valence of 4+, which shows greatly reduced overpotential for the OER. This was realized by a straightforward electroplating–etching method, whereby electroplated CeO<sub>2</sub> coating on a nickel foam (NF) was etched with ferric nitrate solution (Schematic illustration in Fig. 1(a)), installing and stabilizing the high valence Fe<sup>4+</sup> in FeO<sub>x</sub>/CeO<sub>2</sub>. The Fe<sup>4+</sup>-containing OER electrocatalyst (named Fe<sup>HV</sup>/CeO<sub>2</sub>@NF) only requires 238 mV overpotential to deliver 100 mA cm<sup>−2</sup> current density in 1 M KOH (pH = 13.6). Also, the OER potential of Fe<sup>HV</sup>/CeO<sub>2</sub>@NF can be stabilized at about 1.53 V vs. RHE (without *iR* compensation) to reach a high current density of 200 mA cm<sup>−2</sup> for over 500 h. Density functional theory (DFT) calculations uncovered that the effective interfacial d–f electron transfer from the US-FeO<sub>x</sub> to CeO<sub>2</sub> maintains the stable high valence states of Fe sites, which accounts for the high electroactivity for the OER. The optimized electronic structures guarantee the fast conversions of the intermediates on the catalyst, leading to the largely lowered overpotential required for the OER. This advanced catalyst system combined with its simple fabrication method should help to set the stage for the industrialization of sustainable OER catalysts and water-splitting systems.

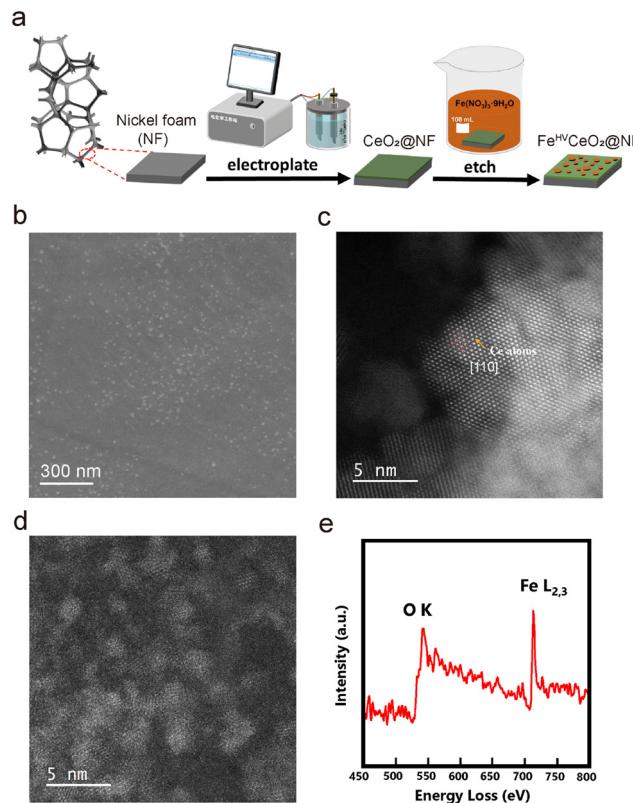


Fig. 1 Morphology and structure characterization of the Fe<sup>HV</sup>/CeO<sub>2</sub>@NF catalyst. (a) Schematic illustration of the synthesis process. (b) SEM image of Fe<sup>HV</sup>/CeO<sub>2</sub>@NF. (c) and (d) HAADF-STEM image of CeO<sub>2</sub> (c) and FeO<sub>x</sub> particles (d) in Fe<sup>HV</sup>/CeO<sub>2</sub>@NF. (e) EELS (electron energy loss spectroscopy) spectrum of d.

## Synthesis and characterization of high-valence Fe oxide

The scanning electron microscope (SEM) morphology of the CeO<sub>2</sub>@NF sample is shown in Fig. S1 (ESI<sup>†</sup>). A uniform electroplated layer, consisting of CeO<sub>2</sub> nanoparticles, is formed on the surface of NF. After being etched in ferric nitrate solution, CeO<sub>2</sub>@NF transforms to Fe<sup>HV</sup>/CeO<sub>2</sub>@NF and the morphology is shown in Fig. 1(b). The CeO<sub>2</sub> layer becomes flat, and there appears dense uniformly dispersed small white spots on the surface of CeO<sub>2</sub>, which are probably iron oxide nanoparticles. The scanning transmission electron microscopy (STEM)-energy dispersive X-ray spectrometry (EDS) mapping images (Fig. S2, ESI<sup>†</sup>) prove the uniform distribution of Ce and Fe in Fe<sup>HV</sup>/CeO<sub>2</sub>@NF. The high-angle annular dark field (HAADF)-STEM images of Fe<sup>HV</sup>/CeO<sub>2</sub>@NF are shown in Fig. 1(c) and (d). The Ce atomic columns belonging to CeO<sub>2</sub> [110] are clearly observed (Fig. 1(c)). In general, the sizes of the FeO<sub>x</sub> nanoparticles are smaller than 5 nm (Fig. 1(d)), which is confirmed by the atomic force microscope (AFM) images (Fig. S3, ESI<sup>†</sup>). The electron energy loss spectroscopy (EELS) spectrum taken from Fig. 1(d) is shown in Fig. 1(e), and it proves that the ultra-small nanoparticles are indeed iron oxide. These iron oxide particles are detached from the CeO<sub>2</sub> substrate, probably because



