The electronic structure of iridium oxide electrodes active in water splitting†

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Iridium oxide based electrodes are among the most promising candidates for electrocatalyzing the oxygen evolution reaction, making it imperative to understand their chemical/electronic structure. However, the complexity of iridium oxide’s electronic structure makes it particularly difficult to experimentally determine the chemical state of the active surface species. To achieve an accurate understanding of the electronic structure of iridium oxide surfaces, we have combined synchrotron-based X-ray photoemission and absorption spectroscopies with ab initio calculations. Our investigation reveals a pre-edge feature in the O K-edge of highly catalytically active X-ray amorphous iridium oxides that we have identified as O 2p hole states forming in conjunction with IrIII. These electronic defects in the near-surface region of the anionic and cationic framework are likely critical for the enhanced activity of amorphous iridium oxides relative to their crystalline counterparts.

Water splitting presents an attractive solution to store excess energy from intermittent renewable sources in the form of hydrogen.1 Iridium oxide is the only known anode electrocatalyst that is both active and stable in the sluggish oxygen evolution reaction (OER) in acidic media,2 which has prompted significant research efforts focused on this conducting oxide. Recently, these efforts have revealed that amorphous forms of iridium oxide have higher activities than their rutile-type crystalline counterparts,3,4 which we refer to simply as rutile IrO2. An explanation for this increased activity has, however, proven elusive because of the challenges associated with understanding the combined influence of band structure, electron correlation, and spin–orbit coupling on the electronic structure of iridium oxide.5 For example, even the Ir 4f spectrum of rutile IrO2 has a line shape that cannot be fit using the standard theoretically derived function, making the origin of this unusual shape a matter of debate.6–8 Thus, while X-ray photoemission and absorption spectroscopies (XPS & XAS) have been used to search for OER relevant surface species in iridium and its oxides, both in situ9–13 and ex situ14,15 unambiguous speciation has remained challenging. Herein, we combine XPS measurements and theory to uncover the origin of the peculiar Ir 4f line shape of rutile IrO2. We use this finding along with the Near-Edge X-ray Absorption Fine Structure (NEXAFS) of the O K-edge to identify the species present in a catalytically more active, amorphous, hydrated IrOx.

All experiments were performed on two commercially available iridium oxide powders. Prior to photoemission experiments, the powders were thoroughly characterized. X-ray diffraction (XRD) confirmed the rutile structure and phase purity of the IrO2 reference material whereas the IrOx powder was found to be X-ray amorphous with a minor metallic contribution (Fig. S1, ESI†). Energy-dispersive X-ray spectroscopy (EDX) yielded the expected stoichiometry of the rutile sample (1:2/Ir:O). In contrast, the amorphous powder had a significantly higher oxygen concentration (Table S1, ESI†) giving rise to a total water loss of ~6 wt% in thermogravimetric (TG) analysis (Fig. S2, ESI†). From temperature-programmed reduction (TPR) (Fig. S3, ESI†) of rutile IrO2, a formal Ir oxidation state of 4.1 ± 0.1 is deduced, demonstrating that iridium and oxygen in rutile IrO2 can be viewed as IrIV and OII−, despite the distorted octahedral environment around each Ir atom and the covalent character of the Ir–O interaction.16 In contrast, the formal oxidation state of oxygen-containing iridium species in the amorphous IrOx is only found to be 3.6 ± 0.1, suggesting that multiple iridium species are present in the amorphous IrOx, some of which are in an oxidation state of less than IV. This result is in line with titration experiments done on hydrated IrOx colloid catalysts, for which an average oxidation state of 3.2 was found.17

† Electronic supplementary information (ESI) available: Further details on the experiments, measurement results and the DFT calculations are presented. See DOI: 10.1039/c5cp06997a
The presence of mixed Ir-valence states has been argued to be essential for the OER on amorphous IrO$_x^{12,18}$. In support of this view, activity tests of the two powders confirmed the expected higher OER activity of the amorphous IrO$_x$ with respect to rutile IrO$_2$ (Fig. S4, ESI†). The enhanced activity of amorphous iridium oxides is likely due to surface species that have an intrinsically higher OER activity than those of the rutile IrO$_2$. To uncover the nature of these species, we turn to XPS and NEXAFS.

Fig. 1 shows a superposition of the measured Ir 4f spectra of the rutile and the amorphous powders normalized by area. Photoemission of an Ir 4f$_{7/2}$ (4f$_{5/2}$) electron in IrO$_2$ gives rise to the intensity at 61.8 eV (64.8 eV) with the anomalous, highly asymmetric line shape typical of rutile IrO$_2$. Though the peak positions in the Ir 4f spectrum of amorphous IrO$_x$ are similar to that of rutile IrO$_2$, the line shape is not, as highlighted by the difference spectrum. The amorphous IrO$_x$ can be seen to have less intensity where the main peaks of rutile IrO$_2$ are located and more intensity around 62.4 eV and 65.4 eV. This additional intensity is not related to the metallic component seen in XRD, which would appear at 60.9 eV, suggesting that the metal detected with XRD is covered by a thick oxide layer and does not contribute to the OER activity. The lack of a metallic contribution in XPS makes it tempting to assign the features at 62.4 eV and 65.4 eV to the low oxidation state of iridium seen in TPR. However, the ambiguity in the line shape of IrO$_2$ makes such an assignment questionable without further evidence.

