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An electrochemically engineered layer of γ -NiOOH with FeOOH on nickel foam for durable OER catalysis for anion exchange membrane water electrolysis

An efficient and durable oxygen evolution reaction (OER) catalyst for alkaline/anion exchange membrane water electrolysis has been developed by a simple two-step electrochemical process. The two-step processes involve the selective modification of γ -NiOOH on the surface of Ni foam followed by the deposition of Fe. The synergistic effect of γ -NiOOH and FeOOH on the modified surface is demonstrated to improve OER activity and durability.

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An electrochemically engineered layer of γ -NiOOH with FeOOH on nickel foam for durable OER catalysis for anion exchange membrane water electrolysis†

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An efficient and durable oxygen evolution reaction catalyst is essential for the advancement of anion exchange membrane water electrolyzers. Here we report a simple, rapid, and cost-effective two-step electrochemical synthesis of an efficient and durable oxygen evolution reaction catalyst, which can be produced on an industrial scale. In the first step of catalyst preparation, a ≈ 250 nm thick γ -NiOOH layer was electrochemically developed over nickel foam. Then iron was electrochemically deposited on γ -NiOOH to stabilize the NiOOH species and improve the OER activity. Various material characterization techniques confirmed the presence of a highly OER-active combination of γ -NiOOH and FeOOH at the catalyst surface. The synergism between high valent Ni^{3.6+} in γ -NiOOH and Fe³⁺ in FeOOH helps the catalyst to attain 10 mA cm⁻² at 1.47 V with a very low Tafel slope value of 34 mV per decade. The catalyst attained 1 A cm⁻² at 1.65 V in 1 M KOH at 80 °C under real electrolyzer operational conditions. Furthermore, the catalyst exhibits stable OER activity at high current densities under real electrolyzer testing conditions.

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Sustainability spotlight

Recently, the demand for hydrogen as a carbon-free fuel has been increasing. Among the various methods of hydrogen production, green hydrogen production is the most compatible for a carbon neutral society. For global acceptance, the production cost of green hydrogen must be competitive with fossil fuel and other methods of hydrogen production. Anion exchange membrane water electrolysis is a relatively inexpensive technology for green hydrogen production because non-noble metals can be used as catalysts and electrolyzer components. In this work, a simple, scalable and energy efficient new method for non-noble metal catalyst preparation for water oxidation is described. The catalyst shows durable oxygen evolution activity in anion exchange membrane water electrolysis at high current density.

Introduction

Water electrolysis coupled with renewable energy sources is a sustainable way to produce green hydrogen. Water electrolysis consists of the hydrogen evolution reaction (HER) and oxygen evolution reaction (OER) as two half-cell reactions. Theoretically, the overall water splitting reaction requires 1.23 V at 25 °C, but in reality, another 0.2 V or above overpotential is required to initiate the water splitting process due to poor catalytic activity and ohmic resistance of the system.^{1–3} Fundamentally, the OER is a sluggish electrochemical reaction, involving oxygen–oxygen

bond formation, that occurs *via* four proton-coupled electron transfers.^{1,4–6} Hence the contribution from the OER to the overall energy loss of the electrolyzer is the highest. Further, since the OER takes place in a highly oxidative environment, catalyst degradation and loss of catalytic activity are issues that affect the life of the electrolyzer.⁷ An efficient and stable OER catalyst is inevitable for a durable water electrolyzer.

The first row of transition metals exhibits promising OER activity in an alkaline medium.^{5,8–10} A wide range of transition metal compounds like oxides,^{11–13} nitrides,¹⁴ sulphides,¹⁵ phosphides/phosphates,^{16–19} and carbides²⁰ were reported as superior OER catalysts. Systematic analysis of the OER mechanism of transition metal-based catalysts using different spectroscopic techniques reveals that the real catalytically active species are oxides of transition metals.^{21–23} M–OH, M–O, M–OOH, and M–OO intermediates were formed on all transition metal catalysts during the OER, regardless of the elements present with the metal on the catalyst surface.^{17,21,24–27} Later, it was discovered

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that bi/tri-metallic compounds have augmented OER activity compared to the corresponding mono-metallic catalysts.^{3,28–30} Among various bimetallic oxides, Ni–Fe based catalysts are more efficient and cost-effective for the OER.^{31–33} Even though the exact catalytic center of the Ni–Fe catalyst is still uncertain, all findings unanimously accept that the oxyhydroxide of these metals is important in the OER catalytic cycle.^{22,34–36} Individually, NiOOH and FeOOH are OER active but require a higher overpotential, and FeOOH has higher OER overpotential than NiOOH. However, the addition of a small amount of Fe to the NiOOH lattice was found to reduce the OER overpotential. Different findings have been reported regarding the high OER activity of the Fe doped NiOOH catalyst and its active sites. Combined X-ray adsorption analysis and DFT + U studies suggest that the Fe³⁺ incorporating γ -NiOOH is the active site for the OER.³⁷ Mössbauer spectroscopy and Spectroelectrochemical + DFT studies confirm the formation of Fe⁴⁺ facilitated by NiOOH during the OER and report that the high valent iron oxide might be an important center of OER activity.^{38,39} Other theoretical studies suggest that both Fe and Ni synergistically promote the oxidation state of each other to the +4 state and both high valent metals participate equally to promote the OER activity of Fe doped NiOOH.⁴⁰ Further, it is also reported that Fe addition to the lattice of NiOOH promotes the electronic conductivity of NiOOH.⁴¹