of the sonication process during TEM sample fabrication. The lattice fringes of both the  $\text{CeO}_2$  support and the ultra-small  $\text{FeO}_x$  particles with a size of about 2–5 nm are also observed from the HRTEM image of  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  (Fig. S4, ESI†). For the sake of comparison, LFe/ $\text{CeO}_2@\text{NF}$  (larger  $\text{FeO}_x$  particles) and  $\text{Fe}@\text{NF}$  (without  $\text{CeO}_2$  layer) reference samples were also fabricated. The LFe/ $\text{CeO}_2@\text{NF}$  sample was synthesized by increasing the concentration of  $\text{Fe}^{3+}$  in the etching solution. The morphology of the LFe/ $\text{CeO}_2@\text{NF}$  catalyst is shown in Fig. S5 (ESI†), which presents larger iron oxide particles ( $\sim 100$  nm) than that of the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  catalyst (Fig. 1(c)). As for the  $\text{Fe}@\text{NF}$  sample, what was partially etched away was the NF substrate in the acidic ferric nitrate solution accompanied by the formation of numerous holes, which is shown in Fig. S6 (ESI†).

The X-ray photoelectron spectroscopy (XPS), X-ray absorption fine structure (XAFS) and soft X-ray absorption spectroscopy (SXAS) were carried out to analyze the electronic interaction between  $\text{CeO}_2$  and Fe in  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$ , and the results are shown in Fig. 2. Fig. 2(a) presents the XPS spectra for the Fe 2p region of  $\text{Fe}@\text{NF}$ , LFe/ $\text{CeO}_2@\text{NF}$  and  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$ . While the peaks of  $\text{Fe}@\text{NF}$  and LFe/ $\text{CeO}_2@\text{NF}$

in the range of 710–712 eV correspond to  $\text{Fe}^{3+}$ , the peak of  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  in the range of 712–714 eV can be assigned to  $\text{Fe}^{3+}$  and  $\text{Fe}^{4+}$ .<sup>23,29</sup> The Fe K-edge XANES spectra of the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  catalyst and  $\text{Fe}_2\text{O}_3$  and  $\text{Fe}$  foil reference samples are shown in Fig. 2(b). Clearly, the Fe absorption edge of  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  shifts towards higher energy (inset of Fig. 2(b)) than that of  $\text{Fe}_2\text{O}_3$ , which confirms a higher Fe valence state in  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  than +3 in  $\text{Fe}_2\text{O}_3$ ,<sup>30,31</sup> and also is in agreement with the XPS result. Further support is from the SXAS shown in Fig. 2(c). The Fe L-edge is divided into two peaks (at 714 eV and 726 eV) corresponding to the  $\text{L}_3$  and  $\text{L}_2$  edges. Compared with standard  $\text{Fe}_2\text{O}_3$ , the Fe L-edge in  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  presents a blue shift, again evidencing a higher Fe valence state than +3 in  $\text{Fe}_2\text{O}_3$ .<sup>32,33</sup> Fig. 2(d) shows the Ce 3d XPS spectra, and the total areas of the  $\text{Ce}^{3+}$  and  $\text{Ce}^{4+}$  peaks are in proportion to their contents.<sup>34,35</sup> The  $\text{Ce}^{3+}$  content in the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  catalyst is calculated to be 35%, while the  $\text{Ce}^{3+}$  content in  $\text{CeO}_2@\text{NF}$  is 31%. The above results indicate an electronic transfer between Fe and Ce in  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$ , that is, part of  $\text{Fe}^{3+}$  is oxidized to  $\text{Fe}^{4+}$  associated with the shift of  $\text{Ce}^{4+}$  to  $\text{Ce}^{3+}$ .