It is well known from many-body theory that, as a conductor, IrO$_2$ has asymmetric core level spectra. However, the peculiar asymmetry of the Ir 4f spectrum cannot be fit with only one component when employing the standard Doniach–Sunjic (DS) line shape. Although a variety of hypotheses have been put forward to explain this apparent failure, it should be noted that the DS function is formally exact only at the edge singularity. The widespread applicability of the DS line shape over a wider energy range is simply a consequence of the weakly structured electron–hole pair excitation spectra common in many metals. Clearly such weak structure is not required, nor is it always present. While the true line shape can be computed by perturbation theory, such an approach requires detailed knowledge of the electron–hole pair excitation spectra, hence the atomic structure, and offers little chemical insight. As such, we have chosen to employ a one-electron picture wherein the structured electron–hole pair excitation spectra give rise to “shake-up” satellites, simultaneous core ionization and monopole excitation of a valence electron. While approximate, this approach can be used to understand the experimental spectra and develop robust fit models for complex systems where we lack detailed knowledge of the structure.

To investigate the possibility of shake-up satellites in the Ir 4f spectrum of rutile IrO$_2$, we turned to density functional theory (DFT) calculations of the states involved in the shake-up process. The d projected density of states (PDOS(d)) of an Ir atom with an Ir 4f core hole in rutile IrO$_2$ has a strong narrow feature at 1 eV above the Fermi energy ($E_F$) (see inset in Fig. 2 and Fig. S9, ESI†) to which valence electrons can be excited. Because the ground state is metallic and, assuming constant excitation matrix elements, the probability of valence electron excitation scales with the inverse square of the excitation energy ($1/\Delta E^2$), the most intense shake-up satellites originate from occupied states near $E_F$. Weaker satellites will originate from states with energies below $E_F$, the most prominent of these will be due to shake-up from localized non-bonding Ir d states at 2 eV binding energy.

![Fig. 1](https://example.com/fig1.png)

**Fig. 1** Ir 4f spectra and difference spectrum of rutile IrO$_2$ and amorphous IrO$_x$ measured in UHV at a kinetic energy of the photoelectrons of 900 eV.

![Fig. 2](https://example.com/fig2.png)

**Fig. 2** Theory-based fit models for rutile IrO$_2$ (below) and amorphous IrO$_x$ (above). The inset shows the unoccupied PDOS(d) of rutile IrO$_2$ in presence of an Ir 4f core hole. The sharp feature ~1 eV above $E_F$ causes the shake-up satellite to appear at ~1 eV above the main line.
4f5/2 line was included to account for shake-up from the localized fit, a small component at 3.1 eV higher binding energy than the Ir satellite at 1.0 eV higher binding energy. To slightly improve the function with the peak maximum at 61.8 eV and a Gaussian structure can be fit as shown in Fig. 2. The fit consists of a DS associated with the formation of O 2p holes. 24,25

The presence of principally IrIV and OII– in rutile IrO2 is further corroborated by NEXAFS measurements. The strong resonances at 530 eV and 533 eV are in excellent agreement with the computed O K-edge spectrum of rutile IrO2, where all oxygen is formally OIV (Fig. 3). In the O K-edge spectrum of the amorphous IrOx, however, a new resonance appears at ~529 eV, which is also reflected in the O 1s spectrum (Fig. S5, ESI†). This type of pre-edge feature is commonly observed in other transition metal oxides with strong covalent interactions.24,25 While not an exact comparison, such a pre-edge is perhaps most well-known to emerge when hole doping superconducting cuprates, where it is associated with the formation of O 2p holes.24,25

We can computationally test if such formally OI– species also lead to a resonance at ~529 eV in iridium oxide by introducing iridium vacancies to create hole states in IrO2 (Fig. S6 and S8, ESI†). Doing so, we find that the holes localize on oxygen, which, unlike hole localization on iridium (see ESI†), leads to results consistent with experiment. The OI– species give rise to a new resonance at 529 eV in the calculated O K-edge, though they lack the resonance at 530 eV seen in the OII– spectrum (Fig. S7, ESI†).