Several methods have been reported for the synthesis of NiFe catalysts in either powder form or on the surface of a conducting surface. Among the various synthesis methods, electrochemical synthesis is fairly fast, highly reproducible, and produces minimal waste. Further, since this method commonly requires low temperature and pressure, it is a more energy efficient and economical than other traditional material syntheses like wet-chemical, solid-state and vapor-phase synthesis. Here, we present a modified simple and quick two-step electrochemical method for the scalable synthesis of a durable Fe incorporated γ -NiOOH (γ -NiOOH–Fe) OER catalyst with high performance. The two-step method involves the selective synthesis of a thick γ -NiOOH layer (≈ 250 nm) on a clean Ni foam through electrochemical oxidation followed by electrochemical deposition of a thin layer of Fe on the γ -NiOOH layer. Material characterization of γ -NiOOH–Fe confirms the formation of γ -NiOOH after oxidation of Ni foam and the presence of both γ -NiOOH and FeOOH after Fe deposition. The electrocatalytic analysis of γ -NiOOH–Fe revealed its high OER activity by attaining a current density of 10 mA cm⁻² at 1.47 V with a Tafel slope value of 34 mV per decade. The OER activity of γ -NiOOH–Fe remained stable after the accelerated durability test and long term constant current electrolysis at different current densities at room temperature. Furthermore, an anion exchange water electrolyzer was assembled with γ -NiOOH–Fe as the anode catalyst, and the electrolysis was performed at 80 °C in 1 M KOH solution. The electrochemical performance of the γ -NiOOH–Fe anode was simultaneously evaluated during electrolyzer polarization. Later, the durability of the γ -NiOOH–Fe anode was tested during high current density electrolysis. We also confirm that the described method of catalyst preparation

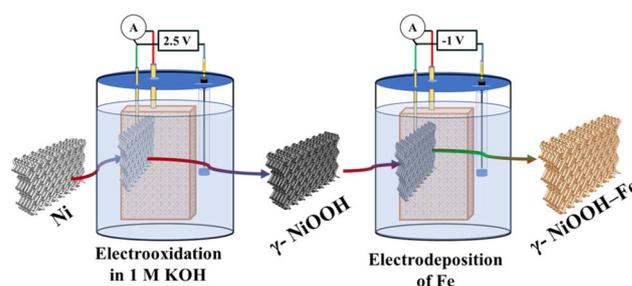
is a highly reproducible, fast, economical, and feasible technique for large-scale production.

Results and discussion

Commercially purchased nickel foam has a surface oxide layer. Acid washing with 1 M HCl followed by water and alcohol cleaning removes the surface oxide layer and grease on the Ni foam.⁴² In the catalyst preparation process (Scheme 1), at first, a layer of NiOOH was fabricated over a cleaned Ni foam surface by oxidizing the Ni foam at a high potential (2.5 V) in 1 M KOH solution. Electrochemical oxidation of Ni in an alkaline medium initially produces Ni(OH)₂ and further oxidation leads to the formation of NiOOH. As per the Bode cycle, (Fig. S1†) when Ni(OH)₂ becomes oxidized, β -NiOOH forms primarily, and further oxidation causes the formation of γ -NiOOH.^{43,44} Here, the formation of γ -NiOOH mainly occurred on surface of the Ni foam, as the Ni foam was oxidized at 2.0 V higher than required for the formation of NiOOH. As it is undoubtedly proved that NiOOH is an important intermediate in the oxygen evolution reaction and the addition of Fe further enhances the OER activity and stability of NiOOH, a thin layer of Fe was electrochemically deposited on γ -NiOOH to obtain γ -NiOOH–Fe. For a comparison, another catalyst with only Ni–Fe composition was also prepared on a unoxidized Ni foam surface through a similar electrochemical Fe deposition process.

The Raman spectrum of the oxidised Ni foam, presented in Fig. 1a, shows two major peaks at 479 cm⁻¹ and 560 cm⁻¹, attributed to the e_g bending vibration and the A_{1g} stretching vibration of Ni–O in γ -NiOOH, respectively. Further, the broad peak band between 850 cm⁻¹ and 1200 cm⁻¹ is ascribed to the active oxygen species (NiOO⁻) in oxyhydroxide.^{22,45} Later, when γ -NiOOH–Fe was derived by electrodepositing a thin layer of Fe on γ -NiOOH, an additional peak at 684 cm⁻¹ was observed in addition to the peaks of γ -NiOOH, which corresponds to the FeOOH.⁴⁶ The presence of FeOOH indicates that the electrochemically deposited Fe was quickly oxidized by the pre-oxidized Ni foam substrate consisting of γ -NiOOH. Meanwhile, the Raman spectrum of Ni–Fe has indicated the peaks at 456 and 534 cm⁻¹ attributed to the Ni–O stretching vibration of NiFe.⁴⁷

The XRD pattern of acid washed Ni foam has two major peaks at 2 θ values of 44.6 and 51.8 corresponding to the (111) and (200) planes of Ni.⁴⁸ Although the Raman spectra showed



Scheme 1 Electrochemical fabrication of catalysts.



