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11634Elucidating the alkaline oxygen evolution reaction
mechanism on platinum†M. Favaro,^{abc} C. Valero-Vidal,^{ad} J. Eichhorn,^c F. M. Toma,^c P. N. Ross,^e J. Yano,^{bf}
Z. Liu^{gh} and E. J. Crumlin^{*ad}

Understanding the interplay between surface chemistry, electronic structure, and reaction mechanism of the catalyst at the electrified solid/liquid interface will enable the design of more efficient materials systems for sustainable energy production. The substantial progress in *operando* characterization, particularly using synchrotron based X-ray spectroscopies, provides the unprecedented opportunity to uncover surface chemical and structural transformations under various (electro)chemical reaction environments. In this work, we study a polycrystalline platinum surface under oxygen evolution conditions in an alkaline electrolyte by means of ambient pressure X-ray photoelectron spectroscopy performed at the electrified solid/liquid interface. We elucidate previously inaccessible aspects of the surface chemistry and structure as a function of the applied potential, allowing us to propose a reaction mechanism for oxygen evolution on a platinum electrode in alkaline solutions.

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Introduction

With the world's energy utilization increasing, new solutions are required to satisfy the energy demand while minimizing the environmental impact. To meet this challenge, new technologies must be improved to achieve sustainable energy conversion and consumption.^{1,2} Fuel cells and electrolyzers³ constitute a promising candidate towards the clean conversion of chemical energy into electricity and electricity into chemical energy, respectively. However, their current use is still partially limited by several factors, such as the limited efficiency and stability of the low cost alternatives to platinum (Pt) for the electrocatalysis of oxygen reduction and evolution reactions (ORR and OER,

respectively).^{4,5} In order to enhance the catalytic performance of next generation fuel cells and electrolyzers, it is necessary to obtain a comprehensive understanding of the chemistry and electronic structure of the catalyst and the reaction intermediates/products under realistic operating conditions.^{6,7} In particular, Pt has been used as a model system to provide surface chemical and structural transformation insights under various (electro)chemical reaction environments.^{8–23} Ultimately, the scope of these studies is to expedite the development of novel and cost-effective catalysts with tailored physical/chemical properties, capable of achieving, if not surpassing, the Pt electrocatalytic performance.^{8,24–27}

Previous studies have been conducted to identify the role of Pt oxides in the oxygen evolution reaction pathway (also known as oxygen discharge).^{8,24–26} Krasilshchikov^{24,26,28} demonstrated that at high overpotentials the corresponding current density (Tafel slope 120 mV dec^{−1}) depends exponentially on the thickness of the Pt surface oxide. Other studies report much higher Tafel slopes (up to 160 mV dec^{−1})^{13,24,26} in both acidic and alkaline environments. Such an increase is often attributed to an increase of the surface oxide layer thickness and Pt dissolution.^{13,15–26,29} In alkaline media, the reversible binding of a hydroxide ion (OH[−]) coupled to a one electron oxidation is thought to precede the removal of one proton and one electron to form surface oxide species.³⁰ This hypothesis has been mostly supported so far by surface structure and chemical characterizations far from *operando* conditions,^{6,31,32} thereby limiting an unambiguous investigation of the role of Pt oxides in the oxygen evolution catalytic cycle. Moving towards realistic operating conditions, Arrigo *et al.* recently studied Pt nanoparticles deposited on an acidic-polymer membrane using ambient pressure X-ray photoelectron spectroscopy (APXPS) in a humidified (solid/gas interface) environment.²⁷ The authors found that the

^aAdvanced Light Source, Lawrence Berkeley National Laboratory, One Cyclotron Rd, Berkeley, CA 94720, USA. E-mail: ejcrumlin@lbl.gov

^bJoint Center for Artificial Photosynthesis, Lawrence Berkeley National Laboratory, One Cyclotron Rd, Berkeley, CA 94720, USA

^cChemical Sciences Division, Lawrence Berkeley National Laboratory, One Cyclotron Rd, Berkeley, CA 94720, USA

^dJoint Center for Energy Storage Research, Lawrence Berkeley National Laboratory, One Cyclotron Rd, Berkeley, CA 94720, USA

^eMaterials Sciences Division, Lawrence Berkeley National Laboratory, One Cyclotron Rd, Berkeley, CA 94720, USA

^fMolecular Biophysics and Integrated Bioimaging Division, Lawrence Berkeley National Laboratory, One Cyclotron Rd, Berkeley, CA 94720, USA

^gState Key Laboratory of Functional Materials for Informatics, Shanghai Institute of Microsystem and Information Technology, Chinese Academy of Sciences, Shanghai 200050, People's Republic of China

^hDivision of Condensed Matter Physics and Photon Science, School of Physical Science and Technology, ShanghaiTech University, Shanghai 200031, China

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To achieve a molecular-level comprehension of the OER mechanism, we studied a polycrystalline Pt electrode surface under anodic conditions in an alkaline electrolyte, by means of *operando* APXPS performed with “tender” X-rays ($h\nu = 4$ keV) directly at the electrified solid/liquid interface.^{35,36} We report the speciation of porous Pt surface oxides as a function of applied potential, thereby elucidating the differences between thermodynamic predictions (Pourbaix diagrams)³⁷ and the actual surface composition under polarization. Moreover, we study the Pt surface chemistry as a function of time during oxygen evolution, thereby observing the evolution of the different Pt oxidation states (as well as their spatial localization in the near-surface region) under non-equilibrium conditions (initial transient) and upon reaching steady state conditions. Based on these findings, we propose an OER mechanism in terms of a catalytic (Tolman) cycle³⁸ for the various oxidized Pt species. We show that the rate-determining step is constituted by activated Pt sites selectively binding OH^- . In addition, by coupling the experimental results with numerical simulations of the photoelectron intensity, we reveal that the active sites during OER are located in the sub-surface region, whereas Pt oxides such as $\text{Pt}^{\text{(II)}}\text{O}$ and $\text{Pt}^{\text{(IV)}}\text{O}_2$ are present in the high-surface outer layer and are not actively contributing to the oxygen evolution.

