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## Recent advances in understanding oxygen evolution reaction mechanisms over iridium oxide

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Water electrolysis driven by renewable energy can produce clean hydrogen, but its efficiency remains low, in part because of slow kinetics at the anode for the oxygen evolution reaction (OER). Learning from the most active catalysts for the OER, iridium oxides, would be the key to the development and establishment of design guidelines for active and stable OER catalysts. This article reviews *in situ* or operando spectroscopic and advanced computational studies in the past decade concerning the OER over iridium oxide for both the oxidation of water molecules and hydroxide ions. By collectively reviewing the reported findings, we illustrate the plausible OER catalytic cycles including the dissolution of iridium during the reaction, which at the same time disclosed discrepancies in the proposed mechanisms. Such discrepancies are thought to originate from variations in the experimental conditions employed in those studies, calling for comprehensive and systematic *in situ* or operando studies in the future. Toward the end, we discuss a recent approach for improving the activity and stability of OER catalysts.

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## 1. Introduction

The transition to a sustainable society largely relies upon the utilization of renewable energy on a large scale. However, low energy density and intermittent availability hamper the widespread implementation of renewable energy, necessitating the

development of systems that allow for its conversion to other forms of energy. The electrocatalytic process is a promising candidate for this purpose, which, using the electricity generated from renewable energy, can convert thermodynamically stable substances such as H<sub>2</sub>O and CO<sub>2</sub> to energy-dense or value-added chemicals.<sup>1,2</sup> Notwithstanding recent advances, the current technology of electrolysis fails to compete economically with fossil fuel-based counterparts,<sup>1</sup> calling for research efforts on the development of efficient and cost-effective electrolysis systems.

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The oxygen evolution reaction (OER) is an anodic half-reaction that can be coupled with various cathodic counterparts, including the hydrogen evolution reaction (HER), the CO<sub>2</sub> reduction reaction, and the N<sub>2</sub> reduction reaction.<sup>3–8</sup> The fundamental study on the OER can be traced back to 1955, when Rüttschi *et al.* first attempted to rationalize the OER performance over a variety of electrodes.<sup>9</sup> Their study assessed the approximate bonding strength of M–OH (where M stands for metal) by analyzing the thermochemical and spectroscopic data,<sup>9</sup> which were plotted with respect to the overpotential at 1 A cm<sup>–2</sup> for the OER reported by Hickling and Hill in 1947.<sup>10</sup> The plot revealed that the OER overpotential linearly scaled with the M–OH bonding energy, suggesting that M–OH bonding energy may serve as a descriptor for the OER activity.<sup>9</sup> Later, this fundamental understanding was put forward by Trasatti in 1980,<sup>9</sup> which revealed a correlation between the OER overpotential and the change of enthalpy from lower to higher oxide transition of transition metal oxides, *e.g.*, from 3+ in Ir<sub>2</sub>O<sub>3</sub> to 4+ in IrO<sub>2</sub>.<sup>11</sup> The disclosed correlation is volcano-shaped; a metal oxide that requires a too large enthalpic change exhibits substantial overpotential, while another metal oxide that requires a too small enthalpic change also demands a large overpotential. This trend indicates that there exists an optimal enthalpy change that minimizes the overpotential for the OER, which was found for iridium oxide and ruthenium oxide having 79.5 and 83.4 kJ mol<sup>–1</sup>, respectively, for the transition from M<sub>2</sub>O<sub>3</sub> to MO<sub>2</sub>.<sup>11</sup> In fact, iridium oxide and ruthenium oxide are regarded even now as one of the most active OER electrocatalysts.<sup>12–14</sup>

These rationalizations of electrocatalysis rely on and are corroborated by experimental evidence. In the first place, the Tafel analysis might be used to elucidate the reaction mechanism and, in particular, to evaluate the rate-determining step (rds), which is the slowest elementary step of a multistep

chemical reaction,<sup>15</sup> of electrochemical reactions.<sup>16</sup> In the case of the OER over iridium oxide, the Tafel slope values under acidic aqueous media were reported to be around 30–45 mV dec<sup>–1</sup>.<sup>17,18</sup> These values, however, could not solely determine the rds because the multiple-step nature of the OER yields a variety of theoretically anticipated Tafel slope values when the electron transfer coefficient deviates from 0.5.<sup>16,19</sup> This fact indicates the need for direct evidence by observing the catalyst surface *via* spectroscopic means to elucidate the surface state of the catalysts, the surface adsorbate on the catalysts, and in turn the reaction mechanism. Early reports in this line of study employed *ex situ* experiments. Typically, the as-made catalysts were characterized by techniques such as X-ray photoelectron spectroscopy (XPS) for the identification of the surface chemical state, electron microscopy for morphologies, and so on. The properties of the catalyst elucidated by these characterization studies were often attempted to be correlated with the catalytic performance. However, such analyses are not capable of providing a solid idea of catalysis because the catalyst material may undergo chemical and morphological changes by being placed in the actual electrolyte environment or during the catalytic reaction, which cannot be seen using characterization studies performed under a vacuum of pre-reaction samples.<sup>20–22</sup> From this viewpoint, it is essential to investigate at least the catalyst of post-reaction. For instance, Kötz *et al.* studied the OER mechanism over iridium oxide using *ex situ* XPS at excitation around the O 1s and Ir 4f regions to track changes of its oxidic nature using post-reaction materials.<sup>21,23</sup> They firstly placed the iridium sample in 1.0 M H<sub>2</sub>SO<sub>4</sub> and applied potentials of 0.0 V, 0.6 V, or 1.25 V *vs.* saturated calomel electrode (SCE) (corresponding to 0.25 V, 0.85 V, or 1.5 V *vs.* reversible hydrogen electrode (RHE) at pH 0.14), before recording the XPS spectra. The obtained spectra captured a decrease in OH contribution and an increase in



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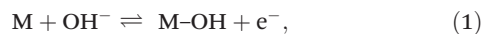
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oxide intensity at increasing anodic potentials applied prior to the XPS measurements. These changes were ascribed to the transition from Ir(OH)<sub>3</sub> to IrO(OH)<sub>2</sub> at 0.22 V vs. RHE that is further oxidized to IrO<sub>3</sub> at 1.5 V vs. RHE,<sup>21</sup> leading to a proposal of a catalytic redox cycle involving Ir<sup>4+/5+/6+</sup>.<sup>21,23</sup> This mechanistic understanding certainly helps in understanding the catalytic cycles. Nevertheless, nowadays, it is commonly considered that such *ex situ* measurements would not be satisfactory to fully rationalize the catalysis; *e.g.*, the catalyst may attain a surface state different from its pristine state only during working conditions.<sup>23–25</sup> This gap between the catalyst states under vacuum and working conditions hampered the establishment of a solid understanding of electrocatalysis.

From a different perspective, computational studies have advanced our understanding of OER electrocatalysis at a molecular level. A milestone report was published by Rossmeisl *et al.*<sup>5,26,27</sup> Their framework of density functional theory (DFT) calculations considered the concerted proton–electron transfer (CPET) steps, where one H<sup>+</sup> and one e<sup>−</sup> are transferred in a single kinetic step:<sup>28</sup>

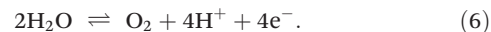


For each step, they computed the free energy changes of the reaction, as well as the binding energy to the surface intermediates.<sup>29</sup> Understanding the changes of the free energy allowed for identifying the potential-determining step (pds) that requires the largest theoretical overpotential among steps and thus is the theoretically predicted bottleneck of the OER.<sup>29</sup>

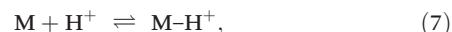
In addition, their study disclosed linear relationships among the binding energies of M, M–OH, M–O, and M–OOH, the so-called linear-scaling relationship.<sup>29</sup> Based on these findings, the theoretical OER overpotential was plotted as a function of the binding energy to the surface intermediates, *e.g.*, M–O, which was found to be volcano shaped.<sup>29</sup> The implication of this figure is that a surface that binds the reaction intermediates neither too strongly nor too weakly possesses the optimal surface, achieving the highest OER performance.<sup>29</sup> This plot demonstrated that iridium oxide and ruthenium oxide sit near the top of the volcano trend, thus corroborating the experimental observations.