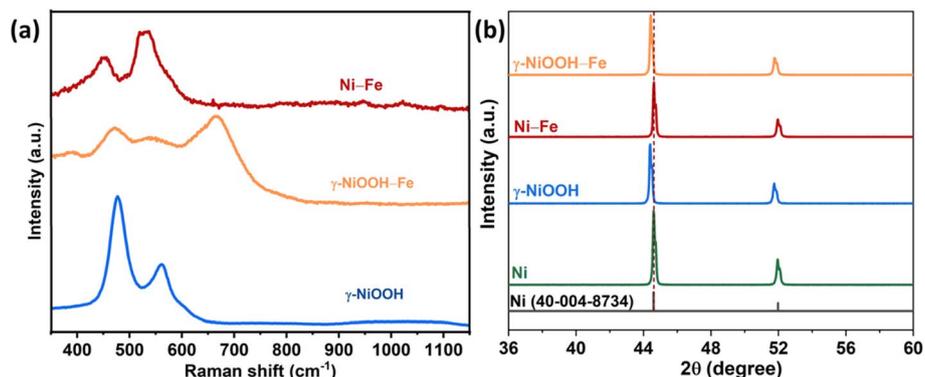


Fig. 1 (a) Raman spectra of γ -NiOOH, γ -NiOOH-Fe and Ni-Fe catalysts; (b) XRD patterns of Ni-foam after acid washing, γ -NiOOH, Ni-Fe, and γ -NiOOH-Fe.

the presence of γ -NiOOH on the surface of the oxidized Ni foam, the X-ray diffraction pattern showed no peaks of γ -NiOOH. But the peak position of γ -NiOOH shifted to lower angles compared to the acid washed Ni foam (Fig. 1b). The shift in peak position is attributed to lattice plane broadening due to the formation of γ -NiOOH. The crystal structure of γ -NiOOH comprises a large interlayer spacing (7 Å) and contains intercalated species such as water or ions that are absorbed between the layers. Further, the electrochemical deposition of Fe shows no characteristic change in the XRD pattern compared to that of γ -NiOOH, which is due to the thin layer deposition of Fe on γ -NiOOH. In the case of the Ni-Fe catalyst, no new peaks and no change in peak

positions were observed when a thin layer of Fe was deposited on the Ni surface.

The surface oxidation states of γ -NiOOH-Fe and Ni-Fe were analyzed by X-ray photoelectron spectroscopy. The fine scan XPS spectra of Ni 2p, Fe 2p and O 1s of γ -NiOOH-Fe are presented in Fig. 2. The deconvoluted Ni 2p_{3/2} spectrum of γ -NiOOH-Fe consists of Ni²⁺ and Ni³⁺ peaks at 856.8 and 858.09 eV respectively with 30% of Ni²⁺ and 70% of Ni³⁺.^{49,50} The amount of Ni³⁺ in the γ -NiOOH-Fe suggests that the catalyst surface majorly consists of NiOOH. The positive shift in the binding energy values of Ni²⁺ and Ni³⁺ compared with normal values is due to the binding with the strong Lewis acid Fe³⁺.⁴⁹ The Fe 2p_{3/2} and 2p_{1/2} peaks at 711.2 and 724.6 eV are ascribed

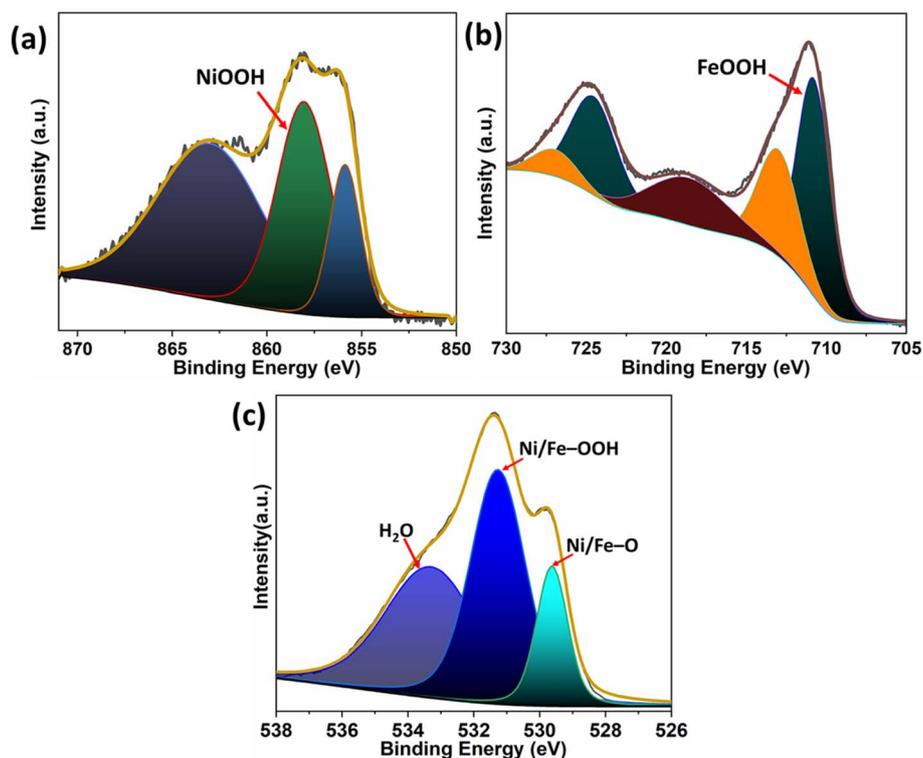


Fig. 2 Deconvoluted XPS spectra (a) Ni 2p, (b) Fe 2p and (c) O 1s spectra of γ -NiOOH-Fe.



