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## Dual active site-mediated Ir single-atom-doped RuO<sub>2</sub> catalysts for highly efficient and stable water splitting†

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The electronic structure modulation through heterogeneous single-atom doping is an effective strategy to improve electrocatalysis performance of catalysts. Here, Ir single-atom doped RuO<sub>2</sub> (Ir<sub>SA</sub>/RuO<sub>2</sub>) is constructed by substituting Ru sites with mono-disperse Ir atoms in RuO<sub>2</sub> crystals. The Ir<sub>SA</sub>/RuO<sub>2</sub>-850 catalyst shows excellent activity for the hydrogen evolution reaction (HER) and oxygen evolution reaction (OER) in alkaline media, with overpotentials of only 37 and 234 mV respectively, at a current density of 10 mA cm<sup>-2</sup>, lower than that of commercial Pt/C (39 mV-HER) and RuO<sub>2</sub> (295 mV-OER). Notably, no significant degradation occurs during the 1000 h HER stability test at 500 mA cm<sup>-2</sup>. Furthermore, Ir<sub>SA</sub>/RuO<sub>2</sub>-850 also demonstrates superior catalytic activity and stability in acidic media. Theoretical calculations show that the interaction between Ir and RuO<sub>2</sub> modulates the electronic structure of both Ru and Ir sites, resulting in the lowest reaction energy barriers of Ru and Ir sites for the HER and OER, respectively, which thermodynamically explains the enhancement of the catalytic activity. Besides, the introduction of Ir atoms also enhances the demetallation energy of Ru atoms and strengthens the structural stability of the crystal, leading to the improved stability of the catalyst. This work provides an effective strategy for construction of high-performing catalysts by precisely controlling the electronic structure and active sites of polymetal atoms.

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## 1. Introduction

Hydrogen energy, as a clean and high-density energy source, occupies an important position in future energy conversion and energy storage.<sup>1,2</sup> Electrochemical water splitting is a clean and sustainable way of obtaining hydrogen energy, which avoids environmental pollution and the greenhouse effect caused by fossil energy sources.<sup>3,4</sup> However, there are still numerous challenges in the industrialization process of water electrolysis. Currently, platinum (Pt) group noble metal catalysts are widely used in electrolysis, in which Pt and iridium (Ir) based catalysts are often used in the hydrogen evolution reaction (HER) and oxygen evolution reaction (OER), respectively.<sup>4–7</sup> However, the limited availability and high cost of Pt and Ir pose a significant constraint on the widespread application of water electrolysis. Additionally, the slow kinetics of oxygen evolution at the anode restricts the overall water splitting (OWS) efficiency.<sup>8–10</sup> Meanwhile, the long-term stability of catalysts at high current

densities still requires further improvement.<sup>11–13</sup> Therefore, the current research objective is to develop low-cost and high-performing catalysts to support industrial applications.

Ruthenium (Ru), as an affordable material among the Pt group noble metals, has been widely investigated and shown promising applications due to the moderate reaction energy barrier towards the intermediates in the water electrolysis reaction,<sup>14–18</sup> and is expected to replace Pt-based and Ir-based catalysts. Among Ru-based catalysts, ruthenium oxide (RuO<sub>2</sub>) has been identified as the optimal catalyst for OWS. In comparison to iridium oxide (IrO<sub>2</sub>), an anodic OER catalyst based on RuO<sub>2</sub> is more cost-effective with lower driving voltages.<sup>19–21</sup> However, Ru is more prone to undergo oxidation under high voltage conditions, leading to deactivation of catalytically active sites and a decline in catalytic performance.<sup>16,22</sup> Moreover, pure RuO<sub>2</sub> as a HER catalyst still requires high overpotentials to drive the reaction.<sup>19,20,23</sup> Thus, to meet the increasingly stringent requirements under industrial conditions, further in-depth research and optimization of Ru-based catalysts are necessary.

Modulating the electronic structure is an effective modification strategy to optimize the catalytic properties of catalysts,<sup>24–28</sup> which includes the construction of heterostructures,<sup>19,21,29,30</sup> alloy engineering,<sup>31–34</sup> crystal phase engineering,<sup>8,35–37</sup> defect engineering,<sup>25,28,32,38</sup> and

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heterogeneous atomic doping strategies.<sup>5,15,16,20,27,38,39</sup> Among these, heterogeneous single atom doping can precisely adjust the electronic structure of catalysts,<sup>38–41</sup> and synergistically balance the catalytic properties of each component, thereby optimizing the adsorption behavior of catalysts towards reaction intermediates and improving reaction kinetics. It has been reported that Ru atoms as electron acceptors in the RuO<sub>2</sub> system can inhibit the transformation of Ru atoms into higher valence states during the reaction process, further enhancing the stability of the material.<sup>15,22</sup> Generally speaking, atoms with higher electronegativity in the crystal structure are more prone to gaining electrons. Ir has the same electronegativity as Ru but has more electrons outside the nucleus. In particular, the Ir element possesses special physical properties that make it less susceptible to oxidation even at high potentials.<sup>38,42</sup> Furthermore, Ir single-atom doping in RuO<sub>2</sub> is of significant importance to construct RuO<sub>2</sub>-based catalysts, which are expected to exhibit high HER/OER and OWS performance under alkaline conditions.

