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# Unlocking OER catalytic potential and chiral Fe<sub>3</sub>O<sub>4</sub> film as a game-changer for electrochemical water oxidation pathway and by-product control†

Wenyan Zhang, \* Chaoqun Jiang, Hangmin Guan, Yuanyuan Wang,   
Yingfei Hu, Wei Wang, Wenjie Tian and Lingyun Hao

Electrochemical water splitting is an attractive technique for hydrogen production, but its development is constrained due to the sluggish kinetics of the oxygen evolution reaction (OER), limited charge transmission efficiency, and the generation of harmful by-products (like hydrogen peroxide) accompanying the OER process. Herein, we constructed an Fe<sub>3</sub>O<sub>4</sub> film with an inherent chiral structure to demonstrate its chiral induced spin selectivity (CISS) effect on modulating the OER pathway and facilitating the transfer of electrons extracted from the OER. Compared with normal Fe<sub>3</sub>O<sub>4</sub>, the chiral Fe<sub>3</sub>O<sub>4</sub> film facilitated the OER pathway for 3  $\sum$  O<sub>2</sub> production via the CISS-induced spin alignment of electrons and radicals, while it suppressed the  $\frac{5}{2}$  pathway for peroxide by-product generation. Meanwhile, as a half-metallic material, chiral Fe<sub>3</sub>O<sub>4</sub> film promoted the transportation of spin-aligned electrons derived from water oxidation to an external circuit. Compared with normal Fe<sub>3</sub>O<sub>4</sub> film which is not capable of aligning electrons, the potential at 10 mA cm<sup>-2</sup> declined by 150 mV, the Tafel slope decreased by 104 mV dec<sup>-1</sup>, the charge transfer resistance decreased by 50%, the free charge carrier density was amplified by 10-fold, and the electron lifetime was prolonged within the chiral Fe<sub>3</sub>O<sub>4</sub> film.

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## 1. Introduction

For decades, the excessive consumption of fossil fuels has led to severe environmental pollution and a global energy crisis, posing a significant threat to the future development of human society.<sup>1–5</sup> Consequently, there is an imperative to explore clean and sustainable energy sources capable of replacing fossil fuels. Among these alternatives, hydrogen has garnered considerable attention due to its status as an environmentally friendly energy carrier, characterized by high enthalpy and absence of carbon emissions (such as CO and CO<sub>2</sub>) during combustion.<sup>6–8</sup>

Electrocatalytic water splitting has emerged as a cost-effective and promising technique for generating hydrogen from water, and it offers the advantage of not requiring extreme conditions like high temperature or pressure.<sup>9–13</sup> A myriad of advanced catalysts for splitting water have been developed, such as MOF-derived catalysts,<sup>14</sup> nickel oxide and nickel phosphide,<sup>15</sup> NiFe selenide,<sup>16</sup> CoNi hydrogen phosphate

nanotubes,<sup>17</sup> nickel nitride,<sup>18</sup> and 2D-layered catalysts like NiCo hydroxide nanosheets.<sup>19</sup>

However, the efficiency of electrochemical water splitting falls short of expectations.<sup>20–23</sup> The primary hurdle in this process lies in the sluggish oxygen evolution reaction (OER) that involves a complex four-electron process.<sup>24–28</sup> Furthermore, an undesirable by-product, hydrogen peroxide, tends to form at the anodes when electrons derived from water oxidation are not spin-polarized. The presence of this by-product poses a detrimental impact on the OER reaction system, as it tends to poison the surface of electrocatalysts, thereby diminishing their catalytic activity and shortening their operational lifespan.<sup>29–31</sup> To facilitate efficient water splitting, it has become imperative to accelerate OER kinetics while mitigating the production of hydrogen peroxide by-products during the OER process.

It is noted that the OER is inherently spin-sensitive, because prior to the formation of O=O, the active sites must facilitate the extraction of three out of four electrons with identical spin orientations. Consequently, electrocatalysts that are capable of ensuring spin selectivity at the catalyst/electrolyte interface and subsequently facilitating the seamless transport of these spin-aligned electrons to the external circuit have become highly desirable for enhancing OER efficiency.<sup>23–25,32,33</sup>

The chiral induced spin selectivity (CISS) effect represents a promising approach to spin control in the OER process.

College of Material Engineering, Jinling Institute of Technology, Nanjing 211169, China. E-mail: wiseyanyan@jit.edu.cn; Fax: +86-25-86188966; Tel: +86-25-86188966

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For example, the attachment of organic chiral molecules (such as proteins or DNA) to semiconductors like TiO<sub>2</sub> or CdSe, induces a **CISS** effect, reducing the OER barrier while suppressing the formation of by-products.<sup>26,27,34</sup> Similarly, Inorganic materials characterized by intrinsic chiral configurations, such as chiral CuO and chiral Co<sub>x</sub>O, have showcased the **CISS** effect on lowering the OER barrier and mitigating the generation of by-products. Notably, their current density can surpass that achieved with semiconductors decorated with organic chiral molecules by up to 1000-fold.<sup>35–37</sup>

As an inorganic material, magnetite (Fe<sub>3</sub>O<sub>4</sub>) is renowned for its exceptional catalytic activity resulting from its redox characteristics, narrow band gap, cost-effectiveness, low toxicity, and remarkable durability.<sup>38,39</sup> It is plausible that incorporating Fe<sub>3</sub>O<sub>4</sub> with an inherent chiral structure could lead to a **CISS** effect on spin control for the OER. Beyond these benefits, Fe<sub>3</sub>O<sub>4</sub> stands out as a half-metallic material, exhibiting nearly 100% spin polarization in the electronic density of states (DOS) near the Fermi energy level. Consequently, the electrical conductivity of Fe<sub>3</sub>O<sub>4</sub> is fundamentally dependent upon the spin states of electrons, with one spin channel exhibiting metallic behavior, while the other spin channel demonstrates insulating properties.<sup>40–45</sup> Such a distinguished feature holds potential for enhancing the transfer of spin-aligned electrons from the catalyst to the external circuit.