Probing the solid/liquid interface

To approximate the current density of the thin electrolyte layer under *operando* conditions formed by the “dip & pull” procedure, we conducted a separate electrochemical experiment that preserved the same “dip & pull” geometries, we masked the bottom of the electrode that was immersed in the bulk electrolyte and compared to an unmasked electrode (see schematization reported in ESI Fig. 1†). The thickness of electrolyte layer on the Pt WE was equal to 21.3 nm. This value was calculated *via* the inverse Beer–Lambert relation using the attenuation of the Pt 4f core level signal passing from hydrated conditions (saturated environment before “dip & pull” procedure) to the final solid/liquid interface configuration.^{31,36} We have determined that a bulk (unmasked) current density of about 1.0 mA cm⁻² at +900 mV (overpotential η = +695 mV, OER conditions) corresponds to *ca.* 0.3 mA cm⁻² in the nanometric-

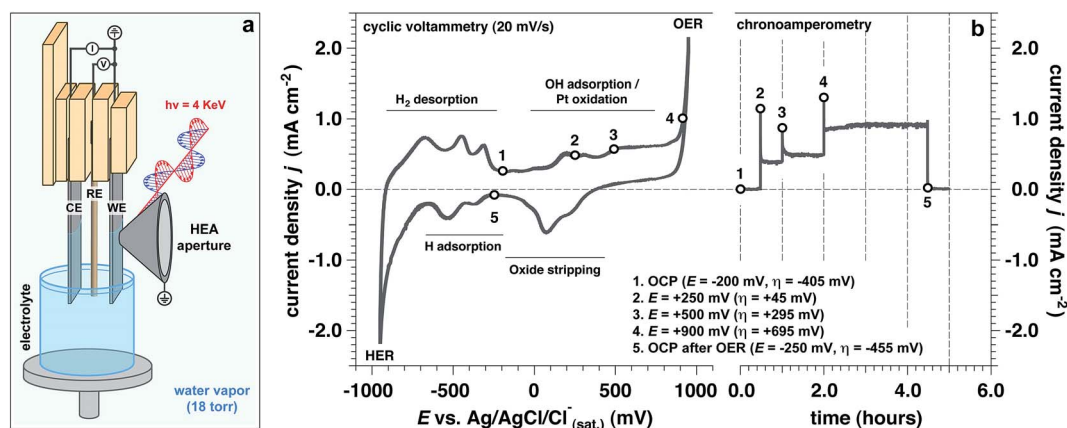
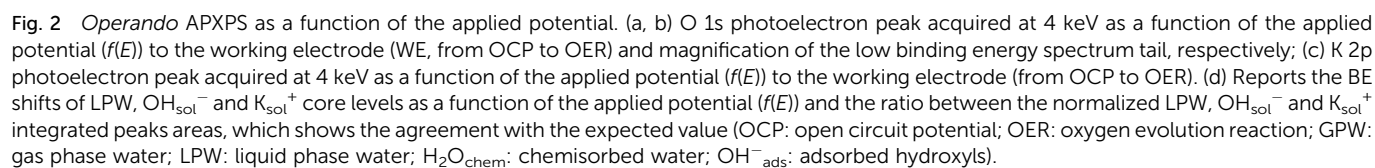


Fig. 1 Experimental set-up and electrochemical characterization. (a) Three electrode electrochemical set up, used in this work (CE: counter electrode; RE: reference electrode; WE: working electrode, HEA: hemispherical electron analyzer); (b) cyclic voltammetry (20 mV s⁻¹) and chronoamperometry for a polycrystalline Pt electrode in alkaline conditions (measured pH 13.9) at room temperature and for a water pressure in the chamber equal to 18 Torr (OCP: open circuit potential; HER: hydrogen evolution reaction; OER: oxygen evolution reaction).

The binding energy (BE) trends of the O 1s LPW, OH_{sol}⁻ and K 2p_{3/2} core levels present in the liquid layer as a function of the applied potential is reported in Fig. 2d. The BEs for elements in the electrolyte (BE_{electrolyte}) obey the relation:

$$\text{BE}_{\text{electrolyte}} = \text{BE}^0 - eE \quad (1)$$

E is the applied potential and BE^0 is the BE measured in absence of an applied potential. According to eqn (1), the aforementioned core levels shift toward lower BEs with respect to the Fermi level of the sample (grounded to the analyzer, see



schematization reported in Fig. 1a) for applied anodic potentials, thereby confirming that potential control is achieved in the measuring spot (1 : 1 shift).

From the O 1s low BE tail in Fig. 2b, it is possible to observe that the area of the Pt–O component (centered at 530.4 eV) increases as well as the adsorbed OH[−] (OH_{ads}[−] at 531.3 eV).

Finally, the area of the chemisorbed water component (H₂O_{chem} centered at 532.4 eV) also remains constant as the potential changes from OCP to +500 mV, with an increase under OER at +900 mV. Interestingly, upon removing the potential (“OCP after OER”), the peak area does not appreciably change. Such observation is consistent with an increase of the electrode surface area under catalytic conditions, which will be detailed in a following section.