Here, the Ir single-atom doped RuO<sub>2</sub> (Ir<sub>SA</sub>/RuO<sub>2</sub>) catalyst is synthesized by solvothermal and subsequent pyrolysis processes. Structure characterization results indicate that Ir atoms replace Ru in the crystal lattice of RuO<sub>2</sub> in a single dispersed form. There exists a pronounced interaction between Ir single atoms and RuO<sub>2</sub> as a matrix, which alters the electronic structure of both Ru and Ir sites, endowing the electronic structure-modulated Ir<sub>SA</sub>/RuO<sub>2</sub> catalysts with enhanced HER/OER and OWS activity and stability, superior to those of commercial Pt/C and RuO<sub>2</sub> catalysts. Theoretical calculations further show a significant electron transfer between Ru and Ir atoms, which suggests a strong interaction between Ir and RuO<sub>2</sub>. The electronic modulation of Ru and Ir sites optimizes the adsorption/desorption behavior of reaction intermediates, providing Ru and Ir sites with the lowest rate-determining step (RDS) reaction energy barriers when they act as active centers for the HER and OER, respectively. Furthermore, the introduction of Ir single-atoms enhances the demetallation energy of Ru atoms, thereby stabilizing the catalyst in electrolytic water.

## 2. Results and discussion

### 2.1 Catalyst synthesis

Fig. 1a illustrates the preparation process of catalysts *via* a two-step method. First, 0.2 g of urea and 0.1 g of ammonium fluoride were added to 40 mL of water and 20 mL of ethanol solution until completely dissolved, followed by the addition of 2 mmol RuCl<sub>3</sub> and 0.06 mmol IrCl<sub>3</sub> and stirring until completely dissolved; then the above solutions were transferred to a 100 mL Teflon-lined autoclave and maintained at 160 °C for 10 h, and the sample obtained was centrifuged and dried to obtain the precursor. The obtained precursor was then subjected to heat treatment under different conditions (650 °C, 4 h; 750 °C, 4 h; 850 °C, 4 h; and 950 °C, 4 h) to obtain the final catalysts, which were named Ir<sub>SA</sub>/RuO<sub>2</sub>-650, Ir<sub>SA</sub>/RuO<sub>2</sub>-750, Ir<sub>SA</sub>/RuO<sub>2</sub>-850, and Ir<sub>SA</sub>/RuO<sub>2</sub>-950, respectively.

**Synthesis of RuO<sub>2</sub>.** The precursor is obtained in the same way as described above except that IrCl<sub>3</sub> was not added, and the

synthesized samples were centrifuged as well as dried in a vacuum. Subsequently, the precursor was heat-treated at 850 °C for 4 h to obtain the final RuO<sub>2</sub> catalysts.

### 2.2 Characterization of the catalyst

First, the phase analysis of the samples was performed using X-ray diffraction (XRD), as depicted in Fig. 1b and S1.† It is evident that all diffraction peaks correspond to the standard card phase of RuO<sub>2</sub> (PDF # 40-1290) for pure RuO<sub>2</sub> and Ir-containing RuO<sub>2</sub>, while no discernible peaks related to Ir or its compounds can be observed. The Ir content in the material was determined to be 1.5% by inductively coupled plasma (ICP) analysis, confirming the successful incorporation of Ir. Microscopic morphology and distribution of Ir within the catalyst were characterized using transmission electron microscopy (TEM). Fig. 1c reveals that Ir<sub>SA</sub>/RuO<sub>2</sub>-850 exhibits a nano-spherical morphology with diameters ranging from 50 to 100 nm. The spacing of the lattice fringe is 0.315 nm, consistent with the (110) crystal face of tetragonal RuO<sub>2</sub> (PDF # 40-1290). Furthermore, the morphology of pure RuO<sub>2</sub> (Fig. S3†) has no significant alteration compared to Ir-doped RuO<sub>2</sub> catalysts. Atomic-level mapping images shown in Fig. 1d indicate a uniform distribution of Ru, Ir and O elements on the catalyst surface, in which the Ir element is present in a mono-dispersed form.

For further characterization of the presence and location of Ir, aberration-corrected high-angle annular dark-field scanning transmission electron microscopy (AC HAADF-STEM) was employed. Fig. 1e reveals a periodic uniformly arranged atomic surface without lattice distortion. Furthermore, numerous isolated bright atoms marked by yellow circles are presented with contrast distinct from the surrounding atoms on the surface, which are attributed to Ir atoms due to higher atomic numbers. Atomic line profiling shown in Fig. 1f demonstrates that the bright spots possess stronger peak intensities and larger atomic diameters, further confirming their attribution to Ir atoms. From this, the element Ir exists in single atomic sites by substitution of Ru in RuO<sub>2</sub> crystals.