To verify these hypotheses, we constructed an Fe<sub>3</sub>O<sub>4</sub> film with an inherent chiral structure to investigate its **CISS** effect on spin alignment for the OER and its effectiveness in transferring spin-aligned electrons to the external circuit. We observed that, compared with normal Fe<sub>3</sub>O<sub>4</sub> film without a chiral structure, the utilization of a chiral Fe<sub>3</sub>O<sub>4</sub> film as an OER electrocatalyst resulted in notable catalytic improvements, encompassing the facilitation of the OER pathway while hindering hydrogen peroxide by-products, a 150 mV reduction in the applied potential required to achieve 10 mA cm<sup>−2</sup>, a 42% increase in current density at 1.8 V (vs. SCE), a decrease of 104 mV dec<sup>−1</sup> in the Tafel slope, a 50% reduction in charge transfer resistance, and a 10-fold amplification in the free charge carrier density. The benefits brought about by the chiral Fe<sub>3</sub>O<sub>4</sub> film are intricately

intertwined with the **CISS** effect induced by its chiral structure and the capabilities of half-metallic materials to facilitate electron transfer.

## 2. Results and discussion

Fig. 1(a) shows the XRD patterns of the chiral Fe<sub>3</sub>O<sub>4</sub> film coated on FTO by electrochemical deposition. (Details are provided in ESI†). The XRD patterns exhibit diffraction peaks corresponding to SnO<sub>2</sub> (JCPDS 2-1337) and Fe<sub>3</sub>O<sub>4</sub> (JCPDS 7-322). The intensity of the Fe<sub>3</sub>O<sub>4</sub> diffraction peak is lower than that of SnO<sub>2</sub>, due to the relatively small amount of Fe<sub>3</sub>O<sub>4</sub> on the FTO substrate. The XPS spectra were acquired to delve into the surface state of chiral Fe<sub>3</sub>O<sub>4</sub> on FTO. In Fig. 1(b), the XPS survey spectra reveal that the surface of chiral Fe<sub>3</sub>O<sub>4</sub> is primarily composed of Fe and O elements. The chiral Fe<sub>3</sub>O<sub>4</sub> film has a regular laminate morphology, as demonstrated in Fig. 2(a). Furthermore, the laminate structure of chiral Fe<sub>3</sub>O<sub>4</sub> is hierarchically constructed from nanoparticles, as Fig. 2(b) shows.

The Fe 2p XPS spectrum of the chiral Fe<sub>3</sub>O<sub>4</sub> film is presented in Fig. 3. Within this spectrum, the binding energy peak centered at approximately 710.6 eV corresponds to the Fe 2p<sub>3/2</sub> state, while the peak at 724.0 eV corresponds to the Fe 2p<sub>1/2</sub> state.<sup>46–49</sup> These peaks are fitted using a shakeup satellite and two spin-orbit doublets, which originate from the Fe<sup>3+</sup> and Fe<sup>2+</sup> states.<sup>46–48</sup> The area ratio of Fe<sup>3+</sup>/Fe<sup>2+</sup> in the spectrum is approximately 2, indicating that the material deposited on the FTO substrate is Fe<sub>3</sub>O<sub>4</sub>.

The chirality of the catalyst surface was assessed through an electrochemical selectivity test, a sensitive technique for discerning the surface chirality of materials (details in ESI†).<sup>50,51</sup> As shown in Fig. 4(a), chiral Fe<sub>3</sub>O<sub>4</sub> exhibited a higher current density and smaller reduction potential when it was scanned in a chiral electrolyte comprising 10 mM L-tartaric acid than in a chiral electrolyte containing 10 mM D-tartaric acid. These results confirmed that chiral Fe<sub>3</sub>O<sub>4</sub> exhibited contrasting chiral selectivity responses in electrolytes with opposite chirality, specifically preferring charge exchanges with L-tartaric acid.

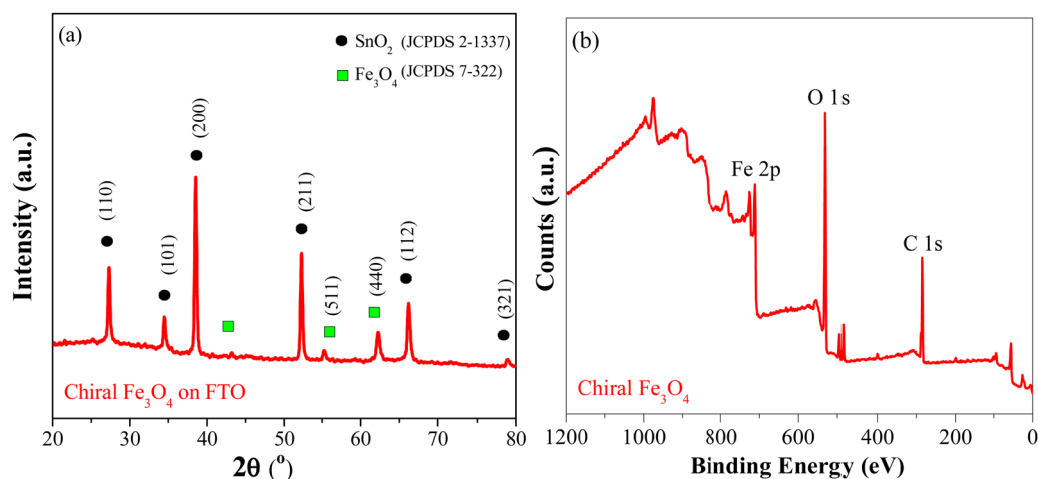


Fig. 1 (a) XRD patterns and (b) XPS survey spectra (b) of chiral Fe<sub>3</sub>O<sub>4</sub>.