Evolution of Pt surface chemistry as a function of the applied potential

The Pt 4f spectra (Fig. 3) can be de-convoluted into four different spectral components: metal Pt (71.2 eV), Pt^δ–OH_{ads} (71.8 eV), Pt^(II) (72.5 eV), and Pt^(IV) (74.6 eV).^{27,41} Among these different components, Pt^δ–OH_{ads} provides insights into the OER mechanism at the Pt/electrolyte interface.

As proposed by Krasilshchikov²⁸ and Damjanovic,^{24,26} it is well-accepted that under alkaline conditions the OER on Pt starts with the nucleophilic attack of OH[−] ion on Pt followed by

a fast electron transfer to the metastable configuration where OH[−] is then chemisorbed on an “activated” platinum site (Pt^δ).^{24,26} The formal positive charge of the Pt^δ (set between Pt metal and Pt^(II)) in the Pt^δ–OH_{ads} active site, as suggested by Wagner *et al.*,⁴² could explain the spectral position of its corresponding photoelectron peak, centered between the metal and Pt^(II) spectral components.

Interestingly, while the integrated area of this component remains unchanged with applied potentials below the OER threshold, it drastically increases under OER conditions. This finding is supported by a similar trend observed in the O 1s spectrum for the OH_{ads}[−] (Fig. 2a and b). At OCP after OER, the intensity of this component is comparable to the initial OCP state. This reversibility suggests that the formation of this component is driven by high overpotentials (*i.e.* under OER conditions). Since oxygen evolution in alkaline environments is proceeded by the deprotonation of adsorbed OH[−], we conclude that the observed spectral feature, Pt^δ–OH_{ads}, is representing the active species in the OER catalytic cycle.

Together with Pt^δ–OH_{ads}, the amount of Pt^(II) species increases with the applied potentials, where we observe a sharp increase once the potential is raised to +900 mV (OER conditions). Under this applied potential, the formation of Pt^(IV)O₂ oxide becomes prominent concurrently with the previously discussed Pt species (Fig. 3).

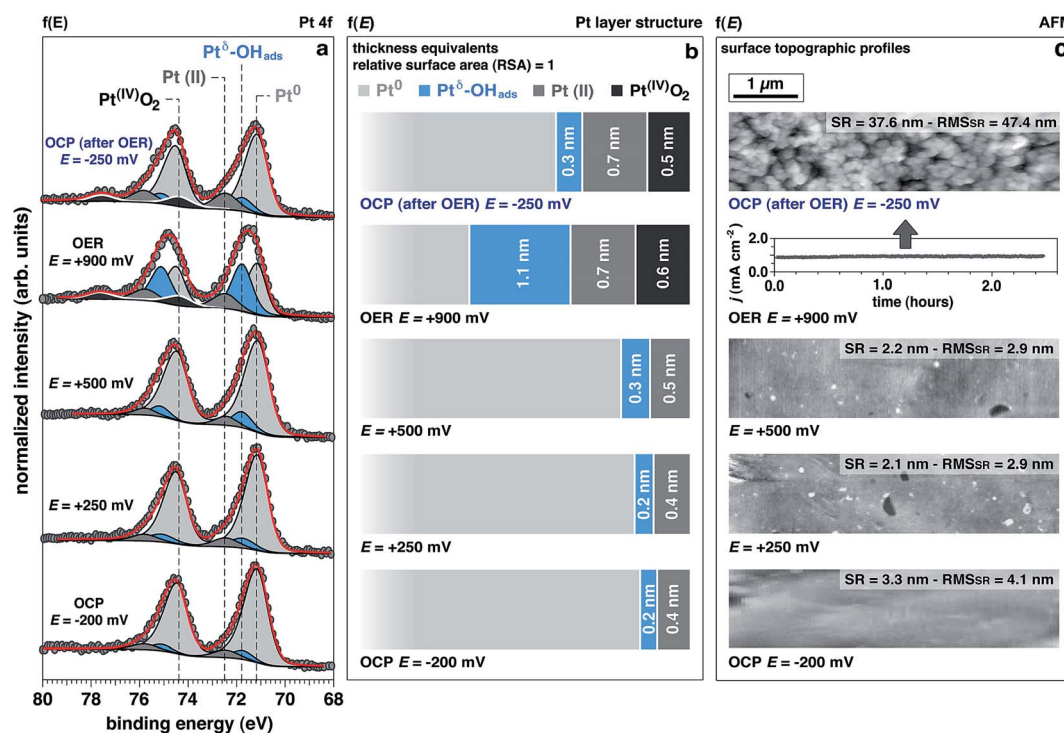


Fig. 3 Evolution of the surface chemistry studied by *operando* APXPS, as a function of the applied potential. (a) Pt 4f spectra acquired at 4 keV as a function of the applied potential ($f(E)$) to the working electrode (from OCP to OER) (OCP: open circuit potential; OER: oxygen evolution reaction; OH_{ads}: adsorbed hydroxyls). (b) Evolution of the surface structure as a function of the applied potential ($f(E)$), from OCP to OER. The numerical simulations allowed to retrieve information about the spatial localization of the different Pt chemistries. Thickness values are provided in “thickness equivalents” as the sample is not atomically flat, and the APXPS gives average information over a surface region of 300 μm in diameter (OCP: open circuit potential; OER: oxygen evolution reaction); (c) peak force tapping mode *ex situ* AFM topographies of the Pt surface taken after 120 min resting at the reported potentials (SR: surface roughness; RMSR: surface roughness root-mean-square).