The electronic structure and coordination environment of Ir were further investigated by X-ray absorption spectroscopy (XAS). The X-ray absorption near edge structure (XANES) at the Ir-L<sub>3</sub> edge (Fig. 2c) reveals a significant positive shift for the white line of Ir<sub>SA</sub>-RuO<sub>2</sub> compared to Ir foil and a weak positive shift compared to the pure IrO<sub>2</sub>. These results suggest that the valence state of Ir is slightly larger than +4, indicating that Ir exists in the form of single atoms while not in the metal state. In addition, the Ir-L<sub>3</sub> edge reveals a positive shift compared to pure RuO<sub>2</sub> combined with the negative shift of the Ru 3p XPS spectrum, indicating an electron transfer from Ir to Ru in the crystal structure. This demonstrates that the Ir atom has a pronounced interaction with RuO<sub>2</sub> as a matrix. The extended X-ray absorption fine structure (EXAFS) of Ir<sub>SA</sub>/RuO<sub>2</sub> (Fig. 2d) presents a prominent peak at 1.2 Å corresponding to the Ir–O first shell coordination, and the peak at 3.2 Å is attributed to the Ir–Ru second shell coordination, and no peaks resemble those of Ir foil. These can prove the absence of Ir–Ir bonds, further confirming the presence of Ir in a single-atom form. Wavelet



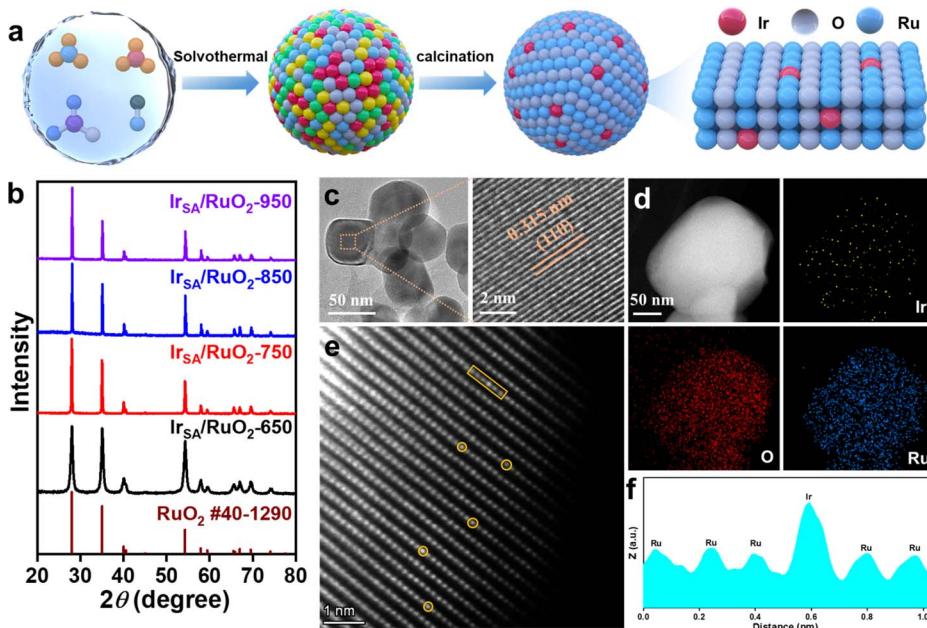


Fig. 1 (a) Schematic diagram for a two-step route to prepare Ir<sub>SA</sub>/RuO<sub>2</sub>; (b) XRD pattern of Ir<sub>SA</sub>/RuO<sub>2</sub>; (c) TEM image of Ir<sub>SA</sub>/RuO<sub>2</sub>-850; (d) HAADF-STEM image of Ir<sub>SA</sub>/RuO<sub>2</sub>-850 and the corresponding elemental mapping images of Ir, O and Ru; (e) AC HAADF-STEM image of Ir<sub>SA</sub>/RuO<sub>2</sub>-850; (f) corresponding line intensity profile of the area in the yellow box in (e).

transformation of Ir<sub>SA</sub>/RuO<sub>2</sub>-850 (Fig. 2e) reveals two characteristic peaks, in which the peak at *R*-space 1.5 Å is attributed to the first shell of Ir–O bonds, while the peak at *R*-space 2.8 Å belongs to the second shell of Ir–Ru bonds, indicating the existence of Ir–O–Ru bonds in the crystal structure. In conclusion, advanced photoelectron spectroscopy techniques confirm the existence of Ir atoms in the form of single atoms within the host crystal structure of RuO<sub>2</sub>.