Notwithstanding the success of the past computational approaches, the employed model omitted several considerations for the sake of simplicity and versatility. Firstly, those models did not compute kinetics. Analysis of the free energy of the reaction intermediates to elucidate the bottleneck was justified by the linear scaling relationship between the reaction energy and the activation energy, the so-called Brønsted–Evans–Polanyi (BEP) relationship.<sup>30</sup> However, the BEP relationship is not always valid, and the pds and rds can differ from each other.<sup>31</sup> Further complicating this kinetic aspect is the variation in the OER reactant. Depending on the electrolyte pH

levels, the OER can proceed either as the oxidation of the hydroxide ion (OH<sup>−</sup>) or the water molecule (H<sub>2</sub>O), as shown in eqn (5) or eqn (6), respectively:<sup>32,33</sup>



Dissociation of the O–H bonding of H<sub>2</sub>O molecules is kinetically more difficult than that of the OH<sup>−</sup> reactant, which would lead to reactant switching from H<sub>2</sub>O to OH<sup>−</sup> when shifting the pH from near-neutral to alkaline values.<sup>34,35</sup> In fact, such reactant switching was experimentally observed at around pH 10–11.<sup>19,36</sup> Secondly, the surface state of the catalyst can vary depending on the reaction environment, which was also not considered in the simplified model. For instance, due to the acid–base equilibria at the catalyst surface under the reaction conditions, the surface is deprotonated [protonated] when the pH of the electrolyte solution is larger [smaller] than the pK<sub>a</sub>.<sup>37</sup> Likewise, the surface is positively [negatively] charged when the pH of the electrolyte solution is smaller [larger] than the isoelectric point (IEP) of the oxide. These changes in the surface state of catalysts can lead to subsequent changes in the reaction mechanism.<sup>38</sup> Thirdly, while the introduction of the computational hydrogen electrode (CHE) has simplified the treatment of potential, which makes this DFT-based approach quite versatile,<sup>5,26,27</sup> the model cannot consider the decoupled pathways. More specifically, although the reaction scheme of eqn (1)–(4) merely contains CPET, the reactions possibly proceed by a sequence of decoupled proton transfer (PT) or electron transfer (ET), which is the transference of only H<sup>+</sup> or only e<sup>−</sup>, respectively:<sup>28,39</sup>



All in all, while the model without these considerations has been successful in rationalizing the general trend for electrocatalysis, further detailed understanding and development of active electrode materials require sophistication and complication of the framework.

In this context, the recent decade has witnessed the development of spectroscopic techniques that enable the direct observation of the surface state *in situ* or operando<sup>14,40</sup> and the advancement of computational calculations that can simulate and map the catalytic cycles in more detail.<sup>41</sup> This review herein discusses the recent progress in understanding the OER mechanism based on *in situ* or operando experimental evidence as well as computational results. We limit ourselves to iridium oxide, one of the most active OER electrocatalysts. Readers interested in other catalysts can refer to other published studies.<sup>42–44</sup> We illustrate plausible OER cycles *via* H<sub>2</sub>O oxidation in section 2 and review its spectroscopic and computational support, while section 3 deals with those for OH<sup>−</sup> oxidation. Section 4 discusses the dissolution mechanisms of iridium during the OER mechanism, corroborating the catalytic cycle.



## 2. OER mechanism via H<sub>2</sub>O oxidation over iridium oxide

Iridium oxide sits at or near the top of the volcano-shaped trend for the OER,<sup>11,29</sup> and has been employed in commercial polymer electrolyte membrane (PEM) electrolyzers despite its high cost and scarcity.<sup>45</sup> Developing active OER catalysts composed of Earth-abundant and cost-effective elements is the key to cost reduction and thus the introduction of hydrogen *via* electrolysis in the market. The rational design of such catalysts would require identification of the reaction mechanism and the working state of the best-performing iridium oxide catalysts. Accordingly, the recent decade has witnessed significant advancements in the understanding of OER electrocatalysis. Herein, this section collectively reviews the proposed mechanisms of H<sub>2</sub>O oxidation over iridium oxide based on spectroscopic evidence and advanced theoretical calculations. We firstly introduce an overall picture of plausible OER catalytic cycles, and subsequently review the spectroscopic and computational data supporting the scenarios in the later subsections.

### 2.1. Proposed mechanism of H<sub>2</sub>O oxidation over iridium oxide

The existing literature allows for the elucidation of OER catalytic cycles over iridium oxide, illustrated in Fig. 1. There have been four reported cycles: (1) a cycle involving a surface adsorbate driven by the redox of the iridium centre (red colored in Fig. 1a), (2) a cycle involving a surface adsorbate driven by the redox of the adsorbed O species (blue colored in Fig. 1b), (3) a cycle involving the redox of the Ir=O state (green colored in Fig. 1c), and (4) the one that involves lattice oxygen (orange colored in Fig. 1d). Below we describe each cycle in more detail.

The scenario of the red-colored cycle in Fig. 1a starts with the oxidation of the initial state, Ir<sup>3+</sup> (1 in the figure), with the H<sub>2</sub>O reactant, forming Ir<sup>4+</sup>-O<sup>2-</sup>-H surface species (2) and H<sup>+</sup> *via* CPET. This species is oxidized to the Ir<sup>5+</sup>=O<sup>2-</sup> state (3) *via* a CPET step, which upon further oxidation is converted into Ir<sup>4+</sup>-O<sup>2-</sup>-OH (4). By releasing O<sub>2</sub>, this state relaxes to Ir<sup>3+</sup> (1). This cycle is consistent with the single-site mechanism commonly adopted for DFT calculations.<sup>16,50,54</sup>

The second scenario (blue colored in Fig. 1b) is similar to the first one except that the surface redox species is not only iridium but also oxygen. Firstly, the Ir<sup>3+</sup> (1) is attacked by the H<sub>2</sub>O reactant, and forms Ir<sup>4+</sup>-O<sup>2-</sup>-H surface species (2) releasing H<sup>+</sup> *via* CPET. This species is oxidized *via* a CPET step into the Ir<sup>4+</sup>=O<sup>1-</sup> state (6), which contains the electrophilic oxygen species (O<sup>1-</sup>) as a reactive intermediate.<sup>48-51</sup> Ir<sup>4+</sup>=O<sup>1-</sup> experiences further oxidation with a H<sub>2</sub>O molecule to form Ir<sup>4+</sup>-O<sup>2-</sup>-OH (4) *via* a CPET step. This state relaxes to Ir<sup>3+</sup> (1) with releasing O<sub>2</sub> *via* CPET.

In contrast to the first two scenarios that involve the redox of either iridium or oxygen, the green-colored cycle in Fig. 1c proceeds by the redox of the Ir=O states. The initial Ir<sup>4+</sup>=O (8) changed the oxidation state to Ir<sup>5+</sup>=O (3) *via* ET. This Ir<sup>5+</sup>=O species (3) is the precursor to the reactive oxyl

species<sup>31,52-57</sup> of Ir<sup>4+</sup>-O<sup>1-</sup> (7) that reacts to form Ir<sup>3+</sup>-O<sup>1-</sup>-OH (4) *via* PT. The formed Ir<sup>3+</sup>-O<sup>1-</sup>-OH (4) releases O<sub>2</sub> with H<sub>2</sub>O reactant and relaxes to Ir<sup>4+</sup>=O (8) *via* CPET. This route likely appears at a high overpotential (*e.g.*, >2.4 V *vs.* RHE),<sup>58</sup> where the accumulation of charge is considered to occur.<sup>59</sup>