Understanding the surface and sub-surface structure of Pt electrode under OER conditions

Using numerical simulations of the individual Pt component photoelectron attenuation (averaged over a 300 μm -diameter surface area region, equal to the analyzer cone aperture), we were able to qualitatively determine the surface and sub-surface structure as a function of the applied potential (Fig. 3b). We evaluated a wide range of Pt species configurations ranging in thickness and relative locations, finding that the structure that best matches with the experimental data is constituted by stacked layers of Pt species. In an effort to simplify the simulations, dense layers referred to as “equivalent thickness” are used to represent the actual porosity of the Pt electrode upon OER reconstruction (see detailed explanations about the numerical procedure reported in the Experimental section and ESI Fig. 4†). At OCP and increasing the potential to +500 mV, the Pt WE possesses a sub-surface domain of $\text{Pt}^{\delta}\text{-OH}_{\text{ads}}$ and an outer (electrolyte facing) layer of Pt^{III} , present in the form of $\text{Pt}^{\text{III}}(\text{OH})_2$ and $\text{Pt}^{\text{III}}\text{O}$. Under OER conditions at +900 mV, the outermost layer presents an additional layer of $\text{Pt}^{\text{IV}}\text{O}_2$, while the $\text{Pt}^{\delta}\text{-OH}_{\text{ads}}$ increases significantly. The spatial continuity between the Pt^{III} and the Pt^{IV} layers is ensured by an interfacial region of mixed $\text{Pt}^{\text{II, IV}}$ oxides. As reported in literature,^{24,25,27,43,44} our data and simulations confirm that the highly electroactive layer constituted by $\text{Pt}^{\delta}\text{-OH}_{\text{ads}}$ is localized sub-surface, between the metal substrate and an overlayer constituted by Pt^{III} and Pt^{IV} species.

The evolution of the outer surface morphology as a function of the applied potential was studied by *ex situ* AFM (Fig. 3c). The surface undergoes a clear reconstruction upon OER conditions transitioning from a flat metallic surface at OCP and low applied potentials (as confirmed by the low surface roughness (SR) values) to a high surface area oxidic system after catalytic conditions, characterized by a much higher SR than the pristine surface. Therefore, we believe that the so-called “gelly structure” reported in literature for Pt surfaces under OER conditions in acidic electrolytes,^{27,43,44} also occurs when the reaction is driven in alkaline environments. Using the previously described simulations, we can investigate the chemistry of the mesoporous (“gelly”) structure, which is composed of a $\text{Pt}^{\text{IV}}\text{O}_2$ outer layer and an inner layer (on top of the $\text{Pt}^{\delta}\text{-OH}_{\text{ads}}$ active phase) constituted by a mixture of Pt^{III} phases consisting of $\text{Pt}^{\text{III}}(\text{OH})_2$ and $\text{Pt}^{\text{III}}\text{O}$.

Experimental surface observations compared to thermodynamics predictions

Interestingly, the measured Pt electrode properties differ drastically from the predicted Pt/ H_2O Pourbaix diagram⁴⁵ (ESI Fig. 5†). In alkaline conditions (pH 13.9, KOH 1.0 M) at the OCP, Pt should partially oxidize to a $\text{Pt}^{\text{III}}(\text{OH})_2$ phase. Upon increasing the potentials to +250 mV, +500 mV, and +900 mV (+447 mV, +697 mV, and +1097 mV vs. normal hydrogen electrode, respectively), the only thermodynamically stable phase the Pourbaix diagram predicts is $\text{Pt}^{\text{IV}}\text{O}_2$.

On the other hand, the experimental results clearly shows that up to +500 mV $\text{Pt}^{\delta}\text{-OH}_{\text{ads}}$ and Pt^{III} species are present on the surface (Fig. 3). Notably, the Pt^{IV} spectral component is

eventually observed only under the OER potential (+900 mV), therefore at a much higher potentials than the thermodynamically predicted in the Pourbaix diagram at the same pH. Such a deviation can be due to kinetic effects that are not taken into account in a merely thermodynamic investigation of the stability of Pt phases. In addition, at OCP after OER, the surface chemistry exhibits a partial chemical hysteresis. While the $\text{Pt}^{\delta}\text{-OH}_{\text{ads}}$ feature decreases (as a result of the potential removal, as previously discussed), it is still possible to observe both Pt^{III} and Pt^{IV} spectral fingerprints. On the contrary, the Pourbaix diagram predicts only Pt^{III} (present as $\text{Pt}^{\text{III}}(\text{OH})_2$) as the stable phase at this pH value.

Pt surface chemistry evolution under OER conditions

The time evolution of the Pt 4f spectrum and accompanying Pt^0 -normalized integrated areas of the Pt spectral components under OER conditions at +900 mV is reported in Fig. 4a and b, respectively. The same time evolution for the O 1s and K 2p core levels is reported in ESI Fig. 6,† together with the ionic concentration and the thickness of the electrolyte layer determined as a function of the observation time. Between 0 to 60 minutes under OER conditions we can observe the initial transitions for $\text{Pt}^{\delta}\text{-OH}_{\text{ads}}$, Pt^{III} , and Pt^{IV} before stabilizing and reaching steady state conditions (>60 minutes). During this period, it can be seen that the $\text{Pt}^{\delta}\text{-OH}_{\text{ads}}/\text{Pt}^0$ and $\text{Pt}^{\text{IV}}/\text{Pt}^0$ ratios increase accompanied by a slight decrease of the $\text{Pt}^{\text{III}}/\text{Pt}^0$ ratio with respect to time, Fig. 4b. Using our previously established layered model, we used numerical simulations to estimate the respective “equivalent thicknesses” for each Pt phase (i) and