### 2.3 Electrocatalytic performance evaluation

The HER and OER performances of the catalysts were probed by using a three-electrode system under alkaline conditions. Fig. 3a show the linear sweep voltammetry (LSV) curves of the HER for different catalysts in 1 M L<sup>-1</sup> KOH. It can be seen that the HER activity of Ir<sub>SA</sub>/RuO<sub>2</sub>-850 is superior to that of commercial Pt/C and pure RuO<sub>2</sub>. Moreover, this advantage gradually increases with the increase in current densities. The overpotential of Ir<sub>SA</sub>/RuO<sub>2</sub>-850 is 37/88 mV at current densities

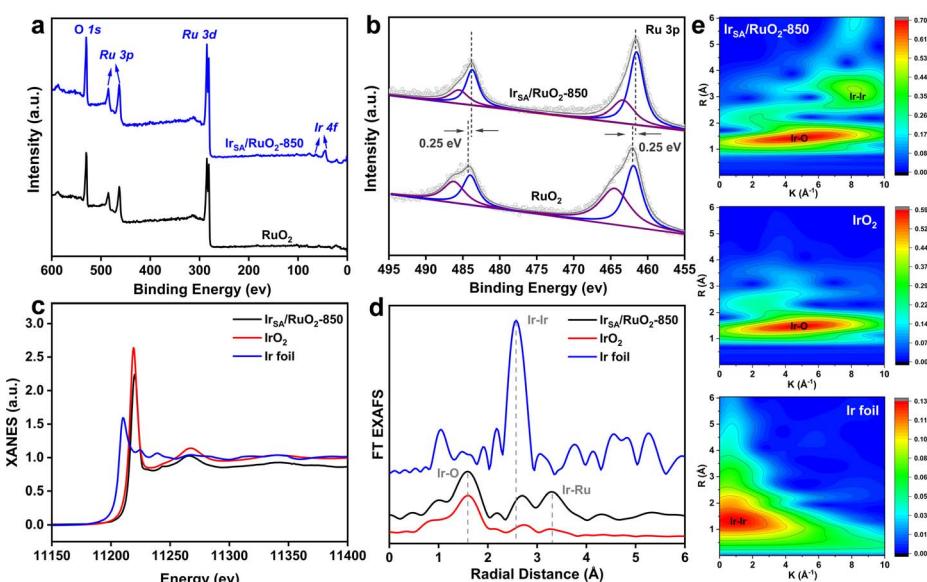


Fig. 2 (a) XPS survey spectra, (b) high-resolution Ru 3p XPS spectra of pure RuO<sub>2</sub> and Ir<sub>SA</sub>/RuO<sub>2</sub>-850, (c) XANES spectra, (d) EXAFS fitting curve and (e) EXAFS wavelet transform plots of Ir foil, IrO<sub>2</sub> and Ir<sub>SA</sub>/RuO<sub>2</sub>-850.

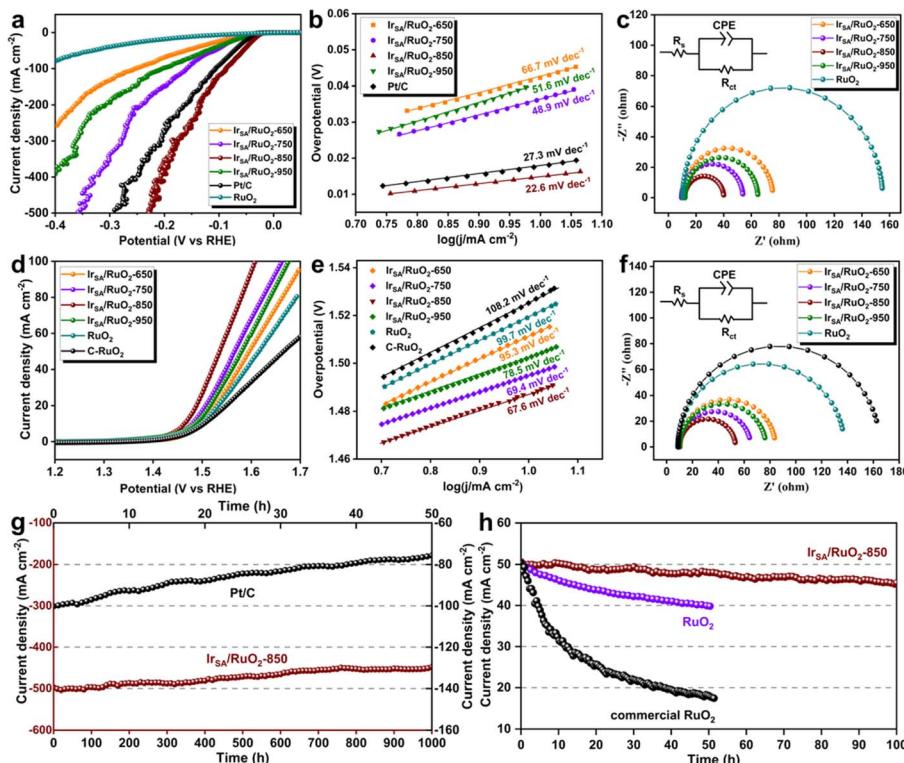


Fig. 3 The performance of catalysts in  $1\text{ M L}^{-1}$  KOH: (a) LSV curves and (b) Tafel slopes of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ ,  $\text{RuO}_2$  and commercial Pt/C; (c) equivalent circuit and fitted EIS curves of the catalysts; (d) LSV curves, (e) Tafel slopes and (f) equivalent circuit and fitted EIS curves of the catalysts, the black curves is  $\text{C}-\text{RuO}_2$ ; (g) HER stability test curves of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 and commercial Pt/C; (h) OER stability test curves of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850,  $\text{RuO}_2$  and  $\text{C}-\text{RuO}_2$ .