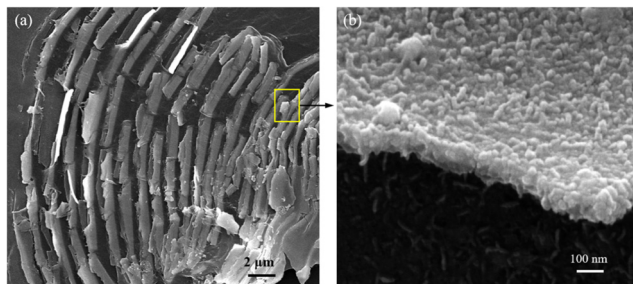


Fig. 2 (a) SEM image of chiral  $\text{Fe}_3\text{O}_4$ ; (b) magnification image of the rectangular region in (a).

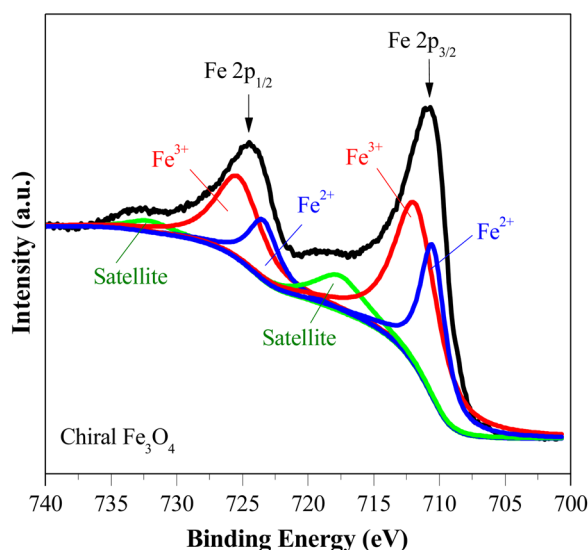


Fig. 3 Fe 2p XPS spectra of chiral  $\text{Fe}_3\text{O}_4$ .

To complement the findings from the electrochemical selectivity test, we further elucidated the surface chirality of the chiral  $\text{Fe}_3\text{O}_4$  film using circular dichroism spectroscopy (CD) and polarized optical microscopy (POM) measurements performed in reflection mode. Fig. 4(b) demonstrates that the chiral  $\text{Fe}_3\text{O}_4$  film prominently displays optical activity within the visible light region, whereas normal  $\text{Fe}_3\text{O}_4$  exhibits no discernible optical activity within this region, suggesting inherent chirality with the chiral  $\text{Fe}_3\text{O}_4$  film. This observation strongly implies the intrinsic chirality of the chiral  $\text{Fe}_3\text{O}_4$  film. The results of POM are shown in Fig. S2 (ESI<sup>†</sup>), which also support the formation of a chiral film.

The chiral  $\text{Fe}_3\text{O}_4$  film, which has inherent chiral structure, demonstrated a CISS effect on spin alignment during the OER. This was verified by monitoring the generation of the singlet-state by-product  $\text{H}_2\text{O}_2$  throughout the OER process.<sup>32–34</sup> The UV-vis spectra of the electrolyte in the water oxidation system are presented in Fig. 5(a), employing *o*-tolidine as an indicator to track the production of  $\text{H}_2\text{O}_2$ . When normal  $\text{Fe}_3\text{O}_4$  was used as an electrocatalyst, an absorption peak emerged at approximately 436 nm, as  $\text{H}_2\text{O}_2$  oxidized *o*-tolidine, leading to the formation of a yellow compound. However, this peak was substantially diminished when the chiral  $\text{Fe}_3\text{O}_4$  film was utilized as an electrocatalyst, signifying the inhibition of  $\text{H}_2\text{O}_2$  generation alongside OER.

As depicted in Fig. 5(b), the OER pathway of  $\text{Fe}_3\text{O}_4$  in alkaline media follows the adsorbate evolution mechanism (AEM), involving a sequential series of concerted four-proton-coupled electron transfer (PCET) processes that take place on active sites. These processes create multiple oxygen intermediates, including  $^*\text{OH}$ ,  $^*\text{O}$ , and  $^*\text{OOH}$ , to establish the pathway,  $^*\text{OH} \rightarrow ^*\text{O} \rightarrow ^*\text{OOH} \rightarrow \text{O}_2$  for the formation of  $\text{O}_2$ . It is recognized that the OER process is spin sensitive, because the four electrons extracted from its four steps should have the same spin direction before the formation of the  $\text{O}=\text{O}$  bond.