to the Fe^{3+} of FeOOH and those at 713.5 and 727.7 eV correspond to the Fe^{2+} in $\gamma\text{-NiOOH-Fe}$.^{50,51} The calculated percentage of Fe^{3+} peaks out of the Fe 2p peak area is around 70% indicating that most of the Fe in $\gamma\text{-NiOOH-Fe}$ is FeOOH . The deconvoluted O 1s spectrum of $\gamma\text{-NiOOH-Fe}$ has three peaks centered at 529.6, 531.3, and 533.2 eV. The peak at 529.6 eV is referred to as the oxygen bound to the metal atom, such as from Ni–O and Fe–O bonds. The peak at 531.3 eV is from the oxygen in the OH^- and that at 533.2 eV corresponds to the oxygen from the chemisorbed water molecules.^{42,52,53} Deconvoluted XPS spectra of Ni 2p, Fe 2p and O 1s of Ni–Fe are presented in Fig. S2.† The Ni $2p_{3/2}$ spectrum shows only the presence of Ni^{2+}

at 854.9 eV.⁴⁹ Fe $2p_{3/2}$ and Fe $2p_{1/2}$ spectra confirm the peaks of Fe at 705.6 and 721.2 eV, Fe^{2+} at 712.7 and 725.3 eV and Fe^{3+} at 710.3 and 723.5 eV.^{54,55} The O 1s spectrum has two peaks corresponding to the metal oxide peak at 529.4 eV and oxygen in OH^- at 530.6 eV.⁵² Compared to the XPS spectra of Ni–Fe, peaks of Ni 2p, Fe 2p and O 1s appeared at higher binding energy because of the presence of a highly oxidized Ni surface.

The surface morphology of the catalysts was analysed by FE-SEM and the elemental composition (Ni, Fe and O) at the surface was measured by EDX. No characteristic surface feature was observed for the bare Ni foam (Fig. 3a); on the other hand, very fine, homogeneous nanoflake-like structures were observed