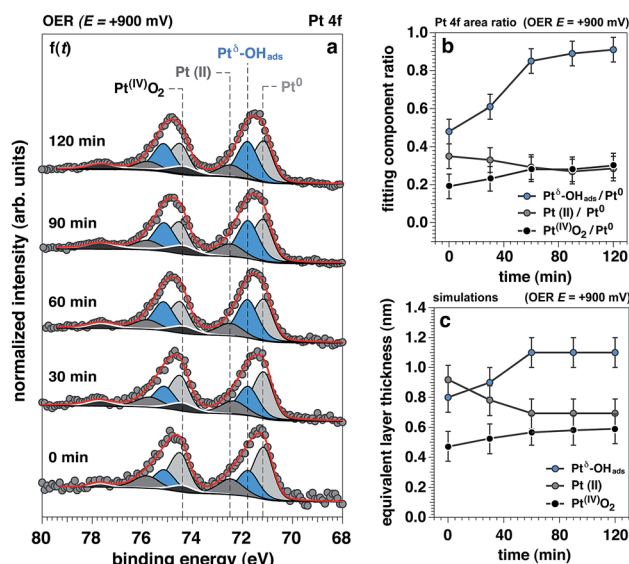


Fig. 4 Chemical and structural evolution of platinum surface under oxygen evolution conditions, as a function of time. (a) Evolution of Pt 4f APXPS photoelectron peak acquired at 4 keV under oxygen evolution reaction (OER) conditions (+900 mV), as a function of the observation time ($f(t)$); (b, c) experimental trends and simulated layer thicknesses for the oxidized Pt components as a function of the observation time under OER conditions, respectively.



retrieve their in-depth localization as a function of time under catalytic conditions (ii) (Fig. 4c).

Proposed Pt structure under catalytic conditions and OER mechanism in alkaline environments

We can now propose a more complete mechanism for platinum OER under alkaline conditions (Fig. 5) by combining our steady state and dynamic information detailed in Fig. 3 and 4 with previous literature. The mesoporosity^{27,43,44} of the Pt electrode (as seen in Fig. 3c and reported in ref. 27, 43 and 44) is present throughout the near surface structure, and it is composed of 3 regions. Moving from the Pt⁰ substrate to the outer surface, we have demonstrated the presence of Pt^δ-OH_{ads}, Pt^(II) species (Pt^(II)(OH)₂ and Pt^(II)O) and Pt^(IV)O₂. As proposed by Krasilshchikov²⁸ and Damjanovic^{24,26} for oxygen evolution on Pt in alkaline conditions, the formation of the Pt^δ-OH_{ads} phase is promoted by the nucleophilic attack of hydroxide ions to the metal Pt surface, followed by a fast electron transfer to the metastable configuration where OH⁻ is chemisorbed on an “activated” platinum site (Pt^δ).^{24,26} Overall, this can be rationalized by a mixed electrostatic and thermodynamic effect. As the potential applied to the WE becomes more anodic, the local increase of hydroxyl concentration (within the inner Helmholtz layer, IHL) becomes more pronounced. At the same time the high positive potential at the electrode surface lowers the Pt/OH⁻ surface binding energy whilst the high pH value (13.9) decreases the electrochemical potential of the electrolyte,⁴⁶ thereby favoring the activation of the Pt surface *via* hydroxyl adsorption followed by the electron transfer and the consequent

formation of the Pt^δ-OH_{ads} species.^{47–49} The final surface coupling and deprotonation of the adsorbed OH⁻ leads to the evolution of oxygen. Interestingly, as it can be seen from the time trends reported in Fig. 4b and c, only when a considerable amount of Pt^δ-OH_{ads} is present on the sub-surface region the surface coupling process and the consequent O₂ evolution can occur. This result is extremely important since it directly shows a kinetic limitation that contributes to the observed OER overpotential. The proposed mechanism is schematically represented in Fig. 5.

It is important to consider the possibility for Pt^(IV) dissolution in the form of a Pt^(IV) aquocomplex as indicated by the Pourbaix diagram. Based on our data, we do not observe any Pt dissolution during the potentiostatic measurements. However, one must consider that reversing (or removing) potentials could induce Pt dissolution on the order of tens of nM as reported by Lopes *et al.*¹⁵ The reported dissolution rates are considerably below the detection limit of the APXPS (about 100 mM). Therefore, although we would not be able to detect this process, a Pt dissolution cannot be excluded.^{50,51}

Conclusions

This study revealed the complexity and significant surface chemical reconstruction that takes place on a Pt electrode, in particular on the nature of Pt^δ-OH_{ads} species, and its crucial role on the OER mechanism. Our findings show that the presence of Pt^δ-OH_{ads} is triggered by potentials within the OER region: upon removing the applied potential, the concentration of this species drops to similar values observed for the initial OCP whilst the Pt(II) and Pt(IV) oxides chemistries remained essentially unchanged. This observation, together with the fact that the OER in alkaline conditions occurs *via* the oxidation of the hydroxyl, allows us to conclude that the Pt^δ-OH_{ads} constitutes the actual rate determining species for oxygen evolution. The combination of APXPS, AFM, and numerical simulations allowed us to provide new insights into the chemical structure of the Pt electrode under OER conditions, showing that the Pt^δ-OH_{ads} layer is located sub-surface. The presence of the Pt^δ-OH_{ads} layer leads to the formation of a high surface area oxidic outer layer in which OH⁻ need to diffuse in order to activate the oxygen evolution.

This observation allows us to propose that the applied overpotential (above the thermodynamic oxidation potential of water) has two main effects. First, it positively polarizes the electrode surface, which lowers the binding energy of hydroxyl ions and increase their adsorption rate. Second, the OH⁻ may electrostatically be driven through the oxidic porous passivation layer.⁵² Therefore, from the experimental observations reported here we provide motivation for reevaluating the kinetic limitations (overpotentials) of OER on Pt to take into account both adsorption/electron transfer rate and mass transport effects through the modified electrode under catalytic conditions.