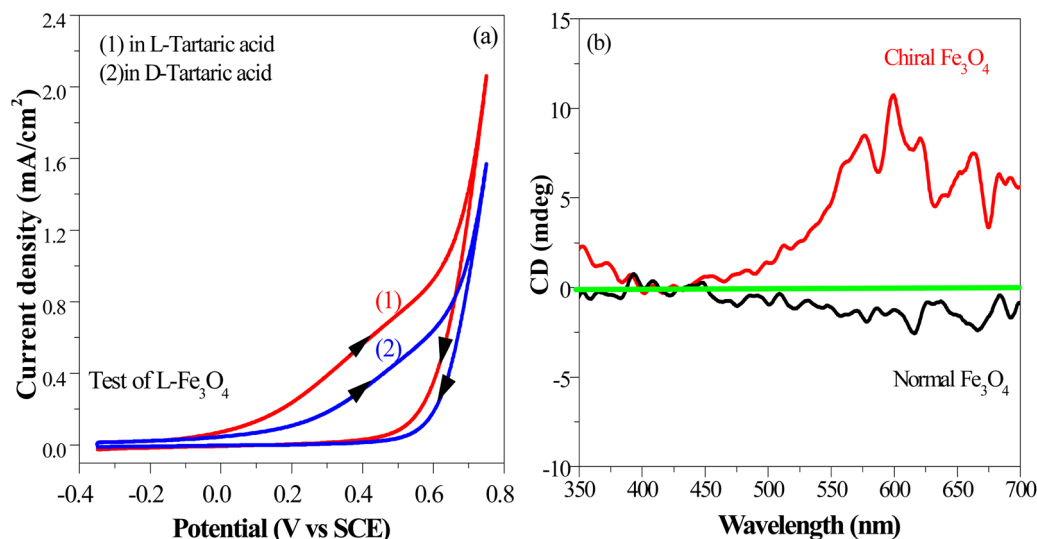
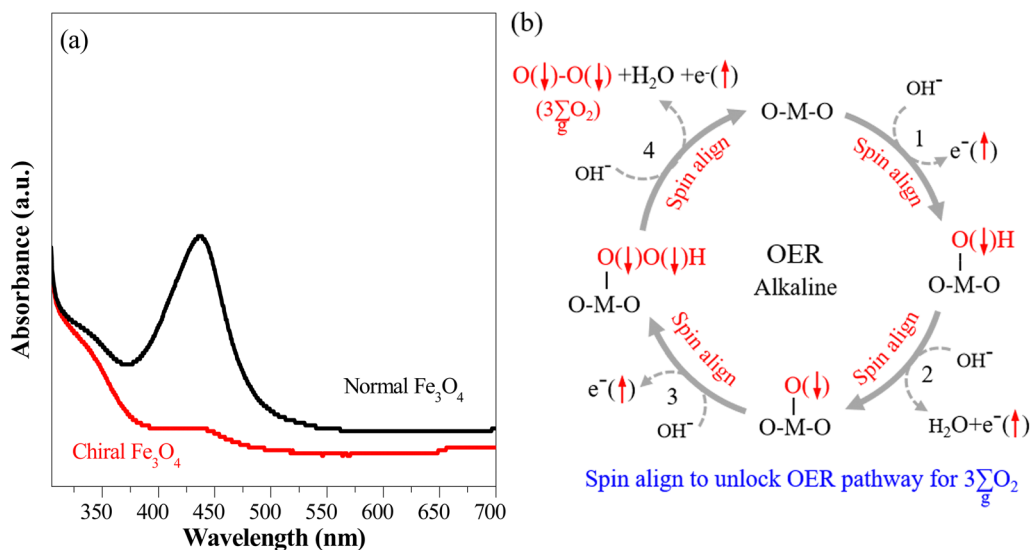


Fig. 4 (a) Electrochemical selectivity test for chiral  $\text{Fe}_3\text{O}_4$  (scanned in a chiral electrolyte involving 10 mM L-tartaric acid and in another chiral electrolyte involving 10 mM D-tartaric acid; arrows indicate scan direction). (b) CD spectra of the chiral  $\text{Fe}_3\text{O}_4$  film and normal  $\text{Fe}_3\text{O}_4$  film.





**Fig. 5** “3Σ<sub>g</sub>O<sub>2</sub>” (a) Visible absorption spectra for the utilized electrolyte titrated with *o*-tolidine as an indicator to assess H<sub>2</sub>O<sub>2</sub> by-product formation in systems employing the chiral Fe<sub>3</sub>O<sub>4</sub> and normal Fe<sub>3</sub>O<sub>4</sub> films as electrocatalysts. (b) Scheme for the **CISS** effect of chiral Fe<sub>3</sub>O<sub>4</sub> on facilitating the water oxidation pathway of 3Σ<sub>g</sub>O<sub>2</sub> formation and forbidding the production of H<sub>2</sub>O<sub>2</sub> during the OER in alkaline electrolyte.