of  $10/100\text{ mA cm}^{-2}$ , lower than that of Pt/C ( $39/105\text{ mV}$ ),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -650 ( $64/254\text{ mV}$ ),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -750 ( $48/159\text{ mV}$ ),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -950 ( $52/205\text{ mV}$ ) and pure  $\text{RuO}_2$  ( $138/421\text{ mV}$ ). The corresponding Tafel slope obtained from the LSV curves (Fig. 3b and S6†) demonstrates that  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 possesses the lowest Tafel value ( $22.6\text{ mV dec}^{-1}$ ), significantly lower than that of Pt/C ( $27.3\text{ mV dec}^{-1}$ ),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -650 ( $66.7\text{ mV dec}^{-1}$ ),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -750 ( $48.9\text{ mV dec}^{-1}$ ),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -950 ( $51.6\text{ mV dec}^{-1}$ ) and  $\text{RuO}_2$  ( $218.4\text{ mV dec}^{-1}$ ), indicating the fastest reaction kinetics among the catalysts.

The equivalent circuit and fitted electrochemical impedance spectroscopy (EIS) results of catalysts are shown in Fig. 3c, in which  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 exhibits a smaller  $R_{\text{ct}}$  value ( $30.3\text{ }\Omega$ ) than that of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -650 ( $65.3\text{ }\Omega$ ),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -750 ( $43.8\text{ }\Omega$ ),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -950 ( $54.7\text{ }\Omega$ ) and  $\text{RuO}_2$  ( $153.5\text{ }\Omega$ ), which indicates the fastest charge transfer capability, promoting the HER kinetics. Overall,  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 has the fastest reaction kinetics and the smallest transfer resistance of all prepared catalysts, resulting in the optimal HER performance. Moreover, as shown in Fig. 3g,  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 shows no significant degradation (10%) at a larger current density of  $500\text{ mA cm}^{-2}$  even up to  $1000\text{ h}$  reactions, indicating better stability than Pt/C whose current density which was initially  $100\text{ mA cm}^{-2}$  decays by 50% after  $100\text{ h}$ . This allows  $\text{Ir}_{\text{SA}}/\text{RuO}_2$  to be a potential HER catalyst for industrial applications.

The LSV curve test results for the catalytic OER performance are depicted in Fig. 3d. It is evident that  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 exhibits

an overpotential of  $234\text{ mV}$  at  $10\text{ mA cm}^{-2}$ , surpassing  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -650 ( $271\text{ mV}$ ),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -750 ( $256\text{ mV}$ ),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -950 ( $264\text{ mV}$ ), pure  $\text{RuO}_2$  ( $286\text{ mV}$ ) and commercial  $\text{RuO}_2$  ( $\text{C}-\text{RuO}_2$ ,  $295\text{ mV}$ ). Calculated Tafel slopes (Fig. 3e) show that  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 exhibits the lowest Tafel value ( $22.6\text{ mV dec}^{-1}$ ), significantly lower than that of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -650 ( $95.3\text{ mV dec}^{-1}$ ),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -750 ( $69.4\text{ mV dec}^{-1}$ ),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -950 ( $78.5\text{ mV dec}^{-1}$ ),  $\text{RuO}_2$  ( $99.7\text{ mV dec}^{-1}$ ) and commercial- $\text{RuO}_2$  ( $108.2\text{ mV dec}^{-1}$ ), which demonstrated that  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 maintained the optimal OER process reaction kinetics. The comparison of overpotentials and Tafel slopes of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 with those of the reported Ru-based OER catalysts in alkaline media (Table S1†) demonstrates a performance advantage for  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850.

The equivalent circuit and fitted EIS results of the catalysts are shown in Fig. 3f.  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 possesses a smaller  $R_{\text{ct}}$  value of  $43.7\text{ }\Omega$  than that of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -650 ( $74.2\text{ }\Omega$ ),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -750 ( $54.3\text{ }\Omega$ ),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -950 ( $66.5\text{ }\Omega$ ),  $\text{RuO}_2$  ( $128.7\text{ }\Omega$ ) and  $\text{C}-\text{RuO}_2$  ( $153.3\text{ }\Omega$ ), suggesting that the incorporation of Ir into  $\text{RuO}_2$  does accelerate the charge transfer. Double layer capacitance ( $C_{\text{dl}}$ ) of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 and  $\text{C}-\text{RuO}_2$  was tested by using CV curves in the non-faradaic region (Fig. S7a and b†). As shown in Fig. S7c,†  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 possesses a larger  $C_{\text{dl}}$  value of  $4.15\text{ mF cm}^{-2}$  than  $\text{C}-\text{RuO}_2$  ( $1.96\text{ mF cm}^{-2}$ ). The electrochemical active surface area (ECSA) was deduced from the  $C_{\text{dl}}$ , and the results are displayed in Fig. S7d,† in which the ECSA of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 and  $\text{C}-\text{RuO}_2$  is 69.2 and 32.6, respectively, indicating that  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 has a higher density of active sites, favorable for the adsorption/



desorption of more reactive intermediates, therefore enhancing the catalytic activity. As shown in Fig. S8,† the mass activity of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 (0.223 mA mg Ru<sup>-1</sup>) increases by 2.75 times compared with that of C-RuO<sub>2</sub> (0.081 mA mg Ru<sup>-1</sup>), indicating that the introduction of Ir improves the intrinsic activity of Ru sites.