Moreover, we have elucidated the differences between thermodynamic predictions of the Pt phase stability and the actual surface chemistry under polarization. The high chemical sensitivity of XPS probing the solid/liquid electrified interface

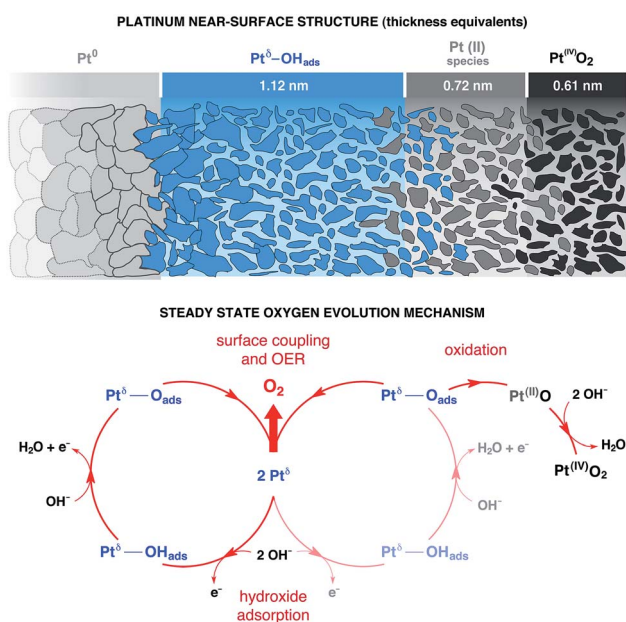


Fig. 5 Proposed catalytic mechanism and phase structure for oxygen evolution on platinum under alkaline conditions. The evaluation of the catalytic loop for oxygen evolution reaction (OER) on Pt (pH 13.9), as well as the structural description of the catalytic state, have been performed by synergistically coupling the experimental results from the *operando* APXPS and the corresponding numerical simulations.



has allowed us to re-interpret the Pourbaix diagram of Pt. Thus motivating the need for these types of *operando* experiments and the development of new thermodynamic, kinetic, and computational models for electrified interfaces under realistic operating conditions.⁵³ Expanding this experimental strategy for the study of different materials and catalytic reactions under *operando* conditions,^{31,32,54} will open a path for new discoveries that will deepen our knowledge towards the accelerated development of novel tailored materials.

Experimental

Dip & pull method and *operando* measurements

The three electrodes were mounted into a PEEK electrode housing that was attached to a multi-axis manipulator. Electrical feedthrough within the manipulator connected the WE, RE, and CE to an external potentiostat/galvanostat (Biologic SP 300) to perform *operando* electrochemistry measurements. During the operation, the WE and the analyzer front cone were commonly grounded.

Prior its introduction into the experimental chamber, the electrolyte (KOH 1.0 M aqueous solution) was outgassed for at least 30 min at low pressure (around 10 Torr) in a dedicated off-line chamber. Then, once the manipulator (Fig. 1a) and the outgassed electrolyte were placed into the APXPS experimental chamber, the pressure was carefully lowered down to the water vapor pressure (between 16 and 20 Torr, at r.t.).

To create a solid/liquid interface (WE/electrolyte), all three electrodes were immersed into the electrolyte. Then they were slowly extracted from the electrolyte solution by rising the manipulator at a constant vertical rate. Following this procedure, a thin layer of liquid electrolyte film is formed on the Pt electrode (typical values ranges between 10 and 30 nm); the three electrodes were then positioned at the focal point of the analyzer, giving the possibility of performing XPS investigation of the solid/liquid interface as a function of the applied potential. During the measurements, the bottom parts of the electrodes were kept into the bulk electrolyte, in order to ensure electrical continuity between the thin electrolyte layer on the WE surface and the bulk electrolyte itself.

Electrochemical measurements

The chemicals employed here were high purity reagents and they were used as received without further operations. MilliQ water (DI, $\rho = 18.2 \text{ M}\Omega \text{ cm}$) and potassium hydroxide (KOH, 99.99%, Aldrich) were used as solvent and supporting electrolyte, respectively.

The working (WE) and counter electrodes (CE, Pt polycrystalline foils, 99.99%, thickness 0.5 mm, Aldrich) were polished to a mirror finish with silicon carbide papers of decreasing grain size (Struers, grit: 2400 and 4000). The samples were then cleaned by two cycles of ultrasonic treatment in a mixture of MilliQ water/ethanol (Aldrich, 1 : 1) for 10 min. A third ultrasonic cleaning was then conducted in pure MilliQ water for 15 min, followed by a thoroughly rinsing and drying in N_2 stream. As reference electrode (RE), a miniaturized leakless

$\text{Ag}/\text{AgCl}/\text{Cl}_{(\text{sat.})}^-$ RE (ET072-1, eDAQ) was used (standard electrode potential $E_{\text{Ag}/\text{AgCl}_{(\text{sat.})}}^\circ = 197 \text{ mV}$ with respect to the normal hydrogen electrode, NHE). All the potentials reported in this work are referred to this RE unless differently stated.

An electrochemical cleaning procedure was conducted by holding the Pt WE at -1200 mV (within the HER region) for 30 minutes, in order to obtain a pure metallic surface and a homogeneous surface, increasing in this way the experimental reproducibility. The cleaning procedure was stopped after reaching the current open circuit potential (OCP) of the cell from the cathodic side. The data acquisition started only when the OCP reached a stable value over the observation time (about 30 minutes). All the voltammetric measurements were conducted at a scan rate of 20 mV s^{-1} .