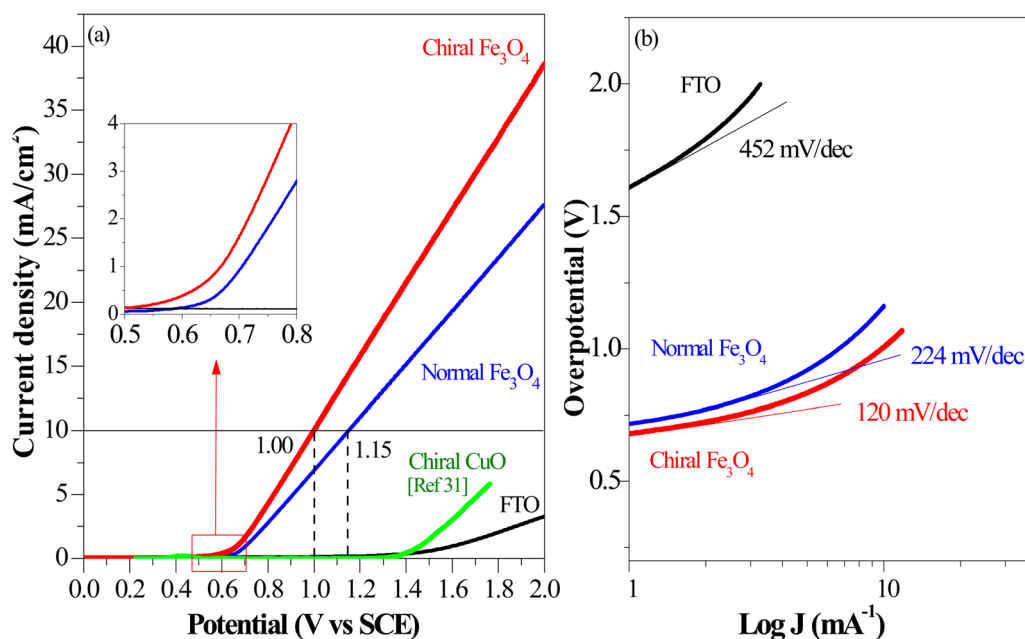
Thereby, the spin orientations of electrons and radicals are beneficial for delivering 3Σ<sub>g</sub>O<sub>2</sub>.

The inhibition of H<sub>2</sub>O<sub>2</sub> by-product suggests that the chiral Fe<sub>3</sub>O<sub>4</sub> film filtered the spin orientation of electrons acquired from water oxidation through the **CISS** effect,<sup>29,30,32,33</sup> synchronously ensuring the alignment of spins in the newly generated •OH radicals that subsequently attached to the M sites of O-M-O species on the Fe<sub>3</sub>O<sub>4</sub> surface during step 1 and step 3 of the AEM process, as demonstrated in Fig. 5(b). This facilitated the

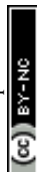
water oxidation pathway for the formation of the triplet-state product, 3Σ<sub>g</sub>O<sub>2</sub> while promoting the alignment of electrons

extracted from water oxidation throughout the four steps of AEM. Conversely, the formation of H<sub>2</sub>O<sub>2</sub>, which occurs on the singlet potential energy surface, was hindered.<sup>29,30,33</sup>

In accordance with the **CISS**-induced preference for the 3Σ<sub>g</sub>O<sub>2</sub> formation pathway during the OER, chiral Fe<sub>3</sub>O<sub>4</sub> exhibited a lower water oxidation barrier than normal Fe<sub>3</sub>O<sub>4</sub>. As



**Fig. 6** (a) Linear sweep voltammetry characterization of chiral Fe<sub>3</sub>O<sub>4</sub>, normal Fe<sub>3</sub>O<sub>4</sub>, FTO, and chiral CuO (the LSV of chiral Fe<sub>3</sub>O<sub>4</sub>, normal Fe<sub>3</sub>O<sub>4</sub>, and FTO were recorded in 0.1 M KOH as electrolyte at room temperature with calomel as reference; the LSV of chiral CuO refers to ref. 31).



depicted in the inset of Fig. 6(a), the LSV onset potential was reduced in comparison with normal  $\text{Fe}_3\text{O}_4$ . Additionally, the potential required to achieve a current density of  $10 \text{ mA cm}^{-2}$  decreased by 150 mV when the chiral  $\text{Fe}_3\text{O}_4$  film was used as an electrocatalyst.

As mentioned above, being a half-metallic material,  $\text{Fe}_3\text{O}_4$  should be fundamentally efficient for transferring spin-aligned electrons, even displaying metallic behavior. Here, it is noticed that the chiral  $\text{Fe}_3\text{O}_4$  and normal  $\text{Fe}_3\text{O}_4$  films demonstrated superior performance in OER current density compared to the chiral  $\text{CuO}$  film, which is not a half-metal, as shown in Fig. 6(a).

In addition, as seen in Fig. 6(a), the current density generated by the chiral  $\text{Fe}_3\text{O}_4$  film at 1.8 V (vs. SCE) was 42% higher than that of normal  $\text{Fe}_3\text{O}_4$ , indicating that the chiral  $\text{Fe}_3\text{O}_4$  film facilitated electron transfer compared to normal  $\text{Fe}_3\text{O}_4$  film. Furthermore, as shown in Fig. 6(b), the Tafel slope of the chiral  $\text{Fe}_3\text{O}_4$  film was  $104 \text{ mV dec}^{-1}$  smaller than that of normal  $\text{Fe}_3\text{O}_4$ , which suggests that the presence of chirality on the  $\text{Fe}_3\text{O}_4$  film enhances OER kinetics and carrier transport. These results demonstrate that, in addition to its CISS effect in promoting the OER pathway for  $3 \sum_g \text{O}_2$  production, the chiral

$\text{Fe}_3\text{O}_4$  film facilitates a more efficient transfer of spin-aligned electrons, derived from water oxidation, from the catalyst to the external circuit compared to the normal  $\text{Fe}_3\text{O}_4$  film.