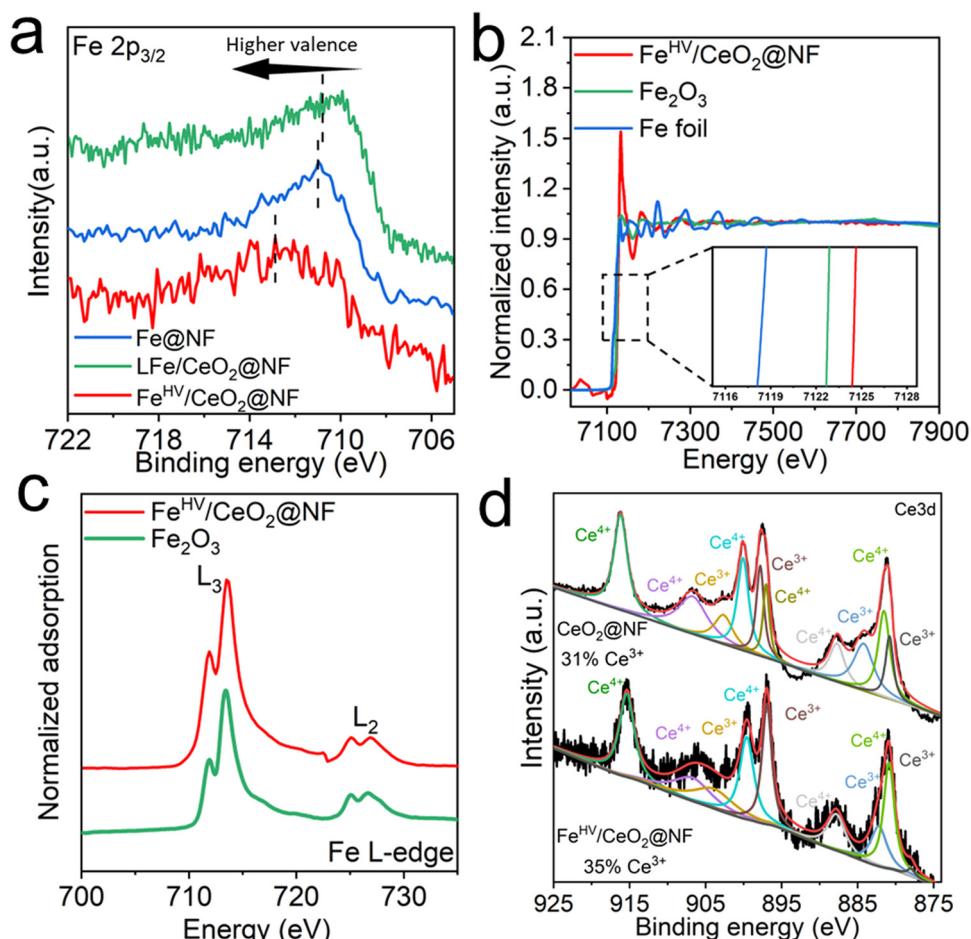


Fig. 2 Electronic structure characterization of the catalysts. (a) Fe 2p XPS spectra of the catalysts. (b) XANES of the samples at the Fe K-edge. The inset enlarges the absorption edges and displays the valence changes of Fe in the samples. (c) Soft X-ray absorption spectra (SXAS) of the catalyst and the  $\text{Fe}_2\text{O}_3$  control sample at the Fe L-edge. (d) Ce 3d XPS spectra of the catalysts.



# Electrochemical performance of $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$

The electrochemical performances of the electrodes were investigated using a three-electrode system in an alkaline environment (1 M KOH, pH = 13.6). As can be seen from Fig. S8 (ESI<sup>†</sup>), the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode displays much better OER performance than the  $\text{Fe}@\text{NF}$ ,  $\text{CeO}_2@\text{NF}$  LFe/CeO<sub>2</sub>@NF and RuO<sub>2</sub>@NF electrodes. The faradaic efficiency is calculated to be 98% at the current density of 30 mA cm<sup>-2</sup> (with the overpotential as low as 210 mV). To reach the high current densities of 100 mA cm<sup>-2</sup> and 200 mA cm<sup>-2</sup>, the required overpotentials are only 238 mV and 261 mV, respectively. Comparatively speaking, under the overpotential of 238 mV (261 mV), the current density of the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode is 5.7(2.6), 8.1(4) and 42.5(43.4) times larger than those of the  $\text{Fe}@\text{NF}$ , LFe/CeO<sub>2</sub>@NF and CeO<sub>2</sub>@NF electrodes, respectively, as can be seen clearly in Fig. 3(a). The Nyquist plot in Fig. 3(b) shows that the semicircle radius of the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode is much smaller than that of  $\text{Fe}@\text{NF}$  and LFe/CeO<sub>2</sub>@NF, indicating the smallest charge transfer resistance of the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode.<sup>36,37</sup> The Tafel plots in Fig. 3(c) show that the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode has the smallest Tafel slope of 40 mV dec<sup>-1</sup>, which is beneficial to accelerate the reaction kinetics.<sup>38,39</sup> The OER activity comparison between the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode and recently reported OER electrode<sup>26,33,34,40-45</sup> by comparing

the overpotential to reach the geometric current density of 30, 50 and 100 mA cm<sup>-2</sup> (Fig. 3(d) and Table S1, ESI<sup>†</sup>) indicates the superior OER activity of the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode. As shown in Fig. S9 (ESI<sup>†</sup>), the OER performance of the reference sample on a Cu foam substrate illustrates the positive role of Ni foam.

In order to explore whether the high catalytic activity of the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode comes from the high specific surface area or the high intrinsic activity of the active sites, we conducted cyclic voltammetry (CV) tests on the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$ , LFe/CeO<sub>2</sub>@NF and  $\text{Fe}@\text{NF}$  electrodes (Fig. S10, ESI<sup>†</sup>) in the non-Faraday potential range (0.32–0.40 V vs. Hg/HgO) and calculated the electrochemical double-layer capacitance ( $C_{\text{dl}}$ ) of the three electrodes. The ECSA was calculated according to the formula  $\text{ECSA} = C_{\text{DL}}/C_s$ , where a specific capacitance of  $C_s = 0.040 \text{ mF cm}^{-2}$  was used in this work. The calculated ECSA values are shown in Fig. 3(e) and summarized in Table S2 (ESI<sup>†</sup>), as well as other relevant electrochemistry parameters. The ECSAs of the LFe/CeO<sub>2</sub>@NF and  $\text{Fe}@\text{NF}$  electrodes are about 5 times and 2.2 times as large as that of  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$ , which demonstrates that the high OER activity of the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode derives from the high intrinsic activity of  $\text{Fe}^{\text{HV}}$  active sites considering that the lower ECSA means an electrode has less active sites. The specific activity ( $j_{\text{ECSA}}$ ) is further obtained and shown in Fig. 3(f), where it is evident that the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode has much higher intrinsic OER activity than LFe/CeO<sub>2</sub>@NF and  $\text{Fe}@\text{NF}$  electrodes. Fig. S11 and