The last cycle (orange-colored) shown in Fig. 1d proceeds *via* the releasing and filling of an oxygen vacancy at the iridium oxide surface, which was named the lattice oxygen evolving reaction (LOER) mechanism or lattice oxygen participated mechanism (LOM).<sup>60,61</sup> This route likely appears at highly anodic potentials (*e.g.*, >1.6 V *vs.* RHE for electrochemically prepared oxide from metallic iridium).<sup>62,63</sup> In this scenario, two pathways are considered depending on the number of lattice oxygen atoms participating in the reaction. The first one depicted in the outer circle starts with the oxidation of Ir<sup>3+</sup>=O (1) to form O-Ir<sup>4+</sup>-O (9) *via* ET. Lattice O bridging iridium sites of this O-Ir<sup>4+</sup>-O (9) are attacked by the H<sub>2</sub>O molecule to form O-Ir<sup>5+</sup>(-OH)-O (10) *via* PCET, which transforms into O-Ir<sup>6+</sup>(=O)-O (11) *via* CPET. This species releases oxygen to form O-Ir<sup>4+</sup>-□ species (12) where □ represents a vacant site, and this formed vacancy is subsequently filled by the attack of H<sub>2</sub>O to form O-Ir<sup>4+</sup>-O (9) *via* PCET. In the second pathway depicted in the orange-colored inner circle of Fig. 1d, evolved O<sub>2</sub> consists of two lattice oxygens.<sup>64</sup> In detail, the lattice O bridging the iridium sites (9) leaches out from the lattice to form □-Ir<sup>6+</sup>(=O)-O (13). Subsequently, this □-Ir<sup>6+</sup>(=O)-O state (13) releases O<sub>2</sub> to form □-Ir<sup>4+</sup>-□ (14), where the two oxygen vacancies are filled by the attack of H<sub>2</sub>O to form O-Ir<sup>4+</sup>-O (9) *via* CPET.

### 2.2. Spectroscopic evidence to support the claimed catalytic cycle for H<sub>2</sub>O oxidation over iridium oxide

The OER catalytic cycles detailed in the previous section were obtained based on the recent *in situ* and operando characterization studies, which allowed for determining the working states of catalysts.<sup>65</sup> The employed techniques include X-ray absorption near-edge structure (XANES), near-edge X-ray absorption fine structure (NEXAFS), X-ray absorption (XAS), XPS, and Raman spectroscopy. The key observations in the literature are highlighted and shown in Fig. 2 that elucidated OER electrocatalysis over iridium oxide.

Initially, the spectroscopic evidence supporting the catalytic cycle in Fig. 1a was provided by an *in situ* XPS study in 2014.<sup>66</sup> The group of Nilsson conducted *in situ* XPS measurements using IrO<sub>2</sub> nanoparticles with the applied potential of 1.75 V *vs.* RHE in the presence of 10 Torr of H<sub>2</sub>O, *i.e.*, neither acidic nor alkaline.<sup>66</sup> By analyzing the XPS spectra in the Ir 4f region, they found that the IrO<sub>2</sub> sample at open-circuit potential (OCP) retained the Ir<sup>4+</sup> state upon exposure to H<sub>2</sub>O. Interestingly, the spectra exhibited a shoulder when an anodic potential of 1.75 V *vs.* RHE was applied, ascribed to the appearance of Ir<sup>5+</sup>. In the same study, they also examined the excitation around the O 1s region, and found that when the potential was switched from OCP to 1.75 V *vs.* RHE, the relative intensity of the peak ascribable to hydroxide species decreased while that of oxide increased.<sup>66</sup> Taken together, their observation pointed to a







**Fig. 2** Spectroscopic data for  $\text{H}_2\text{O}$  oxidation over iridium oxide. (a) *In situ* Ir-L<sub>III</sub> X-ray absorption near-edge structure (XANES) spectra using hydrous iridium oxide with the applied potential from 0.2 to 1.6 V vs. reversible hydrogen electrode (RHE) in 0.5 M  $\text{H}_2\text{SO}_4$  solution. White lines of the absorption in the Ir-L<sub>III</sub> region are shown for iridium oxide at various potentials in the panel (i), and over the reference  $\text{IrCl}_3$  and  $\text{IrO}_2$  in panel (ii), and the second derivatives of the spectra shown in panel (i) are plotted in panel (iii). Reproduced from ref. 67 with permission from the Royal Society of Chemistry. (b) *In situ* near-edge X-ray absorption fine structure (NEXAFS) spectra in the O K-edge region in 0.1 M  $\text{H}_2\text{SO}_4$  at 0.3 Pa at open circuit potential (OCP), 1.6, 1.8, and 2.0 V vs. standard hydrogen electrode (SHE) (corresponding to 1.66 V, 1.86 V, or 2.05 V vs. RHE at pH 0.97) using a sputtered iridium catalyst. Reproduced from ref. 48 with permission from the Royal Society of Chemistry. (c) *In situ* shell-isolated nanoparticle-enhanced Raman spectroscopy (SHINERS) spectra using electrochemically deposited iridium oxide in 0.1 M  $\text{NaClO}_4$  solutions at pH 10 at varying potentials from OCP to 1.8 V vs.  $\text{Ag}/\text{Ag}^+$  (corresponding to 2.6 V vs. RHE at pH 10). Reproduced from ref. 58 with permission from the Royal Society of Chemistry. (d) Differential electrochemical mass spectrometry (DEMS) results during a cyclic voltammogram in a potential window from ca. 0.05 V to 1.6 V vs. SHE (corresponding to ca. 0.05 V to 1.6 V vs. RHE at pH 0) in a 1 M  $\text{HClO}_4$  solution containing 10 wt%  $\text{H}_2^{18}\text{O}$  over thermally prepared  $\text{Ir}^{16}\text{O}_2/\text{Ti}$  catalyst. Reprinted with permission from ref. 65. Copyright (2018) Elsevier.

( $t_{2g} + e_g$ ) 5d states, indicating the presence of  $\text{Ir}^{4+}$  and  $\text{Ir}^{5+}$  at 0.7 V and 1.3 V vs. RHE, respectively.<sup>67</sup> Collectively, their study proposed that iridium oxide changes its oxidation state from  $\text{Ir}^{3+}$  to  $\text{Ir}^{4+}$  at around 1.0 V vs. RHE and further to  $\text{Ir}^{5+}$  above 1.3 V vs. RHE (from 1 to 3 in Fig. 1), thus supporting the red- and green-colored cycles displayed in Fig. 1a.

In contrast, a study was reported in 2016 by Schlögl and co-workers that supports, based on NEXAFS, the anion redox mechanism shown as the blue-colored second scenario in

Fig. 1b.<sup>48–51</sup> The group recorded *in situ* NEXAFS spectra in the O K-edge region using a sputtered iridium catalyst at varying potentials of OCP, 1.6, 1.8, and 2.0 V vs. standard hydrogen electrode (SHE) (corresponding to 1.66 V, 1.86 V, and 2.05 V vs. RHE at pH 0.97, respectively) in 0.1 M  $\text{H}_2\text{SO}_4$  solution at 0.3 Pa as shown in Fig. 2b.<sup>48</sup> The spectra exhibited a broad peak, which was deconvoluted into two; one at the smaller energy was thought to originate from the electron-deficient oxygen species, and another one at the larger energy represents  $\text{O}^{2-}$





species. The study labelled the former one as  $O^{1-}$ , although it is still under debate whether the electron-deficient oxygen is electrophilic oxygen ( $O^{1-}$ ) or an oxyl radical ( $O^{1\cdot}$ ).<sup>31,52–57</sup> At more positive potentials, a hump appeared at the peak shoulder, indicating an increased contribution from the  $O^{1-}$  state. In correlating to the OER activity, they found that the OER performance linearly scaled with the fraction of not  $O^{2-}$  but  $O^{1-}$ , and thus concluded that the electrophilic  $O^{1-}$  species (6 in Fig. 1b) is the active species in catalysing the OER on iridium oxide.<sup>48</sup> We note that these species were observed at potentials above 1.6 V in Fig. 2c higher than the potentials employed in other cases, implying that this route may become likely apparent in higher overpotential regions. This view is compatible with a recent study claiming the charging up of iridium species under highly anodic conditions.<sup>59</sup>