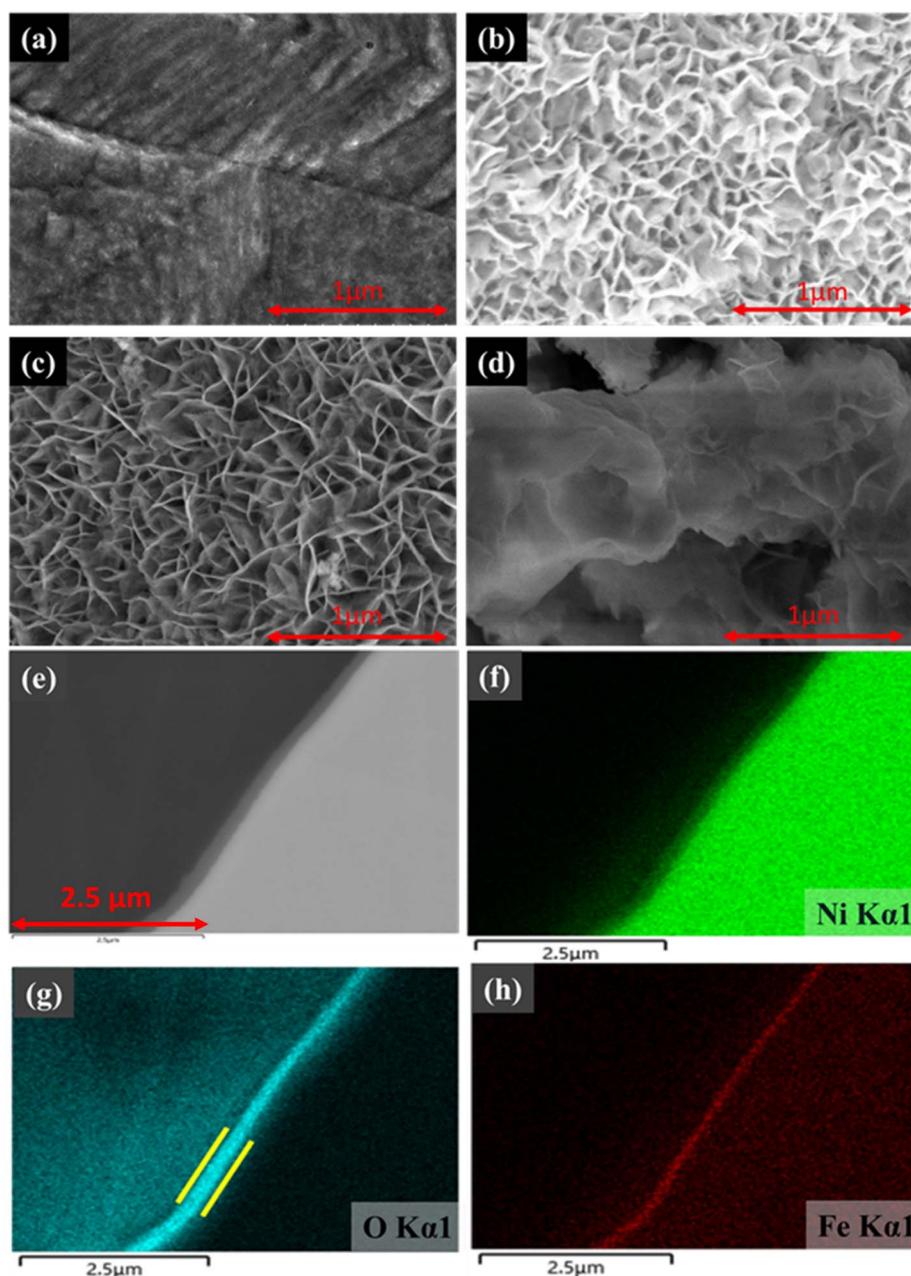


Fig. 3 FE-SEM images of (a) the bare Ni foam surface, (b) Ni foam surface after oxidation ($\gamma\text{-NiOOH}$), (c) $\gamma\text{-NiOOH-Fe}$ and (d) Ni–Fe. (e) Cross-sectional SEM-EDX image of $\gamma\text{-NiOOH-Fe}$, and EDX mapping of (f) nickel, (g) oxygen, and (h) iron.



for the electrochemically oxidized Ni foam due to the formation of γ -NiOOH (Fig. 3b). The nanoflakes grown vertically over the Ni substrate enhance the surface area of the material. Further, when Fe was deposited to create γ -NiOOH-Fe, the surface morphology of the oxidized Ni foam remained the same indicating that the Fe was uniformly deposited over the γ -NiOOH (Fig. 3c). Electrochemical deposition of Fe on the bare Ni foam produced a large petal-like morphology as shown in Fig. 3d. The amount of Fe present on the surface of Ni-Fe and γ -NiOOH-Fe was measured using EDX analysis, and was 17% and 10%, respectively. Cross-sectional SEM-EDX analysis of γ -NiOOH-Fe (Fig. 3e-h) and Ni-Fe (Fig. S3†) catalysts was performed to identify the oxide layer formation. In Fig. 3e, the cross-sectional SEM image of γ -NiOOH-Fe reveals that there is a layer approximately 250 nm thick on the surface, and the oxygen K α 1 map (Fig. 3g) confirms that this 250 nm thick layer is an oxide layer. The Fe K α 1 map of γ -NiOOH-Fe (Fig. 3h) confirms that 50% of the oxide layer contains Fe. In contrast, the cross-sectional SEM-EDX of Ni-Fe (Fig. S3†) shows no such thick oxide layer on the catalyst surface. Consolidating all the material characterisation results, it can be concluded that the γ -NiOOH-Fe catalyst has a γ -NiOOH and FeOOH layer that is \approx 125 nm thick at the top and a \approx 125 nm thick γ -NiOOH layer between the top layer and the bottom Ni substrate.

All electrochemical analyses were performed in 1 M KOH solution to evaluate the electrochemical properties of catalysts. A stabilized voltammogram of Ni foam, Ni-Fe and γ -NiOOH-Fe

was taken with a scan rate of 10 mV s⁻¹ in a deaerated KOH solution, and the results are presented in Fig. 4a. The cyclic voltammogram of Ni-foam exhibited typical redox peaks corresponding to the conversion of Ni(OH)₂ to NiOOH at 1.38 V in the forward scan and back to Ni(OH)₂ at 1.27 V in the reverse scan. The Ni redox peaks of Ni-Fe and γ -NiOOH-Fe were shifted to higher potentials and the peak area and peak current were reduced compared to those of bare Ni, due to the presence of Fe on the catalyst surface.^{56,57} The redox peak of γ -NiOOH-Fe appeared at 1.40 and 1.34 V in the forward and reverse scans respectively. The redox peak of Ni-Fe became more anodic than that of γ -NiOOH-Fe and the oxidation and reduction peaks appeared at 1.41 V and 1.36 V, respectively. This is attributed to the higher surface concentration of Fe in Ni-Fe compared to γ -NiOOH-Fe. The double layer capacitance of the catalysts was measured to calculate the electrochemical surface area. Fig. S4† shows the cyclic voltammogram recorded at different scan rates to measure the double layer capacitance of Ni, Ni-Fe and γ -NiOOH-Fe. As a result of the high surface area, γ -NiOOH-Fe has higher double layer capacitance (1.70 mF) compared to Ni-Fe (1.32 mF) and bare Ni foam (0.6 mF). Further, the OER activity of all catalysts was evaluated by taking the voltammogram, and the current density was normalized with the electrochemical surface area. The OER activity of the catalysts is presented in Fig. 4b. Nickel foam attained 10 mA cm⁻² at 1.61 V. Meanwhile, the potential required to attain the same current density for the Ni-Fe catalyst was 1.49 V. Compared to the other