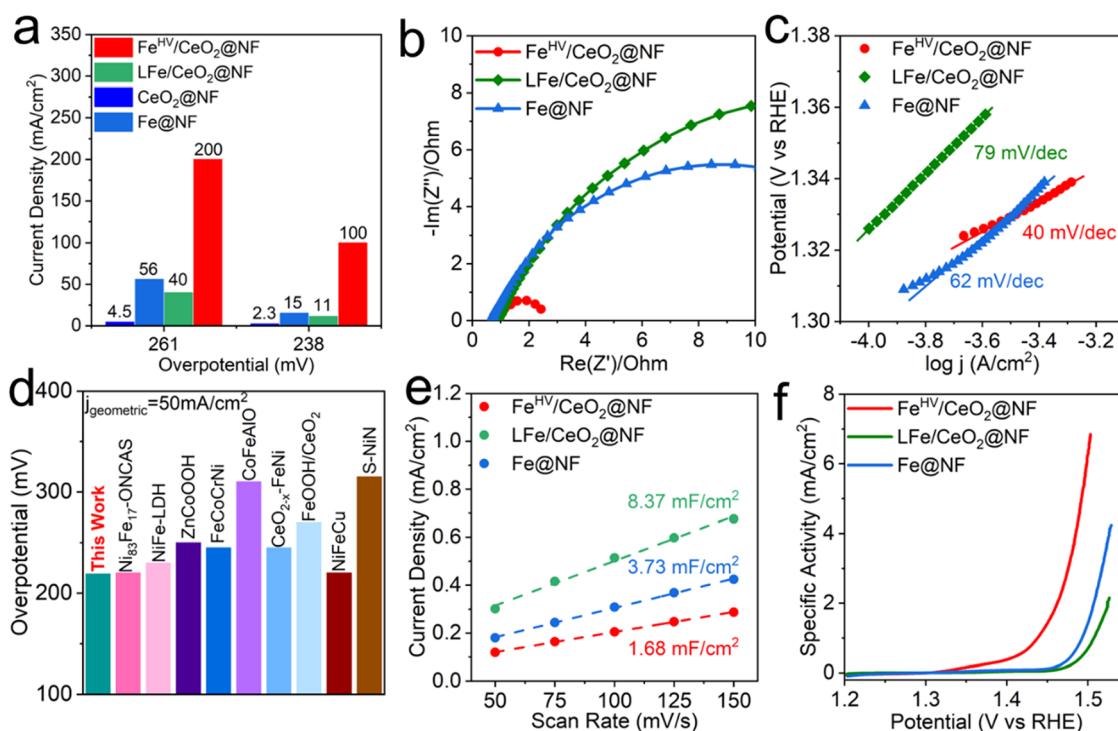


Fig. 3 OER performance characterization of the electrodes. (a) Comparison of the current densities achieved by different electrodes at special overpotentials. (b) EIS curves, measured at frequencies from  $10^{-1}$  to  $10^5$  Hz. (c) Tafel slope plots. (d) Activity comparison among OER electrocatalysts<sup>26,33,34,40-45</sup> based on the overpotential needed to reach the geometric current density of 50 mA cm<sup>-2</sup> (for more details see Table S1, ESI<sup>†</sup>). (e) Electrochemical double-layer capacitance. The slope of current density at OCP vs. scan rate represents the double-layer capacitance. (f) Polarization curves with the current density normalized by ECSA.

Table S3 (ESI†) show the comparison of the specific activity between the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode and a series of recently reported high OER performance electrodes by comparing the overpotential at an ECSA normalized current density of  $1 \text{ mA cm}^{-2}$ . The result shows that the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode has the best OER specific activity among these electrodes.

## Formation of high-valence Fe oxide for the OER by DFT calculations

To further reveal the formation mechanism of high-valence Fe oxide ( $\text{FeO}_x$ ) and its contributions to the OER performances in  $\text{Fe}/\text{CeO}_2$ , DFT calculations have been further introduced to investigate the electronic structures and energetic preferences. For the bonding and anti-bonding orbitals near the Fermi level ( $E_F$ ), the electron-rich feature of the surface  $\text{US-FeO}_x$  is noted (Fig. 4(a)). The electron-rich features indicate a higher electron density of  $\text{Fe}/\text{CeO}_2$  near the  $E_F$ . Since more electrons locate close to  $E_F$ , the electron transfer barriers are much lower, suggesting that the electrons are easier to be transferred from

the  $\text{Fe}/\text{CeO}_2$  surface in accordance with the higher electroactivity of the Fe sites. Meanwhile, the strong orbital couplings at the interface between  $\text{US-FeO}_x$  and  $\text{CeO}_2$  result in favorable electronic modulations. In comparison,  $\text{CeO}_2$  only exhibits an electron-rich feature near the oxygen vacancies while the remaining sites of the surface with dominant contributions from anti-bonding orbitals cannot support the efficient adsorption and conversion of intermediates (Fig. 4(b)). The projected partial density of states (PDOS) have unraveled the electronic structure of  $\text{Fe}/\text{CeO}_2$  (Fig. 4(c)).  $\text{Fe-3d}$  orbitals have shown the  $e_g-t_{2g}$  splitting near the  $E_F$ , where the overlapping with the  $\text{Ce-4f}$  orbitals supports the potential d-f electron transfer. The detailed electronic contributions of surface Ni sites have been revealed, *viz.*, the  $\text{Ni-3d}$  orbitals mainly locate near the  $E_F$  with the dominant peak at  $E_V - 0.85 \text{ eV}$ , thereby pulling up the overall electroactivity as well as the electron transfer rate with an improved  $^*\text{OH}$  adsorption energetics and kinetics for optimum OER performance. In addition, the  $\text{O-2p}$  orbitals have exhibited evident orbital coupling with both  $\text{Ce-4f}$  and  $\text{Fe-3d}$ , indicating the small barriers for electron transfer within  $\text{Fe}/\text{CeO}_2$ . In comparison, the  $\text{Ce-4f}$  orbitals are much flattened