The experimental evidence supporting the green-colored third scenario depicted in Fig. 1c was provided using Raman spectroscopy.<sup>58,69</sup> In 2020, Saeed *et al.* conducted *in situ* shell-isolated nanoparticle-enhanced Raman spectroscopy (SHINERS)<sup>58</sup> using electrochemically deposited iridium oxide in 0.1 M  $NaClO_4$  solution at pH 10 at varying potentials from OCP to 1.8 V vs.  $Ag/Ag^+$  (corresponding to 2.6 V vs. RHE at pH 10).<sup>58</sup> The measured spectra shown in Fig. 2c were divided into four distinct regions depending on the spectra features, (I) OCP to 0.5 V vs.  $Ag/Ag^+$  (corresponding to 1.3 V vs. RHE at pH 10), (II) 0.6 to 1.0 V vs.  $Ag/Ag^+$  (corresponding to 1.4 V to 1.8 V vs. RHE at pH 10), (III) 1.1 to 1.5 V vs.  $Ag/Ag^+$  (corresponding to 1.9 V to 2.3 V vs. RHE at pH 10), and (IV) 1.6 to 1.8 V vs.  $Ag/Ag^+$  (corresponding to 2.4 V to 2.6 V vs. RHE at pH 10), which are respectively red-, orange-, green-, and blue-colored spectra in the figure. The spectra exhibited multiple peaks, ascribed to the Ir–O–Ir twist, Ir–O–Ir stretch, or Ir=O stretch based on the previous Raman studies.<sup>69,70</sup> More specifically, the peaks labelled as  $\alpha$  (252  $cm^{-1}$ ) and  $\beta$  (357  $cm^{-1}$ ) were both assigned to the Ir–O–Ir twist at different iridium oxidation states; the former contains  $Ir^{4+}$ , and the latter contains  $Ir^{>4+}$ .<sup>69,70</sup> The five peaks of  $\gamma$  (504  $cm^{-1}$ ),  $\delta$  (608  $cm^{-1}$ ),  $\epsilon$  (719  $cm^{-1}$ ),  $\zeta$  (773  $cm^{-1}$ ), and  $\theta$  (672  $cm^{-1}$ ) were all ascribed to the Ir–O–Ir stretch;  $\delta$  was observed at the iridium oxidation state of +3, and  $\gamma$ ,  $\epsilon$ , and  $\zeta$  were apparent for  $Ir^{+4}$ , while  $\theta$  indicates the higher oxidation state of  $Ir^{>4+}$ .<sup>69,70</sup> The one at  $\eta$  (813  $cm^{-1}$ ) was thought to originate from the Ir=O stretch at  $Ir^{>4+}$ .<sup>69,70</sup> Regarding the assignment of this peak  $\eta$  in detail, previously the group of Schlögl investigated the influences of isotopes on the peak, and observed that this peak  $\eta$  was insensitive to the isotope H and D, indicative of the absence of H species in the vibration.<sup>69</sup> In addition, in an experiment using solution containing 50 : 50 ( $H_2O^{16} : H_2O^{18}$ ), only two peaks of nearly equal intensity were observed as  $\eta$ , which suggests that this vibration contains a single O atom, or the absence of O–O.<sup>69</sup> Furthermore, although the peak shift of single bonding Ir–O due to the substitution of  $O^{16}$  for  $O^{18}$  was calculated to be 45  $cm^{-1}$ , the observed shift was 59  $cm^{-1}$ . These observations led to a conclusion in the study that the peak  $\eta$  originated from Ir=O.<sup>69</sup>

In region (I) of Fig. 2c, the  $\gamma$ ,  $\delta$ , and  $\epsilon$  peaks were apparent, indicative of the presence of the  $Ir^{4+}$  state. When entering

region (II), an additional feature was observed in the spectra at  $\alpha$  and  $\zeta$ , which suggests that iridium species were still mostly composed of 4+ states. In region (III), the spectra exhibited contributions from  $\beta$  and  $\theta$ , which implies the formation of >4+ states. These  $\beta$  and  $\theta$  states persisted also in region (IV), where additionally the peak  $\eta$  appeared. Importantly, subsequent SHINERS spectroscopy combining the isotope labelling testing using  $D_2O$ -based solutions (1 M  $NaClO_4$  at pD 10) demonstrated almost no change in the peak position of  $\eta$  from 813  $cm^{-1}$  (in  $H_2O$ ) to 817  $cm^{-1}$  (in  $D_2O$ ). Thus, they concluded that all peaks do not contain H elements such as hydroxides, peroxides, or the oxyl radical species, which are considered as the precursor for O–O bond formation.<sup>69</sup> This study proposed that the OER catalytic cycle contains three intermediates, which all have an Ir–O bond, before rate-determining OOH formation, as shown in Fig. 1c.

The orange-colored last scenario of LOER in Fig. 1d was supported by studies using mass spectroscopy in conjunction with isotope labelling.<sup>65,71–74</sup> In 2007, Fierro *et al.* conducted differential electrochemical mass spectrometry (DEMS) measurements during a cyclic voltammogram (CV) in a potential window from *ca.* 0.05 V to *ca.* 1.6 V vs. SHE (corresponding to *ca.* 0.05 V to *ca.* 1.6 V vs. RHE at pH 0) in a 1 M  $HClO_4$  solution containing 10 wt%  $H_2^{18}O$  using thermally prepared  $Ir^{16}O_2/Ti$  catalyst, and their results are shown in Fig. 2d.<sup>65</sup> Their study tracked the identity of evolved  $O_2$  (either  $^{16}O_2$  or  $^{18}O^{16}O$ ) while recording the CVs. Fig. 2d shows the composition of  $^{18}O^{16}O$  and  $^{16}O_2$  at each CV scan, disclosing that the composition of  $^{16}O_2$  decreased with the cycle number while that of  $^{16}O^{18}O$  increased from 17% to 18% until reaching a steady state after the third cycle. To corroborate this observation, they performed an additional CV-DEMS experiment using  $Ir^{18}O_x$  in 1 M  $HClO_4$  containing *ca.* 10%  $H_2^{18}O$ . Their results showed that the evolved  $O_2$  gas was initially composed of  $^{18}O^{16}O$  at 0.9% in the first scan and was <0.5% after the fifth scan with a concomitant increase of  $^{16}O_2$  composition. These results allowed them to analyse the exchanging of lattice O of iridium oxide with O in the solution during the OER at steady state, and thus indicated the participation of lattice oxygen in the OER mechanism as depicted in the orange-colored cycle in Fig. 1d.

In addition to the spectroscopic study, morphological observations such as electron microscopy<sup>75–77</sup> also shed light on the catalysis of iridium oxide. For instance, in 2017, Willinger *et al.* compared the structural features between the more active electrocatalyst, which was hydrothermally prepared and denoted as  $IrO_x$ -FHI, and the less active one, which was commercially available and denoted as  $IrO_x$ -commercial.<sup>75</sup> By analyzing the observed images of high-angle annular dark field scanning transmission electron microscopy (HAADF STEM), they found a larger amount of the tunnel-like motifs of hollandite structure in  $IrO_x$ -FHI than that in  $IrO_x$ -commercial,<sup>75</sup> which was considered to be an active motif for the OER.<sup>78</sup> Their subsequent electron pair distribution function (ePDF) analysis suggested that  $IrO_x$ -FHI consisted of the hollandite structure, whereas  $IrO_x$ -commercial consisted of 60%





hollandite and 40% rutile unit cell.<sup>75</sup> These observations allowed for proposing that the hollandite structure is the key for the highly active OER catalyst, iridium oxide.<sup>75</sup> These studies demonstrate the significance of the morphological aspect on the OER activity.