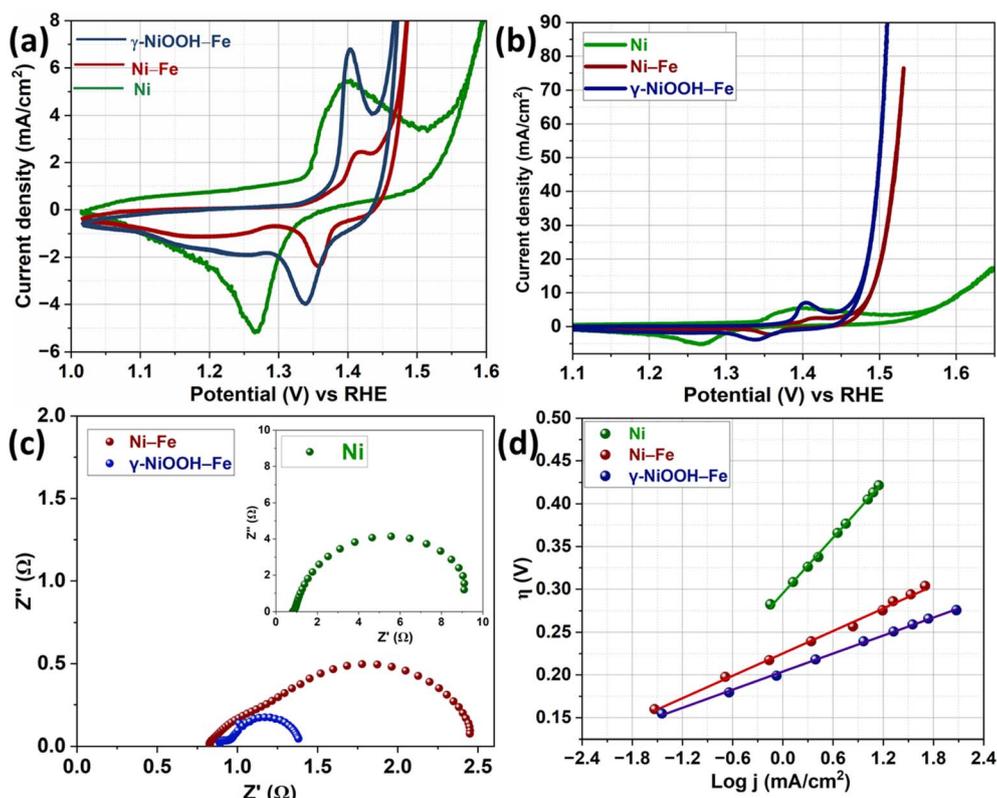


Fig. 4 (a) Cyclic voltammetry responses, (b) OER activity, (c) Nyquist plot measured at 1.5 V, and (d) Tafel slope of Ni, Ni-Fe, and γ -NiOOH-Fe in 1 M KOH at 25 °C.



two catalysts, γ -NiOOH-Fe showed better OER performance by attaining 10 mA cm^{-2} at 1.47 V . Electrochemical Impedance spectra of the catalysts were measured at 1.5 V (Fig. 4c) to analyse the charge transfer kinetics. The low charge transfer resistance of γ -NiOOH-Fe supports its superior OER catalytic activity. Tafel plots of all catalysts were constructed to identify the intrinsic catalytic activity. The potentiostatic method was used to obtain steady state current at different overpotentials to draw the Tafel plot, and the results are presented in Fig. 4c. The γ -NiOOH-Fe has the lowest Tafel slope value of 34 mV dec^{-1} compared to Ni-Fe and Ni with Tafel slopes of 43 and 108 mV dec^{-1} respectively. The lowest Tafel slope of γ -NiOOH-Fe among the OER catalysts confirms that the high OER activity is intrinsic and not due to the surface area. In addition, the turnover frequency (TOF) of the γ -NiOOH-Fe and Ni-Fe was calculated at an overpotential of 300 mV , and the calculation method is given in the ESI.† The number of active sites of the catalysts was calculated from $\text{Ni}^{2+}/\text{Ni}^{3+}$ redox peaks (Ni is considered as the active site for the OER).⁵⁸ The obtained TOF values of γ -NiOOH-Fe and Ni-Fe are 1.3 and 1.1 s^{-1} , respectively. These results indicate that γ -NiOOH-Fe has excellent intrinsic OER activity. The OER activity of γ -NiOOH-Fe is compared with that of other recently reported NiFe based catalysts (Table S1†) to show the significance of this work.

The high intrinsic OER activity of γ -NiOOH-Fe is due to the existence of a synergistic combination of γ -NiOOH and FeOOH on the catalyst surface. In the γ -NiOOH-Fe catalyst, the layer above the Ni surface is γ -NiOOH, in which the Ni exists as $\text{Ni}^{3.6+}$. The surface layer of the catalyst contains the strong Lewis acid Fe^{3+} as FeOOH along with γ -NiOOH. The electrophilic nature of the Fe^{3+} enhances the formation of Ni^{4+} during the polarization.⁴⁰ The Ni^{4+} is crucial for the formation of the important O-O bond for the OER.⁴⁰