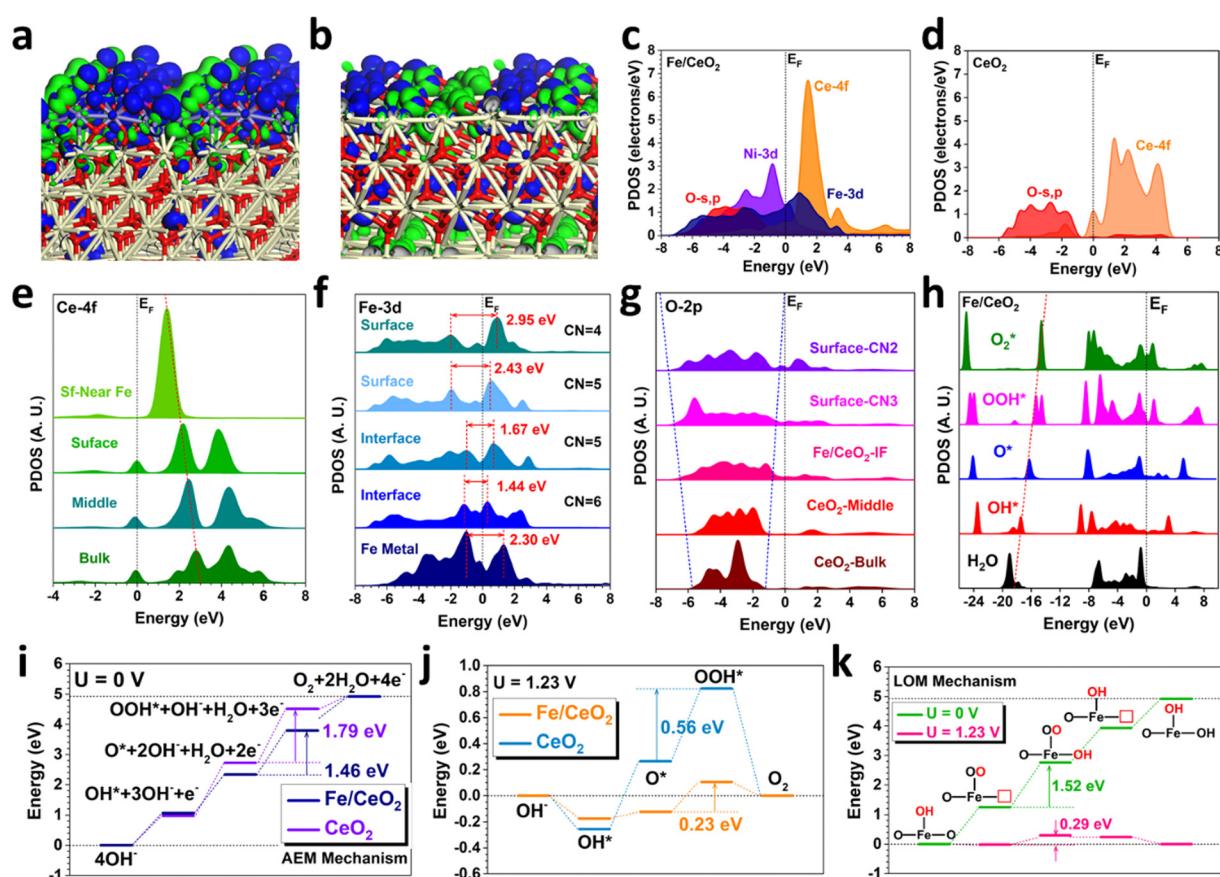


Fig. 4 DFT calculations. (a) The 3D contour plot of electronic distribution near the Fermi level of  $\text{Fe}/\text{CeO}_2$ . (b) The 3D contour plot of electronic distribution near the Fermi level of  $\text{CeO}_2$ . Yellow balls = Ce, Blue balls = Fe, Purple balls = Ni, and Red balls = O. Blue isosurface = bonding orbitals, and green isosurface = anti-bonding orbitals. (c) The PDOS of  $\text{Fe}/\text{CeO}_2$ . (d) The PDOS of  $\text{CeO}_2$ . (e) The site-dependent PDOS of  $\text{Ce-4f}$  in  $\text{Fe}/\text{CeO}_2$ . (f) The site-dependent PDOS of  $\text{Fe-3d}$  in  $\text{Fe}/\text{CeO}_2$ . (g) The site-dependent PDOS of  $\text{O-2p}$  in  $\text{Fe}/\text{CeO}_2$ . (h) The PDOS of key adsorbates during the OER process in  $\text{Fe}/\text{CeO}_2$ . (i) The reaction energy changes of the OER process on  $\text{Fe}/\text{CeO}_2$  and  $\text{CeO}_2$  under  $U = 0 \text{ V}$  through the AEM mechanism. (j) The reaction energy changes of the OER process on  $\text{Fe}/\text{CeO}_2$  and  $\text{CeO}_2$  under  $U = 1.23 \text{ V}$  through the AEM mechanism. (k) The reaction energy changes of  $\text{Fe}/\text{CeO}_2$  through the LOM mechanism.

and O-2p orbitals also become less broadened in  $\text{CeO}_2$  (Fig. 4(d)). The limited overlap between O-2p and Ce-4f orbitals induces an evident barrier of electron transfer, which results in the significantly increased overpotentials of the OER.