Metal single-site catalysts have attracted increasing interest in recent years.<sup>79–82</sup> In 2020, an *in situ* XAS study observing the oxidation state of atomically dispersed iridium oxide on indium tin oxide (ITO) in 0.1 M HClO<sub>4</sub> during the OER was reported by Lebedev *et al.*<sup>79</sup> They measured *in situ* XANES at Ir L<sub>III</sub>-edge at 1.46 V *vs.* RHE and assigned an Ir–O distance of 1.83 ± 0.02 Å as an Ir<sup>+5</sup>=O intermediate, which was the predominant species under OER conditions.<sup>79</sup> Subsequently, the same authors further investigated the reaction mechanism by DFT calculation and suggested the formation of the Ir<sup>+6</sup> dioxo intermediate after the formation of the Ir<sup>+5</sup>=O intermediate.<sup>79</sup> This result of the formation of Ir<sup>+5</sup>=O is consistent with the observations on the heterogeneous solid states as discussed in this review.<sup>67</sup> This type of study would cultivate a better understanding of OER catalysis from a comprehensive perspective, and furthermore would eventually bridge the gap between heterogeneous and homogeneous catalysis.<sup>83–85</sup>

As described above, recent *in situ* and operando characterization studies captured the oxidation state of the active site and surface intermediate during the OER. The experimental evidence in Fig. 2 helps in reasoning the catalytic cycles, shaping the ones in Fig. 1. Notwithstanding the successful applications of these techniques to the electrocatalytic OER, there remains a discrepancy among studies about the oxidation state of iridium and oxygen as well as the surface intermediate, and thus the reaction mechanism. This discrepancy likely arises because of the non-unified catalyst and experimental conditions among reported studies, calling for further systematic investigations of the OER over iridium oxide with these sophisticated techniques.

### 2.3. Theoretical study corroborating the mechanism of H<sub>2</sub>O oxidation over iridium oxide

Theoretical studies help elucidate the reaction mechanism at molecular levels, and thus corroborate the experimental observations. Here in this subsection, we review selected theoretical studies on the OER over iridium oxide considering kinetics<sup>46,47,86</sup> that were omitted in the previously established DFT models.<sup>5,26,27</sup>

Ping *et al.* considered possible reaction pathways over the IrO<sub>2</sub>(110) surface, and computed reaction rates based on the microkinetic models at the constant potential of 1.36 V *vs.* normal hydrogen electrode (NHE) at pH 0 (corresponding to 1.36 V *vs.* RHE at pH 0).<sup>46</sup> Fig. 3a and b show the reaction intermediates and transition state (TS) structures as well as their free energy profile. In Fig. 3a, their considered model proceeds as follows: the initial state having Ir(OH)–O–Ir(OH) species (1'') is partially deprotonated to form Ir(OH)–O–Ir(O) (1'). Then, two scenarios were considered after the formation of this species (1'). In the first scenario, the species (1') experiences deprotonation to form Ir(O)–O–Ir(O) (1) that is

then attacked by H<sub>2</sub>O (TS1) and forms Ir(OH)–O–Ir(OOH) (2). Alternatively, Ir(OH)–O–Ir(O) (1') is attacked by H<sub>2</sub>O (TS1') and is transformed into Ir(OH<sub>2</sub>)–O–Ir(OOH) (2'), which is then deprotonated to form Ir(OH)–O–Ir(OOH) (2). The thus-formed species (2) is deprotonated to form Ir(OH)–O–Ir(OO) (3) and subsequently Ir(O)–O–Ir(OO) (4). The species (4) releases O<sub>2</sub> (TS2) and is protonated to Ir(O)–O–Ir(OH<sub>2</sub>) (5), which *via* internal PT transforms back to the initial state (1''). To assess the feasibility of the pathways, they computed the reaction barrier at TS1 and TS1', which amounted to 0.6 and 0.5 eV, respectively, as compiled in Fig. 3b. Notably, they thus proposed that a thermodynamically less favourable pathway (1' → TS1' → 2' → 2) could be kinetically more favourable.<sup>46</sup> This rationalization deepens the catalyst cycle in Fig. 1 with the proposal of rds being the formation of OOH (4 in Fig. 1a–c).

To investigate the influence of solvent on the conformation of the reaction intermediate during the OER, Gauthier *et al.* studied the interface between rutile IrO<sub>2</sub>(110) and H<sub>2</sub>O.<sup>47</sup> Fig. 3c shows the most stable structures found for the IrO<sub>2</sub>(110) surface with adsorbed \*OOH at the coordinatively unsaturated metal sites (CUS), \*O on the neighbouring CUS, and \*O at the bridge, together with varying numbers of water bilayers (BL).<sup>47</sup> In the absence of an explicit solvent (0 BL), the geometry of adsorbed intermediates was determined by interaction among the surface species; the hydrogen of the \*OOH species was the most stable when \*OOH was directed to the bridging O with hydrogen bonds as illustrated in Fig. 3c (0 BL). The presence of the explicit solvent (1–3 BL) changed the dominant interaction of the H element in the adsorbed \*OOH species from that with the surface to that with the BL. Fig. 3c (1–3 BL) illustrates the conformation in the presence of solvent: a more stable structure was found when a hydrogen bond was formed between the H of \*OOH and the explicit solvent. Notably, as long as BL was taken into account, the number of BLs had little influence on the calculation results. Thus, the bond structure of the active adsorbate was likely determined by the explicit solvent rather than the neighbouring adsorbate. Although by the subsequent thermodynamic calculation with the solvent, they observed that the reaction pathway was relatively unchanged by the inclusion of the explicit solvent, the solvent was found to stabilize the OOH adsorbate by about 0.3 eV. This study was consistent with the reaction intermediates shown in Fig. 1a–c, and provided new insights into the conformation of surface species by considering the effect of the solvent that shall impact the kinetics.

These advanced theoretical calculations demonstrated quantitatively the significance of the kinetic barrier and solvent on the OER electrocatalysis. The insights provided in those studies help in picturing the transition state in the ideal scenario, and would shed light on the rational design of active catalysts. In addition, theoretical studies may be able to assess the feasibility of catalytic cycles proposed by experiments (the four scenarios in Fig. 1) and may refine our view on electrocatalysis in the near future.





**Fig. 3** Theoretical studies for H<sub>2</sub>O oxidation over iridium oxide. (a) Considered reaction pathways for the OER over the IrO<sub>2</sub>(110) surface; red balls are O; blue balls are H; silver balls are Ir. (b) Computed reaction free energy diagram at the constant potential of 1.36 V vs. normalized hydrogen electrode (NHE) at pH 0 (corresponding to 1.36 V vs. RHE at pH 0) corresponding to the mechanism shown in the panel (a). Reprinted with permission from ref. 46. Copyright (2017) American Chemical Society. (c) The most stable structure for the IrO<sub>2</sub>(110) surface with adsorbed \*OOH at the coordinatively unsaturated metal sites (CUS), \*O on the neighboring CUS, and \*O at the bridge, together with varying numbers of water bilayers (BL); blue balls are Ir; red balls are O; white balls are H. Reprinted with permission from ref. 47. Copyright (2017) American Chemical Society.

### 3. OER mechanism *via* OH<sup>−</sup> oxidation over iridium oxide

There have been a limited number of studies on the OER over iridium oxide at alkaline pH levels, plausibly because of its lower OER stability than that under acidic conditions<sup>13,87</sup> and the existence of alternative catalysts such as NiFeO<sub>x</sub> active toward the OER.<sup>13,87</sup> Nonetheless, the investigation and elucidation of the OER over iridium oxide under alkaline pH conditions are of great significance because of two reasons. Firstly, the OER reactant varies with pH levels, H<sub>2</sub>O and OH<sup>−</sup> at acidic and alkaline pH levels, respectively, which corresponds to variation in the transition state and thus kinetics at different pH levels over iridium oxide. Understanding OER kinetics over iridium oxide under distinct conditions forms a generalized view into its catalysis, which would help rationally design OER catalysts. Secondly, increasing attention has recently been given to the OER at the near-neutral pH levels under buffered conditions,<sup>42,88–90</sup> where the OER can proceed also as the oxidation of OH<sup>−</sup>. Insights into the OER at alkaline pH thus certainly help advance the electrocatalysis under near-neutral pH conditions. We in this section describe firstly the mechanism of OH<sup>−</sup> oxidation, before detailing

spectroscopic evidence and theoretical calculations supporting the mechanism.