Electrochemical impedance spectroscopy (EIS) measurements were carried out with the aim of electrochemically evaluate the interfacial properties of the solid/liquid interface. EIS measurements were conducted by measuring the response in current after applying to the WE a sinusoidal wave of potential of 10 mV peak-to-peak voltage amplitude, at OCP (-200 mV) and OER conditions ($+900 \text{ mV}$). The frequency of the sinusoidal potential wave was ranging from 105 Hz to 5 mHz , scanning 10 frequencies per decade.

Beamline 9.3.1 and APXPS experimental details

Beamline 9.3.1 at Advanced Light Source (ALS, Lawrence Berkeley National Laboratory) is equipped with a bending magnet and a Si (111) double crystal monochromator (DCM) having a total energy range between 2.0 keV and 7.0 keV ("tender" X-ray range). The minimal spot size at the beamline is $0.7 \text{ mm (v)} \times 1.0 \text{ mm (h)}$.

For the data acquisition, the pass energy of the Scienta analyzer (R4000 HiPP-2) was set to 200 eV , using a step of 100 meV and a dwell time of 300 ms . Under these conditions, the total resolution (X-rays and analyzer) was equal to about 250 meV at room temperature (r.t.) and at 4 keV . The measurements were taken using a photon energy of 4 keV at r.t. and in normal emission (NE), at a pressure in the experimental chamber matching the water vapor tension at r.t., between 16 and 18 Torr. The detection stage in the analyzer (multichannel plate, MCP) was under high vacuum conditions ($\sim 10^{-7}$ Torr) throughout the all experiment, thanks to the differentially pumped electrostatic lens system based on four independent pumping stages. To limit the evaporation from the electrochemical cell we place a larger outgassed solution buffer in the analysis chamber to facilitate the fully humidified environment at room temperature. The calibration of the binding energy (BE) scale was carried out using the Au 4f photoelectron peak as reference ($4f_{7/2} \text{ BE} = 84.0 \text{ eV}$) as well as the Fermi edge, from a clean gold polycrystalline surface measured under all the investigated experimental conditions.

All the fit reported in this work have been carried out using a Doniach–Šunjić shape for the Au and Pt 4f photoelectron peaks, whereas a symmetrical Voigt function (G/L ratio ranging from 85/15 to 75/25) was used to fit the O 1s photoelectron peaks (after Shirley background subtraction).³⁶ During the