In order to interpret the detailed interactions between surface  $\text{US-FeO}_x$  and  $\text{CeO}_2$ , the site-dependent PDOSs have been supplied to illustrate the electronic structure evolution (Fig. 4(e)). We notice a gradual shifting for the empty Ce-4f orbital towards the  $E_F$  from the bulk to the surface sites near the Fe SNPs, potentially leading to efficient d-f electron transfer from Fe-3d orbitals to Ce-4f orbitals. Thus there appears to be a trend of maintaining the high oxidation states of Fe SNPs induced by the O sites, which is also confirmed by the experimental findings that the surface  $\text{US-FeO}_x$  indeed exists in the high valence states. With the d-f electron transfer, the Fe-3d orbitals display essential contributions to the OER process. Specifically, the Fe sites at the interface show an alleviated  $e_g-t_{2g}$  splitting of 3d orbitals in  $\text{Fe/CeO}_2$ , benefiting the fast electron transfer from Fe SNPs to  $\text{CeO}_2$  (Fig. 4(f)). Compared to the interface, however, the surface Fe sites have shown an enlarged  $e_g-t_{2g}$  splitting. The overall downshifted 3d orbitals of the surface Fe sites significantly enhance the energy band overlap with surface O sites, thus lowering the electron transfer energy costs and further improving the electroactivity of the high valence Fe sites towards the water oxidation process.<sup>46,47</sup> Moreover, the O coordination numbers decrease with increasing  $e_g-t_{2g}$  splitting, especially for the surface Fe sites, supporting the higher electroactivity of Fe sites with high valence states. For the O sites, it is noted that the orbitals become much broadened from the bulk  $\text{CeO}_2$  to the surface, which considerably enhances their couplings with the Fe-3d orbitals

to promote electroactive Fe sites towards the OER (Fig. 4(g)). More importantly, the increased electron density near the Fermi level has improved the electron transfer at the interface. In addition, the PDOSs of the key intermediates have been revealed on the  $\text{Fe/CeO}_2$  (Fig. 4(h)). Notably, the  $\sigma$  orbitals of O-2p in the key intermediates have shown a highly linear correlation for the OER process, which provides an efficient intermediate conversion process with efficient electron transfer. Such a correlation also supports that the surface  $\text{US-FeO}_x$  are the main active sites that contribute to the superior OER performances in  $\text{Fe/CeO}_2$ .

We next examine the reaction trends of the uphill OER process from the kinetic perspectives regarding both the adsorbate evolution mechanism (AEM) and lattice-oxygen-mediated mechanism (LOM). For the conventional AEM, the highest energy barrier occurs in the rate-determining step (RDS) from  $\text{O}^*$  to  $\text{OOH}^*$ . In comparison,  $\text{Fe/CeO}_2$  shows an energy barrier of 1.46 eV for the RDS, which is much lower than that with the  $\text{CeO}_2$  at 1.79 eV and thus guarantees the much-improved OER performance. The overpotential has been estimated by reference to the standard equilibrium potential ( $U = 1.23$  V) (Fig. 4(j)). Although  $\text{Fe/CeO}_2$  shows an uphill trend from  $\text{OH}^*$  to  $\text{OOH}^*$ , the subtle energy barriers lead to a small overpotential of 0.23 V. In contrast,  $\text{CeO}_2$  shows overbinding of  $\text{OH}^*$  in the initial reaction process, leading to high barriers for the following conversions to  $\text{O}^*$  and  $\text{OOH}^*$  with a large overpotential of 0.56 V. For the LOM mechanism (Fig. 4(k)), on the other hand, the highest energy barrier of 1.52 eV is located in the  $^*\text{OH}$  adsorption step on the nearby oxygen vacancy sites of Fe. Compared to the RDS in AEM, the LOM presents a higher energy barrier. Using the standard equilibrium potential of 1.23 V as the reference, the overpotential of OER is determined