#### 3.1. Proposed mechanism of OH<sup>−</sup> oxidation over iridium oxide

There are two proposed OER mechanisms *via* OH<sup>−</sup> oxidation over iridium oxide that are essentially shared with those for the oxidation of H<sub>2</sub>O, depicted in Fig. 1: (1) a cycle involving surface adsorbate driven by the redox of the iridium centre (the red-colored cycle in Fig. 1a), and (2) a cycle involving the redox of the Ir=O state (the green-colored cycle in Fig. 1c). Notably, the difference between those cycles and the corresponding ones at acidic pH levels is that instead of the involvement of H<sub>2</sub>O, the cycles at alkaline pH consume the OH<sup>−</sup> reactant. In the next sections, we review reported studies supporting those scenarios.

#### 3.2. Spectroscopic evidence to support the claimed catalytic cycle for OH<sup>−</sup> oxidation over iridium oxide

In 2017, Ooka *et al.* reported the presence of Ir<sup>5+</sup> at the rds using ultraviolet-visible light (UV-Vis) spectroscopy.<sup>17,91</sup> They employed *in situ* evanescent wave (EW) spectroscopy at 1.2, 1.4, and 1.5 V vs. RHE in solutions at various pH levels from 2 to





**Fig. 4** Spectroscopic data for  $\text{OH}^-$  oxidation over iridium oxide. (a) *In situ* evanescent wave (EW) spectra in a mixed solution (NaOH, Britton–Robinson buffer and  $\text{Na}_2\text{SO}_4$ ) at pH 12 over anodically deposited iridium oxide with varying potential from 1.2 to 1.5 V vs. RHE. (b) Potentials for the OER onset (black), absorption at 450 nm (red), 410 nm, (green), and 580 nm (blue) on the RHE scale compiled as a function of pH, recorded in mixed solutions (NaOH, Britton–Robinson buffer and  $\text{Na}_2\text{SO}_4$ ) from pH 2 to 12 over anodically deposited iridium oxide. Reprinted with permission from ref. 17. Copyright (2017) American Chemical Society. (c) *In situ* surface-enhanced Raman spectroscopy (SERS) at varying potentials from OCP to 1.8 V vs. RHE in 0.1 M NaOH using iridium oxide on an Au substrate. Reproduced from ref. 69 with permission from the Royal Society of Chemistry.

12 over anodically deposited iridium oxide.<sup>17</sup> Fig. 4a shows the EW spectra measured at pH 12, in which two absorption peaks were apparent at approximately 410 and 580 nm at 1.2 V vs. RHE, assigned to  $\text{Ir}^{5+}$  and  $\text{Ir}^{4+}$ , respectively, based on the previous UV-vis study.<sup>38</sup> Interestingly, the former peak continuously shifted toward higher wavelengths at more positive potentials, ending at 456 nm at 1.5 V vs. RHE. Based on their previous time-dependent DFT (TDDFT) calculation result that theoretically elucidated spin states,<sup>91</sup> they considered that this shift from 410 to 456 nm with the potential was due to the change of the spin of two adjacent oxygen ligands from symmetric to asymmetric ones, which they proposed as a state of iridium centre at the rds. Their study also examined the absorption at different pH levels, and the potential at which these absorptions appeared on the RHE scale is shown as a function of pH in Fig. 4b. In the figure, the absorptions at 410, 450 nm (as an approximate representative of 456 nm), and 580 nm are denoted as Abs 410 nm, Abs 450 nm, and Abs 580 nm, respectively, and  $U_{\text{onset},j}$  stands for the onset potential of the OER. Interestingly, the potentials for Abs 580 nm and Abs 410 nm shifted with the pH value with a slope of 30 mV per pH on the RHE scale, implying that the redox reaction proceeds by transferring more than one proton per electron.<sup>92</sup> In contrast, the potential for Abs 450 nm and OER onset was mostly insensitive to the pH values as shown in Fig. 4b. This consistent insensitivity implies that the rds of the OER over iridium oxide was not the valence change of the iridium centre, but the activation of the oxygen ligand, such as the activation of  $\text{O}^{2-}$  to form  $\text{O}^-$ .<sup>17,48,50</sup> Overall, this study considered oxidation states consistent with the transition from  $\text{Ir}^{4+}-\text{O}^{2-}-\text{H}^+$  (2) to  $\text{Ir}^{5+}=\text{O}^{2-}$  (3) in the red-colored cycle of Fig. 1a, and proposed that the activation of the oxygen ligand would take place over  $\text{Ir}^{4+}=\text{O}^{1-}$  (6) as the rds.

In 2017, Pavlovic *et al.* reported a result of *in situ* Raman spectroscopy that supports the green-colored catalytic cycle in Fig. 1.<sup>69</sup> The group conducted *in situ* surface-enhanced Raman spectroscopy (SERS) at varying potentials from OCP to 1.8 V vs. RHE in 0.1 M NaOH using iridium oxide on an Au substrate, and the obtained spectra are shown in Fig. 4c.<sup>69</sup> At OCP, they observed peaks at 465, 527, 600, and 702  $\text{cm}^{-1}$ , which were attributed to the Ir–O–Ir stretching vibrations of  $\mu$ -oxo type oxygen linkages.<sup>70</sup> Upon increasing the potential to 0.4 V, new peaks  $\delta$ ,  $\epsilon$ , and  $\gamma$  appeared at 520, 595, and 476  $\text{cm}^{-1}$ , ascribable to the Ir–O stretch.<sup>70</sup> The spectra remained unchanged until reaching >1.4 V, where an additional peak was observed at 771  $\text{cm}^{-1}$  labelled as  $\eta$ . Their additional experiments employing the isotope D revealed that this peak  $\eta$  was insensitive to the isotopes H and D. In contrast, isotope labelling with  $^{18}\text{O}$  led to a shift of 59  $\text{cm}^{-1}$ , which differed from the expected shift of 45  $\text{cm}^{-1}$  for the surface species containing single bonding of Ir–O. Based on these observations, they hypothesized that the peak  $\eta$  originated from  $\text{Ir}=\text{O}$  vibration as opposed to any other species that contained O–O, or OH vibration,<sup>69</sup> which would serve as a precursor to the reactive oxyl species.<sup>93,94</sup> This  $\text{Ir}=\text{O}$  corresponds the species 3 in Fig. 1a and c, and species 6 in Fig. 1b. The species 3 in Fig. 1c would transform into the reactive oxyl species 7 in Fig. 1c.

### 3.3. Theoretical study corroborating the mechanism of $\text{OH}^-$ oxidation over iridium oxide

A rigorous analysis for the OER mechanism concerning both the kinetics and thermodynamics was provided in 2019 by the group of Exner.<sup>95</sup> Their approach relied on (1) DFT calculations to determine the free energy of the reaction intermediate and construct an *ab initio* Pourbaix diagram,<sup>96–100</sup> and (2) Tafel plots<sup>46,47,96,101,102</sup> to determine the electron transfer coefficient







Fig. 5 Theoretical studies for  $\text{OH}^-$  oxidation over iridium oxide. (a) The *ab initio* Pourbaix diagram for a single-crystalline  $\text{IrO}_2(110)$ . (b) The free energy diagram over  $\text{IrO}_2(110)$  in alkaline media (pH = 12.9) at an OER overpotential of 0.47 V. Reprinted with permission from ref. 95. Copyright (2019) American Chemical Society.

and free energies of the rate-determining transition state,<sup>95,103</sup> under the assumption of the Ir, Ir-OH, Ir=O and Ir-OOH intermediates. These considerations can be combined to construct a free energy diagram at a given overpotential, which allowed for pinning down the variation in the rds with applied overpotentials. Their *ab initio* Pourbaix diagram for  $\text{IrO}_2(110)$  is shown in Fig. 5a, and a representative free energy diagram over  $\text{IrO}_2(110)$  at an OER overpotential ( $\eta_{\text{OER}}$ ) of 0.47 V in alkaline solution at pH 12.9 is presented in Fig. 5b. Interestingly, their analysis disclosed that at  $\eta_{\text{OER}} > 0.3$  V, Ir-OOH is the thermodynamically most stable surface, and its conversion into Ir with releasing  $\text{O}_2$  is the rds at  $\eta_{\text{OER}} > 0.34$  V,<sup>95</sup> in contrast to the common understanding of the rds being the formation of O-O bonding.<sup>29,46,47,96,101,102</sup> This study corroborated the cycle in Fig. 1a-c, with a new insight that not the formation but the decomposition of OOH determines the rate.