In line with the findings from LSV and Tafel characterization, the results from electrochemical impedance spectroscopy (EIS) reveal that the chiral  $\text{Fe}_3\text{O}_4$  film exhibited smaller Nyquist circle diameters than normal  $\text{Fe}_3\text{O}_4$  (see Fig. 7(a)). To gain further insights from these Nyquist plots, we employed an R(RQ) circuit model for fitting, and the fitting outcomes are summarized in Table 1. Herein,  $R_\Omega$  represents solution resistance,  $Q_d$  denotes electrochemical double-layer capacitance, and  $R_{ct}$  corresponds to charge transfer resistance. The  $R_{ct}$  value for normal  $\text{Fe}_3\text{O}_4$  was only 6% of that observed for FTO,

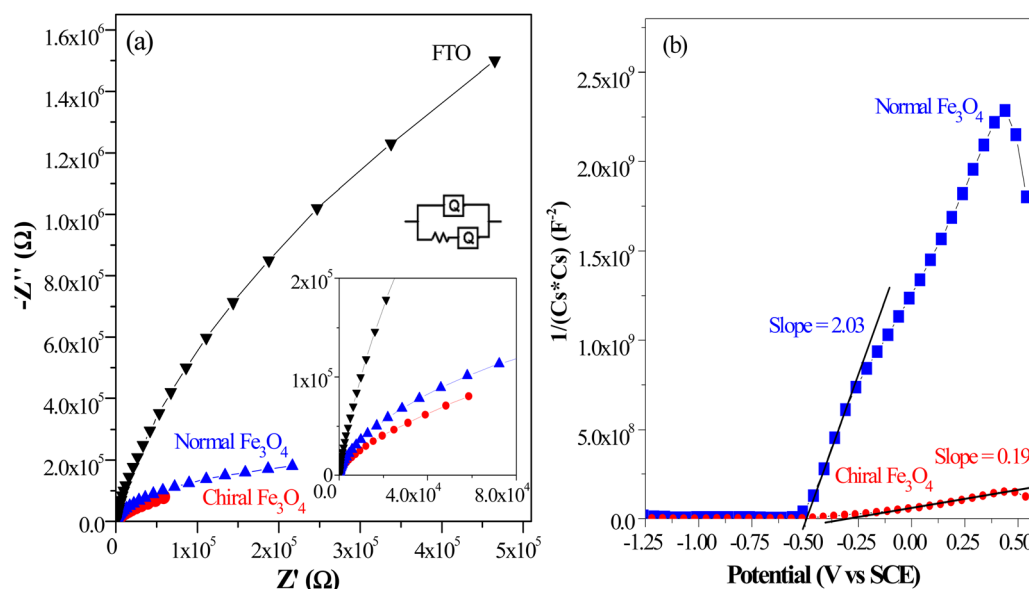
**Table 1** Simulation data of chiral  $\text{Fe}_3\text{O}_4$ , normal  $\text{Fe}_3\text{O}_4$  and FTO with equivalent circuit R(QR)

	$R_\Omega$ ( $\Omega$ )	$Q_d$ ( $\times 10^{-4}$ F)	$n$	$R_{ct}$ ( $\Omega$ )
$\text{Fe}_3\text{O}_4$	42.95	0.2823	0.9434	$3.314 \times 10^5$
Chiral $\text{Fe}_3\text{O}_4$	33.48	1.122	0.95	$1.369 \times 10^5$
FTO	36.45	0.08913	0.9642	$5.135 \times 10^6$

suggesting the significance of  $\text{Fe}_3\text{O}_4$  in promoting electron transfer for the OER. Notably, the chiral  $\text{Fe}_3\text{O}_4$  film exhibits a remarkable 50% reduction in  $R_{ct}$  compared to normal  $\text{Fe}_3\text{O}_4$ , suggesting that electron transport across  $\text{Fe}_3\text{O}_4$  was enhanced due to its intrinsic chiral structure.

Simultaneously, Mott-Schottky measurements provided further evidence of the pivotal role played by chirality in enhancing the catalytic activity of  $\text{Fe}_3\text{O}_4$  film in the OER. The Mott-Schottky equation, expressed as  $(C_{sc})^{-2} = 2(E - E_{fb} - kT/e)/(\epsilon\epsilon_0 A^2 e N_d)$  (details in ESI<sup>†</sup>), dictates that the slope of the linear segments in the Mott-Schottky plot correspond to  $2/(\epsilon\epsilon_0 A^2 e N_d)$ .<sup>52,53</sup> These linear segments are associated with the depleted states of majority carriers located within the space charge region.<sup>54</sup> As depicted in Fig. 7(b), the slope of the linear segment of normal  $\text{Fe}_3\text{O}_4$  was approximately 2.03, while the slope of the chiral  $\text{Fe}_3\text{O}_4$  film was around 0.19, merely 10% of that observed for normal  $\text{Fe}_3\text{O}_4$ . It is calculated that the charge carrier density ( $N_d$ ) of normal  $\text{Fe}_3\text{O}_4$  was about  $3.5 \times 10^{28} \text{ cm}^{-3}$ , while for the chiral  $\text{Fe}_3\text{O}_4$  film, it was approximately  $3.7 \times 10^{29} \text{ cm}^{-3}$ . This result shows that the majority carrier density within the space charge region of  $\text{Fe}_3\text{O}_4$  increased up to 10-fold due to the incorporation of chirality.