The stability test results (Fig. 3h) obtained by a chronoamperometry method show that  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 has a significant improvement in the tolerance of the OER, and the current density decreases by only 15% at 100 h constant voltage, surpassing both C-RuO<sub>2</sub> and pure RuO<sub>2</sub> catalysts, with the current density decreasing by 70% and 20%, respectively, after a 50 h chronoamperometry test. To further explore the stability, the dissolved amount of Ru in the electrolyte for  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 was obtained by ICP testing at different time intervals. As shown in Fig. S9,† the dissolution rate of Ru is small for the initial 60 h, and elevated for the next 60–100 h; the trend is similar to the attenuation of the stability test curve (Fig. 3h). After 100 h of stability testing, the amount of Ru dissolved is about 8.8%, indicating no occurrence of significant dissolution of Ru in the catalyst.

After the OER stability test, the performance degradation of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 was assessed by using LSV curves. It can be found from Fig. S10† that, after the stability test, the overpotential of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 is 248 and 327 mV at the current densities of 10 and 50 mA cm<sup>-2</sup>, respectively, which attenuates only 19 and 28 mV from the initial state, demonstrating that  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 can still maintain good catalytic activity after the stability test.

The OER performance test of all prepared samples and C-RuO<sub>2</sub> was further carried out in acidic media (0.5 M H<sub>2</sub>SO<sub>4</sub>) to better evaluate the OER catalytic activity under broad pH

conditions. As shown in Fig. S11a and b,† when the current density reaches 10/50 mA cm<sup>-2</sup>,  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 reveals a lower overpotential (195/249 mV) than C-RuO<sub>2</sub> (246/335 mV),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -650 (228/303 mV),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -750 (211/272 mV),  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -950 (216/291 mV) and RuO<sub>2</sub> (237/312 mV). This also indicates the high OER activity for  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 in acidic media. Besides, it exhibits the lowest Tafel slope and the smallest  $R_{\text{ct}}$  as shown in Fig. S11c and d,† indicating superior reaction kinetics. These properties result in its excellent OER performance in acidic media. Note that, compared with previously reported RuO<sub>2</sub> catalysts tested in acidic media (Table S2†), the as-fabricated  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 catalyst exhibits outstanding performance in terms of overpotential and Tafel slope values toward the OER. The chronoamperometry test results (Fig. S11e†) demonstrate that  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 can operate stably in acidic media for 100 h (only a decrease of 13%), exhibiting superior stability compared to C-RuO<sub>2</sub> (decrease of 70% for 60 h).

Inspired by the excellent HER and OER performance of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850, it was assembled into a two-electrode cell as both the anode and cathode for OWS. The LSV results depicted in Fig. 4a reveal the outstanding alkaline OWS performance of the  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 ||  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 couple, with a potential of 1.504 V at a current density of 10 mA cm<sup>-2</sup>, lower than that of a commercial Pt/C || RuO<sub>2</sub> couple (1.568 V). Stability evaluations as shown in Fig. 4c demonstrate the 100 h stability at 50 mA cm<sup>-2</sup> for the  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 ||  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 couple, with only 10% degradation, outperforming the commercial Pt/C || RuO<sub>2</sub> couple. Additionally, hydrogen and oxygen gases (H<sub>2</sub>/O<sub>2</sub>) were collected in an H-type electrolytic cell equipped with a proton exchange membrane, and the volumes were measured *via* the gas drainage method