#### 4. Dissolution of iridium oxide during the OER

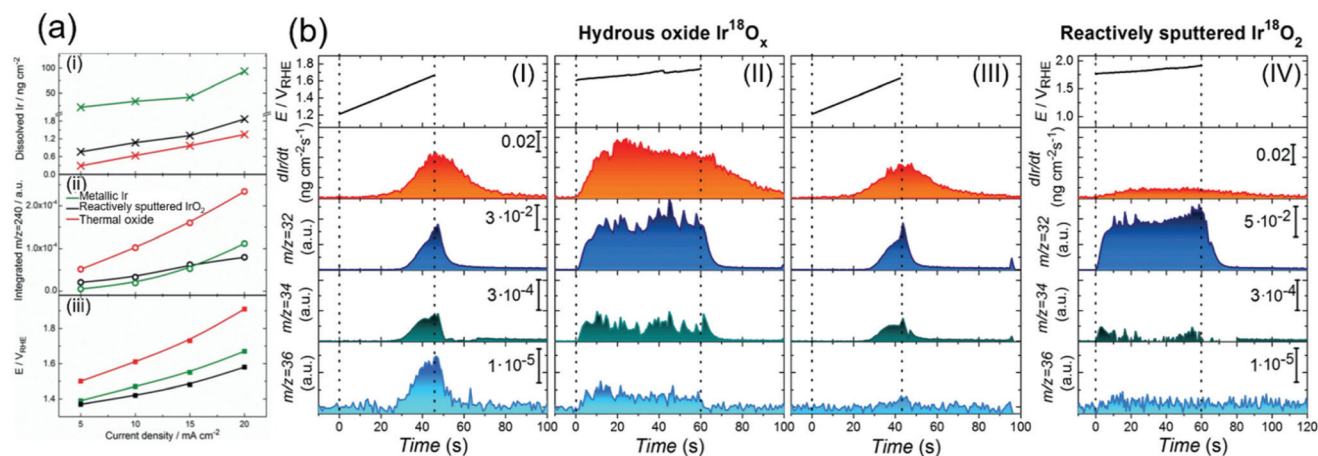
Although the degradation of OER activity over iridium oxide was observed in both acidic and alkaline solutions,<sup>13,101</sup> mechanistic studies on the dissolution of iridium have been reported only in acidic solutions.<sup>62,71,72,104</sup> Two types of dissolution pathways were proposed: (1) dissolving *via* redox of the iridium centre as  $\text{Ir}^{3+}$  (5) in Fig. 1a and b, (2) dissolving with releasing oxygen atom in iridium oxide as  $\text{Ir}^{6+}$  (15) or  $\text{Ir}^{x+}$  (16) during the LOER in Fig. 1d. The measurements were basically performed by electrochemical testing using a scanning flow cell (SFC) with mass spectroscopy such as inductively coupled plasma mass spectrometry (ICP-MS) or with online electrochemical mass spectrometry (OLEMS),<sup>62,71,72</sup> and key results are shown in Fig. 6.

In 2018, Kasian *et al.* reported the dissolution reaction of iridium oxide under acidic conditions using a SFC coupled with OLEMS or ICP-MS.<sup>62</sup> Their measurements were conducted in 0.1 M  $\text{HClO}_4$  solution at a constant current density

of 5, 10, 15, or 20  $\text{mA cm}^{-2}$  for 30 s, using three types of iridium oxide, namely electrochemically prepared oxide from metallic iridium (denoted as metallic Ir), reactively sputtered  $\text{IrO}_2$ , and thermally prepared iridium oxide. Fig. 6a shows, as a function of the applied current density, the amount of dissolved iridium species in **panel i**, integrated signals of mass spectra at mass-to-charge ratios ( $m/z$ ) of 240 as the representative of dissolved  $\text{IrO}_3$  in **panel ii**, and the corresponding potential on the scale of RHE (**panel iii**). The reactively sputtered  $\text{IrO}_2$  exhibited a lower dissolution rate than the metallic iridium similar to thermal oxide according to **panel i**, and the highest activity as in **panel iii**. Its superior activity with a slower dissolution rate was consistent with the previous study,<sup>105</sup> and the lower dissolution rate of the thermal one likely originated from its thermodynamic stability.<sup>106</sup> Notably, metallic Ir exhibited the largest amount of dissolved iridium as shown in **panel i**, and its dissolution rate as  $\text{IrO}_3$  exponentially increased with the current density unlike the other iridium oxides (**panel ii**). They proposed that this increase was due to two dissolution-pathways prevailing at distinct potentials: below 1.6 V *vs.* RHE, the iridium species likely formed  $\text{HIr}^{3+}\text{O}_2$  and dissolved as  $\text{Ir}^{3+}$  *via* the OER catalytic cycles involving  $\text{Ir}^{3+/4+/5+}$  as previously reported,<sup>72</sup> and above 1.6 V *vs.* RHE, the  $\text{IrO}_3$  species react with water and are dissolved in the form of  $\text{Ir}^{6+}\text{O}_4^{2-}$ , which would not be able to be re-deposited on the catalyst surface, *via* the  $\text{Ir}^{4+/5+/6+}$  pathway.<sup>63</sup> This reasoning is reflected in Fig. 1: (1) dissolution of  $\text{Ir}^{3+}$  (5 in Fig. 1a and b) for the pathway of  $1 \rightarrow 2 \rightarrow 3$  (or  $6 \rightarrow 4 \rightarrow 5$ ), and (2) dissolution of  $\text{Ir}^{6+}$  (15 in Fig. 1d) for the pathway through  $9 \rightarrow 10 \rightarrow 11 \rightarrow 15$  or  $9 \rightarrow 13 \rightarrow 15$ .

Recently, the dissolution of iridium oxide is thought to be related to the LOER mechanism<sup>71,73</sup> based on previous observations over various oxide surfaces.<sup>64,107,108</sup> Kasian *et al.* investigated a degradation mechanism using a SFC coupled with OLEMS and ICP-MS in 0.1 M  $\text{HCl}^{16}\text{O}_4$  solution.<sup>71</sup> They used two types of iridium oxide: electrochemically formed hydrous  $\text{Ir}^{18}\text{O}_2$  catalysts, and reactively sputtered  $\text{Ir}^{18}\text{O}_2$ . The detected signals of evolved  $\text{O}_2$  gas were composed of three distinct





**Fig. 6** Experimental data on the dissolution of iridium oxides. (a) Results of a scanning flow cell (SFC) coupled with online electrochemical mass spectrometry (OLEMS) or inductively coupled plasma mass spectrometry (ICP-MS) in 0.1 M HClO<sub>4</sub> solution at constant current densities of 5, 10, 15, or 20 mA cm<sup>-2</sup> for 30 s using three types of iridium oxide: electrochemically prepared oxide from metallic iridium (denoted as metallic Ir, green), reactively sputtered IrO<sub>2</sub> (black), and thermally prepared iridium oxide (red). (i) The amount of dissolved Ir, (ii) the amount of dissolved IrO<sub>3</sub> (*m/z* = 240), and (iii) potential at the end of polarization on the current density. Reprinted with permission from ref. 62. Copyright (2018) Wiley-VCH. (b) Results for the dissolution amount of iridium species and evolved O<sub>2</sub> gas (<sup>16</sup>O<sup>16</sup>O, <sup>16</sup>O<sup>18</sup>O, and <sup>18</sup>O<sup>18</sup>O at *m/z* = 32, 34, and 36, respectively) using a SFC coupled with OLEMS and ICP-MS in 0.1 M HCl<sup>16</sup>O<sub>4</sub> solutions: (I) using hydrous Ir<sup>18</sup>O<sub>x</sub> with sweeping the potential from 1.2 V vs. RHE to the potential where the current density reached a value of 20 mA cm<sup>-2</sup>; (II) using hydrous Ir<sup>18</sup>O<sub>x</sub> at potentials where the current density reached 15 mA cm<sup>-2</sup> for 60 s after the experiment in panel (I); (III) using hydrous Ir<sup>18</sup>O<sub>x</sub> with sweeping the potential from 1.2 V vs. RHE to the potential where the current density reached a value of 20 mA cm<sup>-2</sup> after the experiment in panel (II); and (IV) using reactively sputtered Ir<sup>18</sup>O<sub>2</sub> at potentials where the current density reached 15 mA cm<sup>-2</sup> for 60 s. Reproduced from ref. 71 with permission from the Royal Society of Chemistry.