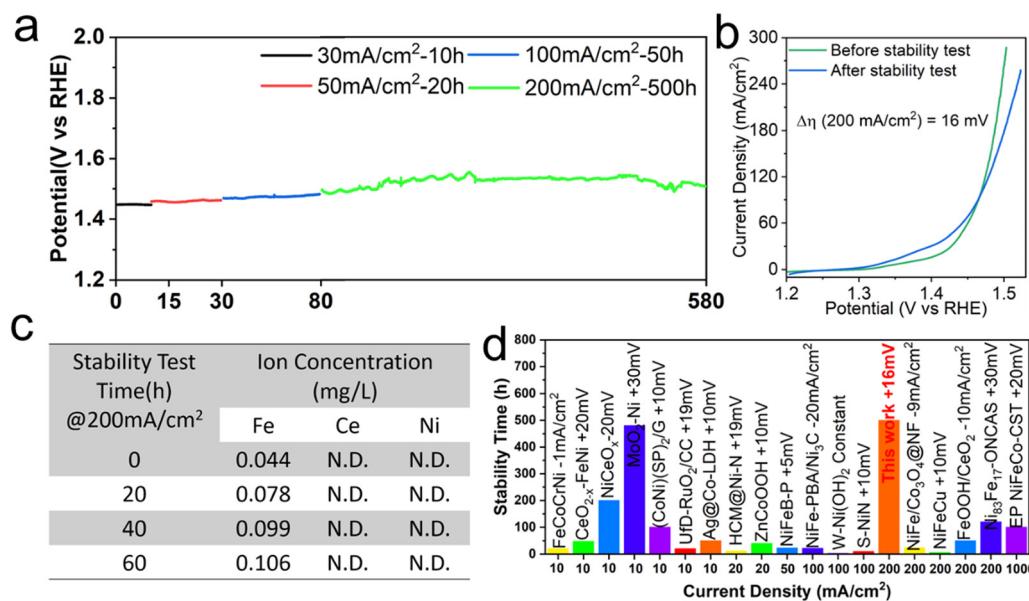


Fig. 5 Electrochemical stability characterization of the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@NF$  electrode in 1 M KOH. (a) The chronopotentiometry measurements. (b) Polarization curves before and after the stability test with a scan rate of  $10 \text{ mV s}^{-1}$ . (c) The ion concentrations of Fe, Ce, and Ni (data acquired by inductive coupled plasma emission spectroscopy) in the electrolyte during the 60 h water electrolysis process. N.D. stands for a value below the measurement limit of  $0.005 \text{ mg L}^{-1}$ . (d) Summary of the stability comparison<sup>26,33-35,41,42,44,45,48-57</sup> of recently reported efficient oxygen evolution electrocatalysts (details in Table S4, ESI†). The electrolyte is 1 M KOH or NaOH.



to be 0.29 V, which is higher than that of the AEM. Therefore, we can conclude that the AEM is preferred over the LOM for our OER catalyst, which agrees with the experimental characterizations (Fig. S12, ESI<sup>†</sup>).

## OER stability of $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$

Besides catalytic activity, operational stability is another key parameter to evaluate the performance of a given electrocatalyst, especially for the industrial water splitting production of  $\text{H}_2$ . The chronopotentiometry measurement in 1 M KOH is carried out to evaluate the long-term stability of  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$ . In the continuous stability test of 30, 50, 100 and 200  $\text{mA cm}^{-2}$ , the potential at every stage remains almost constant and at a high current density of 200  $\text{mA cm}^{-2}$ , the potential can be stabilized at about 1.53 V *vs.* RHE (without *iR* compensation) for up to 500 h (Fig. 5(a)). By comparing the linear sweep voltammetry (LSV) curves of the electrode before and after the stability test, it can be seen that after the long-term (590 h) and high-current stability test, the overpotential increases by only 16 mV at the current density of 200  $\text{mA cm}^{-2}$  (Fig. 5(b)), indicating that  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  is a super-stable electrode for water oxidation. To assess the valence stability of Fe in the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrocatalyst, XPS, SXAS and

Mössbauer spectroscopy were carried out on the samples after a 10 hours OER test at the current density of 30  $\text{mA cm}^{-2}$ . As comparatively shown in Fig. S13–S15 (ESI<sup>†</sup>), the excellent stability of  $\text{Fe}^{4+}$  in the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode is clearly evidenced. The inductively coupled plasma optical emission spectrometer (ICP-OES) is further used to measure the concentration of metal ions in the electrolyte during the water electrolysis of the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode. As shown in Fig. 5(c), only Fe ions are detected and the content tends to be stable after the 60 h stability test. The excellent stability may derive from the fine and highly dispersed iron oxide nanoparticles, which can strongly combine with ceria and are difficult to fall off during the OER test. Therefore, the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode behaves steadily for water oxidation. Fig. 5(d) and Table S4 (ESI<sup>†</sup>) summarize the stability comparison of recently reported efficient oxygen evolution electrodes,<sup>26,33–35,41,42,44,45,48–57</sup> and the stability of the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode is outstanding among them.

## Large-area fabrication and OER performance under industrial conditions

Last but not least, an industrializable water splitting electrode should be equipped with the ability to scale up its production. Currently, common electrode preparation methods in laboratories