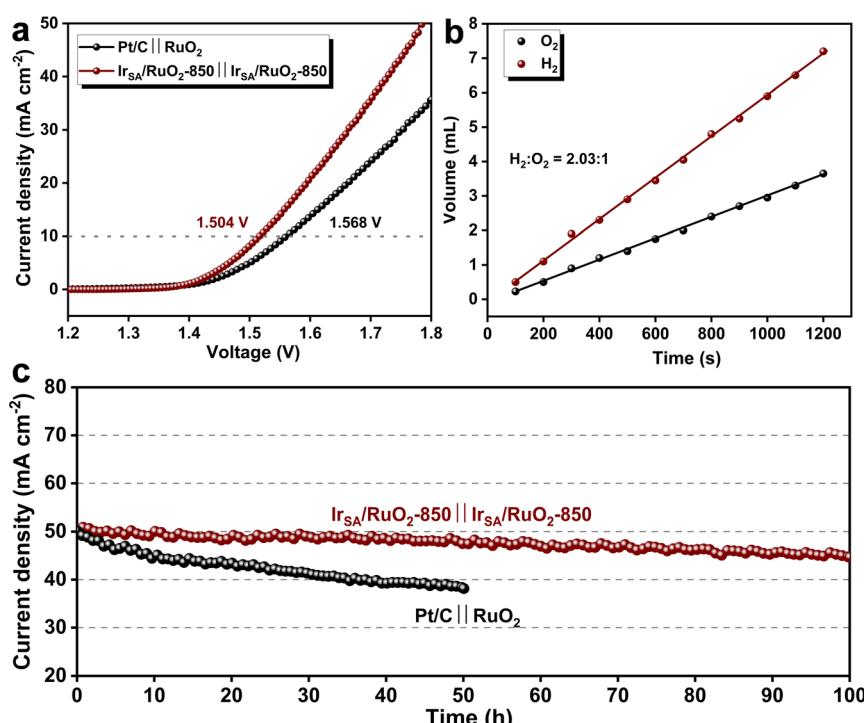


Fig. 4 (a) LSV curves and (c) IT curves of two electrode OWS devices with  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 ||  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 and commercial Pt/C || RuO<sub>2</sub> catalysts; (b) volume of collected H<sub>2</sub> and O<sub>2</sub> versus time.



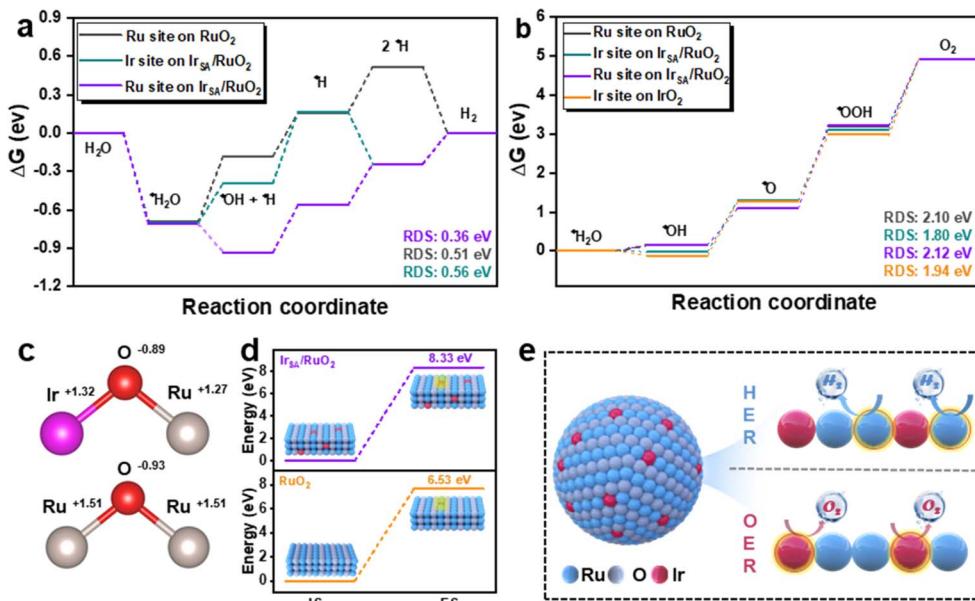


Fig. 5 The Gibbs free energy diagram of (a) the HER; (b) the OER at the Ir or Ru site of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ ,  $\text{RuO}_2$  and the  $\text{IrO}_2$  system; (c) Bader charge of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$  and  $\text{RuO}_2$ . (d) The calculated demetallation energy of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$  and  $\text{RuO}_2$ ; (e) schematic of the  $\text{Ir}_{\text{SA}}/\text{RuO}_2$  catalyst's unique two-site catalysis.

(Fig. 4b). The ratio of  $\text{H}_2$  to  $\text{O}_2$  emitted over a range of time intervals was approximately 2:1, indicating an OWS faradaic efficiency close to 100% for the catalyst. The above results demonstrate that the  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 catalyst possesses a notable reduction in the overall potential for water electrolysis and an enhancement in the stability, which holds significant implications for industrial catalysis towards electrolytic water splitting.

#### 2.4 Density functional theory calculation

The optimization of catalyst performance can be elucidated by density functional theory (DFT) calculations. Fig. S13a and b† show the optimized geometric models for  $\text{RuO}_2$  and Ir single atom doped  $\text{RuO}_2$ . The charge density difference and two-dimensional slice (Fig. S13c†) prove that the introduction of Ir atoms results in a rearrangement of the electronic structure of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ , indicating a strong interaction between Ir and  $\text{RuO}_2$ . Further Bader charge analysis (Fig. 5c) shows that the valence state of  $\text{Ru}^{+1.27}$  in  $\text{Ir}_{\text{SA}}/\text{RuO}_2$  is lower than that of  $\text{Ru}^{+1.51}$  in pure  $\text{RuO}_2$  as well as the  $\text{Ir}^{+1.32}$  in the same structure, suggesting that the Ir acts as an electron donor to transfer electrons to the Ru. This is consistent with the XANES analysis result that the spectral line of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$  slightly shifts towards higher energy compared to those of  $\text{IrO}_2$ , with a negative shift in Ru 3p spectra of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ -850 compared to pure  $\text{RuO}_2$ . In addition, the decrease in the average valence state of Ru is favorable for the enhancement of the OER stability.<sup>12,17</sup> This electronic structure-modulated  $\text{Ir}_{\text{SA}}/\text{RuO}_2$  catalyst can severely affect the adsorption/desorption behavior of reaction intermediates.