species; <sup>16</sup>O<sup>16</sup>O, <sup>16</sup>O<sup>18</sup>O, and <sup>18</sup>O<sup>18</sup>O at *m/z* = 32, 34, and 36, respectively.<sup>71</sup> Fig. 6b shows the detected concentration of dissolved iridium species and evolved oxygen gas. Their experimental protocol for hydrous Ir<sup>18</sup>O<sub>2</sub> was (1) sweeping the potential from 1.2 V vs. RHE to the potential where the current density reached a value of 20 mA cm<sup>-2</sup> in **panel I**, then (2) applying potential where the current density reached 15 mA cm<sup>-2</sup> for 60 s in **panel II**, and finally (3) sweeping the potential again from 1.2 V vs. RHE to the potential reaching 20 mA cm<sup>-2</sup> in **panel III**. Critically, the dissolution of iridium species was observed in every scan. The concentration of dissolved iridium species increased with increasing applied potential in protocol (1) and (3), and the concentration almost remained constant throughout protocol (2). Furthermore, comparison of step (1) shown in **panel I** with step (3) shown in **panel III** revealed that the signals of both <sup>16</sup>O<sup>18</sup>O and <sup>18</sup>O<sup>18</sup>O in **pane III** were smaller than those in **panel I**, implying that oxygen atoms in IrO<sub>2</sub> participated in the OER. In contrast, their experiment using the sputtered IrO<sub>2</sub> provided a distinct picture. Their experimental protocol was to apply a potential reaching 15 mA cm<sup>-2</sup> for 60 s, the same as step (2) of the protocol for the hydrous ones, and the obtained result is provided in **panel IV**. Interestingly, the figure disclosed a lesser signal of iridium dissolution than the hydrous one, and negligible signals of <sup>16</sup>O<sup>18</sup>O and <sup>18</sup>O<sup>18</sup>O.<sup>71</sup>

In their study, Kasian *et al.* detailed the dissolution mechanism and lattice oxygen participation during the OER.<sup>71</sup> Analysing the data in Fig. 6b disclosed that the ratio of the iridium dissolution rate and the <sup>18</sup>O<sup>18</sup>O formation rate was the

same for both hydrous and sputtered IrO<sub>2</sub>, suggesting a link between the participation of lattice oxygen and the dissolution of iridium. They subsequently studied the surface species and morphology of both samples using XPS and atom probe tomography (ATP) in conjunction with reported near ambient pressure XPS (NAP-XPS).<sup>24,49,109</sup> It was observed that hydrous iridium oxide possessed nanopores and -Ir<sup>3+</sup>OOH- species, while the reactively sputtered one did not. They reasoned that the opened spaces provided by the nanopores would allow for frequent attacks by H<sub>2</sub>O molecules, facilitating the dissolution of iridium species as Ir<sup>x+</sup> (16) with the leaching of lattice oxygen atoms. Additionally, the Ir<sup>3+</sup>-OOH species would form HIr<sup>3+</sup>O<sub>2</sub> that dissolves into the solutions as Ir<sup>3+</sup>.<sup>62,72</sup> Taken together, the dissolution was faster for the hydrous one due to its morphological and chemical state, which accompanies the removal of lattice oxygen detected as the formation of <sup>16</sup>O<sup>18</sup>O and <sup>18</sup>O<sub>2</sub>, while such an event was less pronounced for the sputtered one due to the distinct surface state. These findings are reflected in the catalytic cycles described in the orange-colored cycle in Fig. 1d.

## 5. Summary and future outlook

This article reviewed the recent reports on the *in situ* or operando spectroscopic studies as well as advanced computational work for the OER over one of the most active catalysts, iridium oxide. By collectively summarizing the literature, we illustrated the OER catalytic cycles on iridium oxide for the oxidation of



both H<sub>2</sub>O and OH<sup>-</sup> occurring mainly at acidic and alkaline pH levels, respectively. *In situ* or operando spectroscopic studies such as XPS and XANES captured the oxidation state and chemical environment of the catalysts at work, while the studies employing IR and Raman have elucidated the surface binding species in operando. The advanced calculations addressed the kinetic aspects of the OER catalysis, refining our view at the molecular level. Notably, while shaping the scheme of the catalytic cycles, we noticed that there exists some discrepancy regarding the active site, oxidation states, surface intermediates and rds. The discrepancy likely arose from distinct experimental conditions in the literature, *e.g.*, the synthesis protocol for the iridium sample, electrolyte, applied potential, and so on. Therefore, it would be critical in future studies to comprehensively and systematically perform spectroscopic studies that can provide us with a more solid understanding of OER catalysis. In addition, a recent study claimed that the amount of accumulated oxidative charge of electrocatalysts dictates the activation energy of the reaction, which has not been taken into account in the previous studies on the mechanism.<sup>58</sup> This novel concept may link the studies, which can further refine our view on OER catalysis.

The established view on the reaction mechanism helps develop design principles of active and stable OER catalysts. The spectroscopic observation of the Ir<sup>5+</sup> state *in situ* or operando<sup>49,66,67</sup> implies that the rds occurs on this Ir<sup>5+</sup> site, plausibly Ir=O, which is thought to convert into Ir-OOH in one of the cycles.<sup>29,46,47,69,110</sup> This rationale suggests that stabilizing the Ir-OOH state by introducing a second component as a proton acceptor would accelerate this rds.<sup>111</sup> Such an approach was theoretically supported with recent DFT calculations by Busch<sup>112</sup> and has recently been demonstrated experimentally in the case of Mn-based catalysts.<sup>113</sup>

Another rational approach for improving the OER performance based on the rational understanding is the stabilization of the high-valence state of iridium by introducing a second component. It was reviewed in this article that iridium oxide dissolves into the solutions with oxidation states of 3+ or 6+.<sup>62,72</sup> This finding suggests that stabilizing these oxidation states of iridium by introducing a second component would prevent the dissolution, which can prolong the lifetime of catalysts. In fact, improved stability was reported for IrO<sub>x</sub>/SrIrO<sub>3</sub> in 2016,<sup>114</sup> and IrO<sub>x</sub>-TiO<sub>2</sub> in 2017.<sup>115</sup> Thus, the rational design of catalysts based on the understanding of the reaction mechanisms would be an effective approach to improve the performance of iridium-based OER catalysts. Moreover, in such discussions of introducing additional components, the OER activity also improved by leaching the second or third component from the complex such as Sr from the perovskite of SrIrO<sub>3</sub>,<sup>114</sup> Y from the pyrochlore of Y<sub>2</sub>Ir<sub>2</sub>O<sub>7</sub>,<sup>116</sup> Ba and Ln from the double perovskite of Ba<sub>2</sub>PrIrO<sub>6</sub>,<sup>117</sup> Ni and Co from Ir-based nanowire,<sup>118</sup> and Ni from the IrNiO<sub>x</sub> core-shell structure.<sup>119</sup> The thus-derived surface would contribute to the improvement of the OER activity by constructing highly active iridium centers such as iridium hydroxide-dominant surfaces.<sup>51</sup> This review provides the community with a generalized view on the

state-of-the-art understanding of the OER mechanism over iridium and will stimulate further understanding and development of OER catalysts in the future.

## Conflicts of interest

There are no conflicts to declare.

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