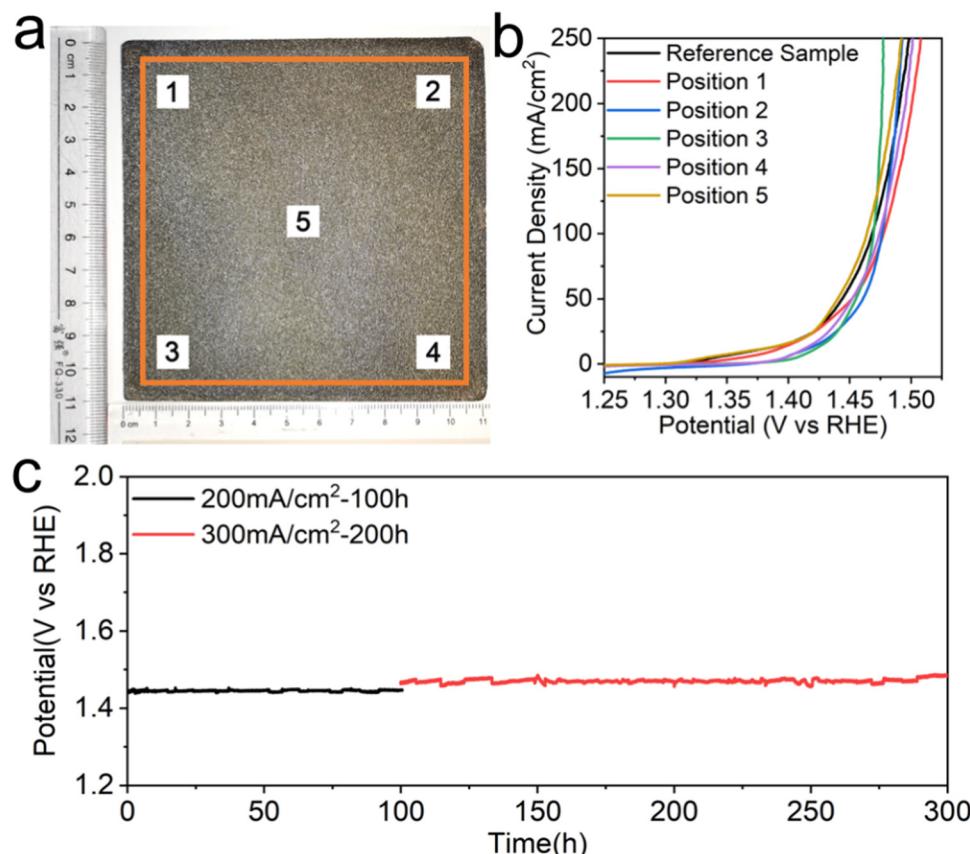


Fig. 6 Industrial application of the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode. (a) Photograph of the large-area ( $100 \text{ cm}^2$ )  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode. (b) Polarization curves of the five different positions ( $1 \text{ cm}^2$ ) in the large-area  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode (the reference sample is directly synthesized with  $1 \text{ cm}^2$ ). (c) The chronopotentiometry measurements in 6 M KOH and at 60 °C.



often involve complex processes and small electrode sizes, making it difficult to meet the industrial requirements for large-scale production. The electroplating-etching method presented here enables us to prepare a large-area electrode due to the simplicity of the preparation process and the scalability of the required experimental equipment, which is of great significance for the realization of industrialized hydrogen production by electrolysis of water. The large-size  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode with  $10 \times 10 \text{ cm}^2$  effective area is synthesized through the electroplating-etching method, and a photograph is shown in Fig. 6(a). We selectively cut out five  $1 \times 1 \text{ cm}^2$  electrodes (labeled in Fig. 6(a)) from the  $10 \times 10 \text{ cm}^2$  electrode to perform the OER test, and compare their performances with the directly prepared  $1 \times 1 \text{ cm}^2$  electrode (reference sample in Fig. 6(b)). The results show that the performance of these six electrodes remained consistent within a certain fluctuation range (Fig. 6(b)). This proves that the large-area electrode has good uniformity, which in turn confirms the scalability of this preparation method. There is no doubt that this is one small but important step towards the realization of industrialized hydrogen production by electrolysis of water. Finally, the industrial-level (6 M KOH, 60 °C) OER activity and stability tests of the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode ( $1 \text{ cm}^2$ ) were conducted. The polarization curve in Fig. S16 (ESI†) shows that the current density can reach  $500 \text{ mA cm}^{-2}$  at the overpotential of 270 mV (without  $iR$  compensation). As shown in Fig. 6(c), the electrode of  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  could deliver large current densities of  $200 \text{ mA cm}^{-2}$  and  $300 \text{ mA cm}^{-2}$  at only 1.44 V and 1.47 V vs. RHE (without  $iR$  compensation) and remain stable up to 300 h in such a harsh environment, further indicating the potential industrial application.

## Conclusions

We have demonstrated an efficient and ultra-stable high valence  $\text{Fe}^{4+}$ -containing OER electrode (named  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$ ) and its scalable preparation ( $100 \text{ cm}^2$  reported here) by a simple electroplating-etching method. The ultra-small size of the  $\text{FeO}_x$  nanoparticles on the  $\text{CeO}_2$  support enabled the formation of  $\text{Fe}^{4+}$ , which has shown high intrinsic activity. Remarkably, the  $\text{Fe}^{\text{HV}}/\text{CeO}_2@\text{NF}$  electrode could deliver large current densities of  $200 \text{ mA cm}^{-2}$  and  $300 \text{ mA cm}^{-2}$  at only 1.44 V and 1.47 V vs. RHE (without  $iR$  compensation) and remain stable up to 300 h under harsh industrial conditions (6 M KOH, 60 °C). DFT calculations unraveled the electronic modulations induced by the d-f electron transfer at the interface of  $\text{Fe}/\text{CeO}_2$ , which held the  $\text{US-FeO}_x$  in a stabilized high valence  $\text{Fe}^{4+}$  state conducive to the OER process. This work provides a guide to designing and preparing large-area high valence transition metal-based electrodes with high OER activity and high stability for industrial production of clean hydrogen from water splitting.

## Author contributions

Shihe Yang and Jun Yu conceived and supervised the project. Hongzhi Liu conducted the experiments. Mingzi Sun performed

the computational study under the supervision of Bolong Huang. Jinghuang Lin performed the STEM analysis with Bin Feng and Yuichi Ikuhara. Kun Qian performed the XAFS characterization. Hongzhi Liu, Jun Yu, Mingzi Sun, Bolong Huang and Shihe Yang wrote the paper. All authors discussed the results and commented on the manuscript.

## Conflicts of interest

The authors declare no conflicts.

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