To investigate the effect of catalysts with a modulated electronic structure on catalytic performance, the Gibbs free energy of different reaction sites during the HER process to reflect the adsorption intermediates was first calculated. By comparing the RDS barrier of different reaction sites in the HER process

(Fig. 5a), it can be seen that the RDS of the Ru site on  $\text{Ir}_{\text{SA}}/\text{RuO}_2$  is  $\text{OH}^* + \text{H}^*$  to  $\text{H}^*$  with an energy barrier of 0.36 eV, lower than that of the hydrolysis of Ru sites on  $\text{RuO}_2$  (0.51 eV) and the barrier of the reaction intermediates  $\text{OH}^* + \text{H}^*$  to  $\text{H}^*$  of Ir sites on  $\text{Ir}_{\text{SA}}/\text{RuO}_2$  (0.56 eV), suggesting that the introduction of Ir enables the Ru site on  $\text{Ir}_{\text{SA}}/\text{RuO}_2$  to drive the HER with a lower overpotential.

Additionally, the Gibbs free energy of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$ ,  $\text{RuO}_2$  and  $\text{IrO}_2$  catalysts at different reaction sites reflecting the OER intermediates was also calculated. Fig. 5b demonstrates that the RDS of these four reaction sites is all the formation of  $^*\text{OOH}$  from  $^*\text{O}$ , in which the Ir site on  $\text{Ir}_{\text{SA}}/\text{RuO}_2$  acting as the active center has the lowest thermodynamic energy barrier (1.8 eV), lower than that of Ru sites on  $\text{Ir}_{\text{SA}}/\text{RuO}_2$  (2.12 eV) and  $\text{RuO}_2$  (2.1 eV), and Ir sites on  $\text{IrO}_2$  (1.94 eV). This reduction in the reaction energy barrier thermodynamically corroborates the enhancement of OER performance.

To explore the improvement in catalyst stability, the demetallation energy of Ru atoms on  $\text{Ir}_{\text{SA}}/\text{RuO}_2$  and pure  $\text{RuO}_2$  was calculated. demonstrates that the demetallation energy barrier of  $\text{Ir}_{\text{SA}}/\text{RuO}_2$  (8.33 eV) is higher than that of  $\text{RuO}_2$  (6.53 eV), suggesting that the introduction of Ir enhances the crystal stability of  $\text{RuO}_2$ , which in turn boosts the stability in catalytic reactions. Fig. 5e shows the schematic diagram of the unique two-site catalysts in the HER and OER, in which the Ru and Ir sites on the catalyst surface act as the active center for the HER and OER, respectively.

### 3. Conclusions

Here, Ir single atom doped  $\text{RuO}_2$  ( $\text{Ir}_{\text{SA}}/\text{RuO}_2$ ) catalysts were successfully constructed for water electrolysis under alkaline conditions. A series of characterization results show that the



interaction between Ir and the RuO<sub>2</sub> matrix simultaneously changes the electronic structures of Ru and Ir atoms as dual active sites for the hydrogen/oxygen evolution reactions (HER/OER), respectively, resulting in excellent catalytic activity and stability. The overpotential of the Ir<sub>SA</sub>/RuO<sub>2</sub>-850 catalyst at 10 mA cm<sup>-2</sup> is 37 mV for the HER and 234 mV for the OER, superior to that of commercial Pt/C (39 mV-HER) and RuO<sub>2</sub> (295 mV-OER), with steady operation for 1000 h at higher current density (500 mA cm<sup>-2</sup>) during the HER. A two-electrode cell with Ir<sub>SA</sub>/RuO<sub>2</sub>-850 demonstrates superior overall water splitting performance compared to those assembled with commercial Pt/C and RuO<sub>2</sub>. Also, Ir<sub>SA</sub>/RuO<sub>2</sub>-850 demonstrates excellent OER activity (195 mV at 10 mA cm<sup>-2</sup>) and stability in acidic media. Theoretical calculations unveil that the strong interaction between Ir–RuO<sub>2</sub> alters the electronic structure and coordination environment of both Ir and Ru sites, optimizing the adsorption/desorption behavior of the reaction intermediates, with the lowest rate-determining step reaction energy barriers when Ru and Ir sites act as the catalytically active centers of the HER and OER, respectively. Additionally, the enhanced demetallation energy of the Ru atom due to single atom Ir insertion is believed to be a crucial factor contributing to the improved stability of the catalyst. This work provides a strategy for designing highly active and stable catalysts and offers insights for the research and development of catalysts under industrial conditions.

## Data availability

The data supporting this article have been included as part of the ESI.†

## Author contributions

Zhenhua Tao: conceptualization, methodology, writing – original draft, data curation, visualization. Ning LV: methodology, formal analysis. Hongyu Zhao: software. Xu Luo: investigation. Jun Yu: supervision. Lei Chen: project administration. Xupo Liu: validation. Shichun Mu: writing – review & editing, funding acquisition, resources.

## Conflicts of interest

There are no conflicts of interest to declare.

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