The electrocatalytic durability of an OER catalyst is the most crucial parameter to be employed in a water electrolyzer. Initially, the durability of γ -NiOOH-Fe and Ni-Fe catalysts was assessed by an accelerated durability test by cycling the potential between 1.2 V and 1.6 V . Both catalysts exhibited stable OER activity after the accelerated durability test. After the constant current electrolysis, performed at 10 mA cm^{-2} , the activity of Ni-Fe was decreased and the required potential to attain 10 mA cm^{-2} increased to 1.51 V , whereas no change in the activity was observed for γ -NiOOH-Fe (Fig. S5†). Later, constant current electrolysis at 150 and at 500 mA cm^{-2} for 12 hours was performed (Fig. 5a). Furthermore, to evaluate the catalyst stability under the real electrolyzer operational conditions, the electrolyzer was assembled with γ -NiOOH-Fe as anode and $\text{Pt}_{32}\text{Ru}_{16}/\text{C}$ as cathode (Scheme S1†). Although we have inhouse made

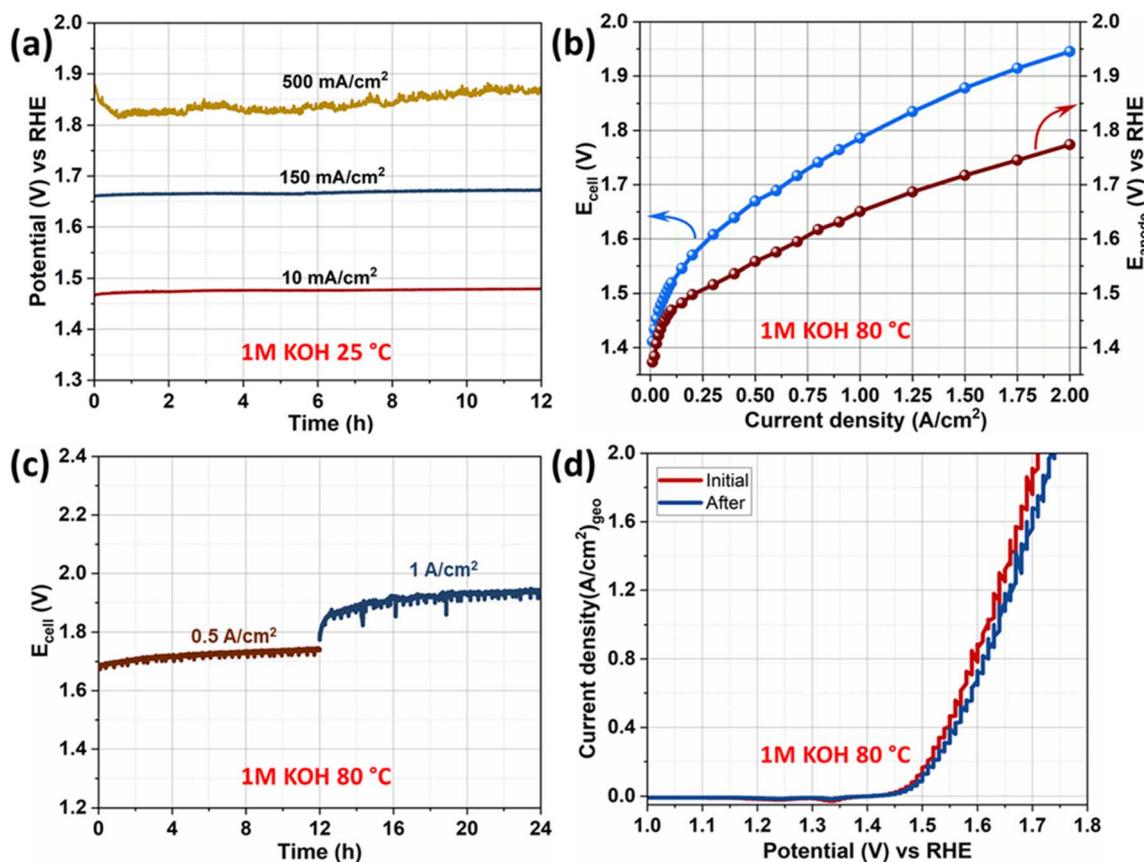


Fig. 5 (a) Chronopotentiometric curves measured at 10, 150 and 500 mA cm^{-2} in 1 M KOH at $25 \text{ }^\circ\text{C}$, (b) polarization curves of the MEA and anode, (c) chronopotentiometric results measured at 0.5 and 1 A cm^{-2} and (d) linear sweep voltammetry responses of γ -NiOOH-Fe before and after the stability tests in 1 M KOH at $80 \text{ }^\circ\text{C}$.