Furthermore, evidence from the Bode and phase angle plots serves to reinforce the enhanced electron transport capabilities observed in the chiral  $\text{Fe}_3\text{O}_4$  film. In Fig. 8(a), the Bode plots illustrate that the chiral  $\text{Fe}_3\text{O}_4$  film displays lower impedance modulus values compared to normal  $\text{Fe}_3\text{O}_4$  and FTO. This



**Fig. 7** (a) Nyquist plots of L- $\text{Fe}_3\text{O}_4$ , D- $\text{Fe}_3\text{O}_4$ , normal  $\text{Fe}_3\text{O}_4$  and FTO. (b) Mott-Schottky plots of L- $\text{Fe}_3\text{O}_4$ , D- $\text{Fe}_3\text{O}_4$  and normal  $\text{Fe}_3\text{O}_4$ .



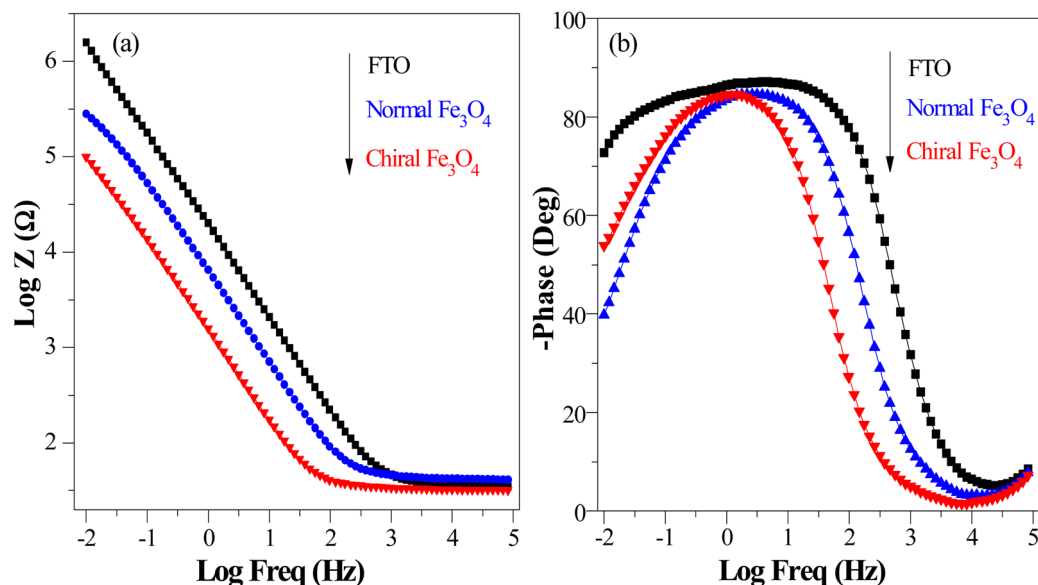


Fig. 8 (a) Bode plots of chiral Fe<sub>3</sub>O<sub>4</sub> film and normal Fe<sub>3</sub>O<sub>4</sub> and FTO. (b) Phase angle plots of chiral Fe<sub>3</sub>O<sub>4</sub> film and normal Fe<sub>3</sub>O<sub>4</sub> and FTO.

observation signifies an increase in carrier mobility on the chiral Fe<sub>3</sub>O<sub>4</sub> film.<sup>55</sup> Moreover, as depicted in Fig. 8(b), the phase angle plot for chiral Fe<sub>3</sub>O<sub>4</sub> shifts towards lower frequencies compared to normal Fe<sub>3</sub>O<sub>4</sub>. This red shift indicates accelerated carrier mobility across the chiral Fe<sub>3</sub>O<sub>4</sub> film. Furthermore, Fig. 8(b) also reveals that the electron lifetime within the chiral Fe<sub>3</sub>O<sub>4</sub> film is prolonged compared to that within the normal Fe<sub>3</sub>O<sub>4</sub> film, taking into account that the electron lifetime ( $\tau_e$ ) can be calculated with the equation  $\tau_e = 1/2\pi f_{\max}$ , where  $f_{\max}$  represents the peak frequency.<sup>49,56</sup>

The elongated electron lifetime implies lower charge recombination probability in chiral Fe<sub>3</sub>O<sub>4</sub> film than in normal Fe<sub>3</sub>O<sub>4</sub> film.

Additionally, hydrogen production in water splitting systems with different OER catalysts was demonstrated, as shown in Fig. 9(a) with chiral Fe<sub>3</sub>O<sub>4</sub> film and normal Fe<sub>3</sub>O<sub>4</sub> film as OER catalysts. The hydrogen production rate with chiral Fe<sub>3</sub>O<sub>4</sub> film as OER catalyst was approximately 20% higher than that with normal Fe<sub>3</sub>O<sub>4</sub> film as OER electrocatalyst and about 5 times of that with chiral CuO film as OER electrocatalyst. Furthermore,