Though this model structure cannot describe all aspects of the amorphous IrOx by assuming that the electronic structure of the computed OI– in IrO2 is similar to that of the analogous coordination defect in the amorphous IrOx, we are able to recover the experimentally observed O K-edge spectrum. Under this assumption, we can take the model computed spectra as representatives of OI– and OII–, allowing us to conclude that amorphous IrOx contains both oxygen species within the probing depth of the measurement. This suggestion is supported by the fact that a linear combination of the two types of computed spectra, using 2:3 for the ratio of OI– to OII–, yields good agreement with experiment (Fig. 3). The physisorbed and chemisorbed water content of the IrOx sample, which was ignored in our computed spectra, leads to a smoothing of the measured O K-edge spectrum at excitation energies between 537 eV and 545 eV,26 along with additional intensity at higher binding energy in the O 1s spectrum (see Fig. S5, ESI†).

The electronic defects in the oxygen seen in NEXAFS also lead to the formation of electronic defects in the iridium framework. Six OI– form with each Ir vacancy, leaving two free electrons that reduce neighboring IrIV to IrIII to maintain local charge neutrality (Fig. 4). Our calculations reveal that this formally IrIII species has an Ir 4f7/2 binding energy of ~62.2 eV, which is in line with binding energy values measured for IrIII in IrClx.27 As before, assuming that the iridium vacancy modeled in our calculations is representative of coordination defects in the amorphous IrOx, we can use our computed result to conclude that IrIII can account for the increased intensity at higher binding energy in the Ir 4f spectrum (see Fig. 1). As with the IrIV species, analysis of the PDOS suggests that a shake-up satellite will appear at ~1.5 eV above the Ir 4f main line of the IrIII species (see Fig. S11, ESI†).

Based on these theoretical findings, we obtain the fit model shown in Fig. 2 for the amorphous IrOx. The Ir 4f spectrum can be deconvoluted into a contribution of formally IrIV and IrIII

**Fig. 3** Measured O K-edge spectra of rutile IrO2 and amorphous IrOx and calculated O K-edges of OI– and a linear combination of 60% OII– and 40% OI– species.

**Fig. 4** IrO6 octahedron in the near-surface region of the IrOx framework (grey lines). Initially, Ir and O are in the formal oxidation states IV and II–, respectively. Upon the creation of an Ir vacancy, surrounding oxygen atoms are oxidized to the formal oxidation state I–. The two additional electrons are distributed among neighboring IrIV reducing them to IrIII.
species along with their respective satellites. An increased full width at half maximum of IrIV is used to recover the fact that the local environment around the IrV atoms in the amorphous structure is expected to be less regular than in the crystalline rutile structure. All fit parameters can be found in Table S3 (ESI).

In conclusion, we report on three major findings. First, we have presented theoretical calculations that offer a convincing explanation for the unusual Ir 4f line shape of phase-pure rutile IrO2. From these results, we developed a fit model, in which a Doniach–Sunjic line shape is combined with Gaussian satellites to account for electron shake-up from ET into unoccupied states 1 eV above ET. In the second part of the manuscript we have shown that caution is required for the speciation of iridium oxides. For though we observe an increase in intensity at higher binding energy in the Ir 4f spectrum of the catalytically more active amorphous iridium oxide powder, we see little indication for the presence of Ir in oxidation state V. Both TPR and our calculations suggest that while two Ir components are present in the amorphous iridium oxide, the lower binding energy feature is identified as formally IrIV, as in rutile IrO2, whereas the higher binding energy species is attributed to formally IrIII, which is associated with the formation of O 2p hole states. Finally, these O 2p hole states give rise to a formally OI species. As a chemically electrophilic oxygen, OI is expected to be susceptible to nucleophilic attack during OER, consistent with Fierro et al.’s findings that the OER is likely based on the consumption and regeneration of lattice oxygen from the near surface region of the catalyst. Our finding that an electrophilic oxygen can be identified by a resonance at ~529 eV in the O K-edge of the amorphous IrOx, in addition to the OII resonance at 530 eV, then allows the study of, not only the active metal centers, but the simultaneous presence of electronic defects in the anionic and cationic framework. These mixed valences of iridium and oxygen likely play a key role in OER catalysis, a finding that might not be restricted to iridium oxides but pertain to OER-active catalysts in general.

To aid in our understanding of the intricate roles these species play in reactions, we are investigating their formation and dynamics as a function of applied potential using in situ XPS and NEXAFS.

**Computational methods**

All Density Functional Theory (DFT) calculations were performed with the Quantum ESPRESSO package29 using the Perdew, Burke, and Ernzerhof (PBE) exchange and correlation potential30 with spin polarization and scalar relativistic corrections. We employed ultrasoft pseudopotentials with a 50 Ry (500 Ry) kinetic energy (charge density) cutoff and a k-point mesh equivalent to (16 × 16 × 16) for the crystallographic unit cell along with Marzari–Vanderbilt cold smearing31 with a smearing parameter of 0.005 Ry. Core level binding energies (BE) were computed using the ASCF method to accurately recover initial and final state effects.32 Oxygen K-edge spectra were computed from Fermi’s golden rule using the XSpectra package.33 Further details are given in the ESI.†

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**References**