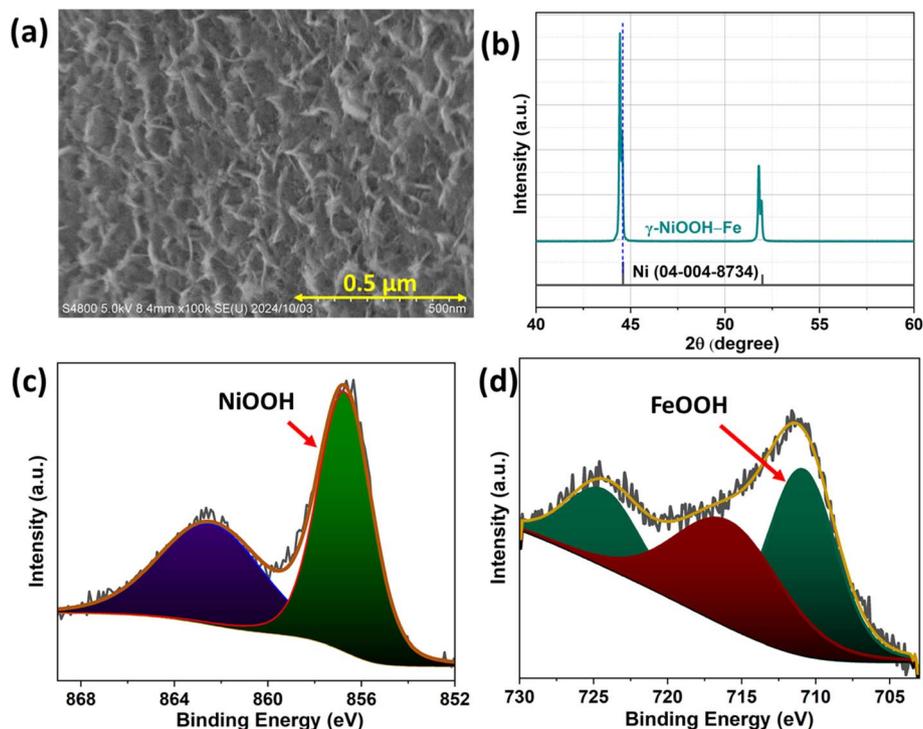


Fig. 6 (a) FE-SEM image, (b) XRD pattern, and (c) and (d) XPS Ni 2p peaks and Fe 2p peaks of γ -NiOOH-Fe respectively, after the electrolysis at 80 °C.

AEMs,^{59–61} in this work we used the commercially available PiperION® AEM, because these results are useful for comparison with other catalysts and with inhouse membranes that will be tested under similar conditions in the near future. Electrolysis was performed at 80 °C in 1 M KOH. Details of electrolyzer setup, activation of membrane electrode assembly, and the method of analysis are provided in the ESI.† During the I - V measurement, the anode potential at different current densities was monitored along with the cell voltage using a reference electrode integrated into the anodic compartment of the electrolyzer and the results are presented in Fig. 5b. Electrolyzer cell voltage (E_{cell}) at 1 and 2 A cm⁻² current density was 1.78 and 1.95 V, respectively and the corresponding anode potential was 1.65 and 1.77 V vs. RHE. Further, the durability of the system was analyzed by constant current electrolysis performed at 0.5 A cm⁻² for 12 hours, followed by 1 A cm⁻² for another 12 hours, and the results are presented in Fig. 5c. Voltammetric response of the γ -NiOOH-Fe anode placed in the electrolyzer measured at 80 °C before and after constant current electrolysis is presented in Fig. 5d. The voltammogram is slightly noisy because of bubble formation. Only a 2.0 mV shift has been observed to attain 1 A cm⁻² current density after the constant current electrolysis indicating the durable OER performance of the γ -NiOOH-Fe electrode.

After electrolysis, SEM, XRD and XPS analyses of the catalyst were conducted to determine the structural and chemical changes of γ -NiOOH-Fe. SEM images show that the surface of the catalyst remains in the nanoflake shape with a difference in the spatial distance between the flakes (Fig. 6a). The XRD

pattern shows no difference in the peaks and peak positions (Fig. 6b). The fine scanning XPS spectra of Ni 2p and Fe 2p of γ -NiOOH-Fe (Fig. 6C & D) reveal that all of the Ni and Fe exists in the NiOOH and FeOOH form. The analysis indicates that although minor structural changes were observed, the nanoflake shape with γ -NiOOH-Fe on the surface was retained after the stability test, suggesting that this structural stability led to the retention of the high electrolyzer performance.

Further, to ensure the scalability of the synthesis of γ -NiOOH-Fe, we made a catalyst with an area of 25 cm² (scale up from 1 cm²), under the same laboratory conditions. The photographs of the modified Ni foam catalyst at different stages are presented in Fig. S6.† Electrochemical oxidation of nickel followed by iron deposition at a controlled potential provides reproducible results. This room-temperature electrochemical route for the preparation of γ -NiOOH-Fe is an industrially viable method because the entire electrochemical process requires less than half an hour compared to the time and energy-consuming conventional heating process.

Conclusions

In summary, we developed a simple, rapid, cost-effective and scalable synthesis of a highly active and durable OER catalyst for AEMWEs. The precise engineering of a layer of γ -NiOOH-FeOOH at the surface and another layer of γ -NiOOH between the surface layer and bottom Ni foam enhances the durability and activity of the catalyst. The catalyst exhibits excellent OER activity in an alkaline medium by attaining 10 mA cm⁻² at



1.47 V ($\eta = 0.24$ V). The low Tafel slope value of 34 mV dec⁻¹ confirms the inherent catalytic activity of the catalyst. The high OER activity of the catalyst might be due to the presence of the high valent Ni (Ni^{3.6+}) layer containing Fe³⁺ at the surface of the catalyst. The Lewis acidic nature of Fe³⁺ endorses the formation of Ni⁴⁺ during the polarization and hence catalyses the critical O–O bond formation in the OER. The catalyst also displayed excellent robustness in its OER catalytic activity under different electrochemical durability analyses including high current density operation under real electrolyzer functioning conditions. The two-step electrochemical method of catalyst preparation is highly reproducible and industrially viable.

Data availability

The authors confirm that the data supporting the findings of this study are available within the article and its ESI.†

Author contributions

The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

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