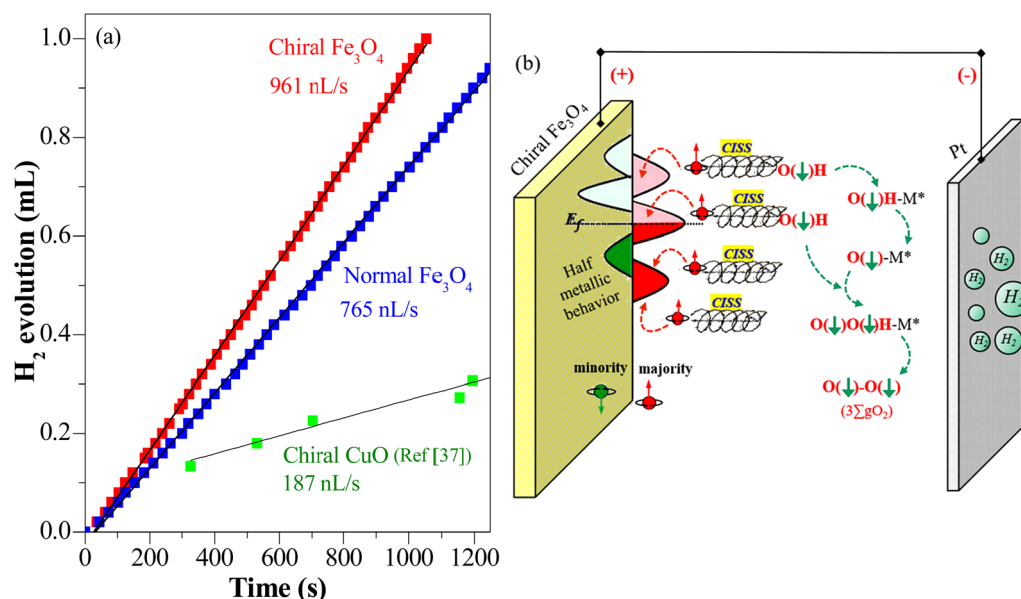


Fig. 9 (a) Hydrogen production recorded in a water splitting system with chiral Fe<sub>3</sub>O<sub>4</sub> and normal Fe<sub>3</sub>O<sub>4</sub> as OER catalysts (conducted at a voltage of 1.5 V (vs. SCE) in 0.1 M KOH electrolyte, at room temperature, with calomel as reference, Pt wire as counter). The hydrogen production of chiral CuO refers to ref. 37. (b) Scheme of the CISS effect in conjunction with the half-metallic nature of chiral Fe<sub>3</sub>O<sub>4</sub> film for aligning the spins of electrons and radicals for the OER and the transportation of spin-oriented electrons.



chiral  $\text{Fe}_3\text{O}_4$  film exhibited robust electrocatalytic activity during the three experimental trials, as shown in Fig. S1 (ESI†). In each trial, the hydrogen production rate was enhanced by at least 20% when chiral  $\text{Fe}_3\text{O}_4$  was employed as electrocatalyst rather than normal  $\text{Fe}_3\text{O}_4$  as electrocatalyst. The growth of the hydrogen production rate is associated with the elevation of water oxidation kinetics and decline of the OER barrier, which was induced by the chiralization of the  $\text{Fe}_3\text{O}_4$  catalyst. The stability also suggests the robustness of the chiral  $\text{Fe}_3\text{O}_4$  film for the OER, as shown in Fig. S4 (ESI†).

The evidence presented above coherently demonstrates the advantages offered by the chiral  $\text{Fe}_3\text{O}_4$  film compared to normal  $\text{Fe}_3\text{O}_4$  film and chiral  $\text{CuO}$ . As illustrated in Fig. 9(b), during the OER, the CISS effect of the chiral  $\text{Fe}_3\text{O}_4$  film aligned the spins of electrons and radicals. This spin-controllable process improved the possibility for the formation of the triplet-state product  $3\sum_g \text{O}_2$ , promoted the AEM water oxidation pathway, and thereby reduced the barrier for the OER. Meanwhile, the half-metallic nature of  $\text{Fe}_3\text{O}_4$  facilitated the transfer of spin-aligned electrons. The combined and synergistic influence of these factors plays a pivotal role in unlocking the catalytic potential of  $\text{Fe}_3\text{O}_4$  film for the OER.

### 3. Conclusion

In summary, this work demonstrated a strategy that harnesses the synergy of chirality and half-metallic materials for unlocking the electrochemical catalytic potential of the OER. Though the current density enhancement of chiral  $\text{Fe}_3\text{O}_4$  over normal  $\text{Fe}_3\text{O}_4$  is not too high, it provides a promising way to facilitate the OER pathway towards  $3\sum_g \text{O}_2$  production through the CISS-induced spin control during AEM. When contrasted with normal  $\text{Fe}_3\text{O}_4$  film which lacks the capability to align electrons, the chiral  $\text{Fe}_3\text{O}_4$  film suppressed the pathway of  $\text{H}_2\text{O}_2$  by-product formation, decreased the potential required to obtain  $10 \text{ mA cm}^{-2}$  current density by 150 mV, reduced the Tafel slope by  $104 \text{ mV dec}^{-1}$ , elevated the current density by 42% at 1.8 V (vs. SCE), lowered the charge transfer resistance by 50%, achieved a 10-fold amplification of the free charge carrier density, and extended the electron lifetime within the catalyst. Furthermore, chiral  $\text{Fe}_3\text{O}_4$  exhibited enhanced OER performance over chiral  $\text{CuO}$  as the transfer of spin-aligned electrons extracted from water oxidation to the external circuit was promoted due to the half-metallic nature of  $\text{Fe}_3\text{O}_4$ .

### Notes

Experimental details and characterization details of this work are shown in our ESI.†

### Data availability

The data that supports the findings of this study are available within the article and its supplementary material. No additional data are available.

### Author contributions

Wenyan Zhang: conceptualization, supervision, writing – reviewing and editing, investigation, project administration, funding acquisition. Chaoqun Jiang: conceptualization, methodology, writing – reviewing and editing, investigation. Hangmin Guan: conceptualization, methodology, writing – reviewing and editing, investigation. Yuanyuan Wang: methodology, investigation, writing – original draft, funding acquisition. Yingfei Hu: methodology, investigation, writing – original draft, funding acquisition. Wei Wang: formal analysis, investigation. Wenjie Tian: visualization, formal analysis. Lingyun Hao: formal analysis, writing – original draft.

### Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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