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- 7 B. K. Carpenter, J. N. Harvey and A. J. Orr-Erwing, *J. Am. Chem. Soc.*, 2016, **138**, 4695–4705.
- 8 B. E. Conway and T. C. Liu, *Langmuir*, 1990, **6**, 268–276.
- 9 N. M. Markovic, H. A. Gasteiger and P. N. Ross, *J. Phys. Chem.*, 1996, **100**, 6715–6721.
- 10 V. Stamenkovic, B. S. Mun, K. J. J. Mayrhofer, P. N. Ross, N. M. Markovic, J. Rossmeisl, J. Greeley and J. K. Nørskov, *Angew. Chem., Int. Ed.*, 2006, **118**, 2963–2967.
- 11 W. Sheng, H. A. Gasteiger and Y. Shao-Horn, *J. Electrochem. Soc.*, 2010, **157**, B1529–B1536.
- 12 I. E. L. Stephens, A. S. Bondarenko, F. J. Perez-Alonso, F. Calle-Vallejo, L. Bech, T. P. Johansson, A. K. Jepsen, R. Frydendal, B. P. Knudsen, J. Rossmeisl and I. Chorkendorff, *J. Am. Chem. Soc.*, 2011, **133**, 5485–5491.
- 13 T. Reier, M. Oezaslan and P. Strasser, *ACS Catal.*, 2012, **2**, 1765–1772.
- 14 I. E. L. Stephens, A. S. Bondarenko, U. Grønby, J. Rossmeisl and I. Chorkendorff, *Energy Environ. Sci.*, 2012, **5**, 6744–6762.
- 15 P. P. Lopes, D. Strmcnik, D. Tripkovic, J. G. Connell, V. Stamenkovic and N. M. Markovic, *ACS Catal.*, 2016, **6**, 2536–2544.
- 16 D. Strmcnik, M. Uchimura, C. Wang, R. Subbaraman, N. Danilovic, D. Van der Vliet, A. P. Paulikas, V. Stamenkovic and N. M. Markovic, *Nat. Chem.*, 2013, **5**, 300306.
- 17 P. Hernandez-Fernandez, F. Masini, D. N. McCarthy, C. E. Streb, D. Friebe, D. Deiana, P. Malacrida, A. Nierhoff, A. Bodin, A. M. Wise, J. H. Nielsen, T. W. Hansen, A. Nilsson, I. E. L. Stephens and I. Chorkendorff, *Nat. Chem.*, 2014, **6**, 732–738.
- 18 D. Strmcnik, K. Kodama, D. Van Der Vliet, J. Greeley, V. R. Stamenkovic and N. M. Markovic, *Nat. Chem.*, 2009, **1**, 466–472.
- 19 D. Strmcnik, V. Tripkovic, D. Van Der Vliet, K.-C. Chang, V. Komanicky, H. You, G. Karapetrov, J. P. Greeley, V. R. Stamenkovic and N. M. Markovic, *J. Am. Chem. Soc.*, 2008, **130**, 15332–15339.
- 20 L. Gan, C. Cui, M. Heggen, F. Dionigi, S. Rudi and P. Strasser, *Science*, 2014, **346**, 1502–1506.
- 21 M. Escudero-Escribano, P. Malacrida, M. H. Hansen, U. G. Vej-Hansen, A. Velazquez-Palezuela, V. Tripkovic, J. Schiøtz, J. Rossmeisl, I. E. L. Stephens and I. Chorkendorff, *Science*, 2016, **352**, 73–76.
- 22 P. P. Lopes, D. Strmcnik, J. S. Jirkovsky, J. G. Connell, V. Stamenkovic and N. M. Markovic, *Catal. Today*, 2016, **262**, 41–47.
- 23 L. F. Sallum, E. R. Gonzalez and J. M. Feliu, *Electrochem. Commun.*, 2016, **72**, 83–86.
- 24 A. Damjanovic, A. Dey and J. O. 'M. Bockris, *Electrochim. Acta*, 1966, **11**, 791–814.
- 25 D. Gilroy and B. E. Conway, *Can. J. Chem.*, 1968, **46**, 875–890.
- 26 I. V. Birss and A. Damjanovic, *J. Electrochem. Soc.*, 1987, **134**, 113–117.
- 27 R. Arrigo, M. Hävecker, M. E. Schuster, C. Ranjan, E. Stotz, A. Knop-Gericke and R. Schlögl, *Angew. Chem., Int. Ed.*, 2013, **52**, 1–6.
- 28 A. L. Krasilshchikov, *Zh. Fiz. Khim.*, 1963, **37**, 531.
- 29 N. Danilovic, R. Subbaraman, K.-C. Chang, S.-H. Chang, Y. J. Kang, J. Snyder, A. P. Paulikas, D. Strmcnik, Y.-T. Kim, D. Myers, V. Stamenkovic and N. M. Markovic, *J. Phys. Chem. Lett.*, 2014, **5**, 2474–2478.
- 30 Y. Surendranath, M. W. Kanan and D. G. Nocera, *J. Am. Chem. Soc.*, 2010, **132**, 16501–16509.
- 31 E. J. Crumlin, Z. Liu, H. Bluhm, W. Yang, J. Guo and Z. Hussain, *J. Electron Spectrosc. Relat. Phenom.*, 2015, **42**, 264–273.
- 32 H.-J. Lewerenz, M. F. Lichterman, M. H. Richter, E. J. Crumlin, S. Hu, S. Axnanda, M. Favaro, W. Drisdell, Z. Hussain, B. S. Brunshawig, A. Nilsson, A. Bell, N. S. Lewis and D. Friebe, *Electrochim. Acta*, 2016, **211**, 711–719.
- 33 Y. T. Law, S. Zafeiratos, S. G. Neophytides, A. Orfanidi, D. Costa, T. Dintzer, R. Arrigo, A. Knop-Gericke, R. Schloegl and E. R. Savinova, *Chem. Sci.*, 2015, **6**, 5635–5642.
- 34 Z. Zhu, F. Tao, F. Zheng, R. Chang, Y. Li, L. Heinke, Z. Liu, M. Salmeron and G. A. Somorjai, *Nano Lett.*, 2012, **12**, 1491–1497.
- 35 S. Axnanda, E. J. Crumlin, B. Mao, S. Rani, R. Chag, P. G. Karlsson, M. O. M. Edwards, M. Lundqvist, R. Moberg, P. N. Ross, Z. Hussein and Z. Liu, *Sci. Rep.*, 2015, **5**, 9788.
- 36 M. Favaro, B. Jeong, P. N. Ross, J. Yano, Z. Hussain, Z. Liu and E. J. Crumlin, *Nat. Commun.*, 2016, **7**, 12695.
- 37 M. Pourbaix, *National Association of Corrosion Engineers*, 1974.
- 38 C. A. Tolman, *Chem. Soc. Rev.*, 1972, **1**, 337–353.
- 39 M. F. Lichterman, S. Hu, M. H. Richter, E. J. Crumlin, S. Axnanda, M. Favaro, W. Drisdell, Z. Hussein, T. Mayer, B. S. Brunshawig, N. S. Lewis, Z. Liu and H.-J. Lewerenz, *Energy Environ. Sci.*, 2015, **8**, 2409–2416.
- 40 A. Eilert, F. Cavalca, F. S. Roberts, J. Osterwalder, C. Liu, M. Favaro, E. J. Crumlin, H. Ogasawara, D. Friebe, L. G. M. Pettersson and A. Nilsson, *J. Phys. Chem. Lett.*, 2017, **8**, 285–290.
- 41 V. A. Saveleva, V. Papaefthimiou, M. K. Daletou, W. H. Doh, C. Ulhaq-Buillet, M. Diebold, S. Zafeiratos and E. R. Savinova, *J. Phys. Chem. C*, 2016, **120**, 15930–15940.
- 42 F. T. Wagner and T. E. Moylan, *Surf. Sci.*, 1987, **191**, 121–146.
- 43 M. Peuckert and H. Ibach, *Surf. Sci.*, 1984, **136**, 319–326.
- 44 A. E. Bolzan and A. J. Arvia, *J. Electroanal. Chem.*, 1994, **375**, 157–162.
- 45 The thermodynamic predictions of the stability of the different Pt species as a function of applied potential and pH have been determined by simulating the corresponding Pourbaix diagram with the Materials Project facility A. Jain, S. P. Ong, G. Hautier, W. Chen, W. D. Richards, S. Dacek, S. Cholia, D. Gunter, D. Skinner, G. Ceder and K. A. Persson, *APL Mater.*, 2013, **1**, 011002; A. Jain, G. Hautier, S. P. Ong, C. Moore, C. Fischer, K. Persson and G. Ceder, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2011, **84**, 045115.



- 46 P. Daubinger, J. Kieninger, T. Unmüssig and G. A. Urban, *Phys. Chem. Chem. Phys.*, 2014, **16**, 8392–8399.
- 47 G. B. Fisher and B. A. Sexton, *Phys. Rev. Lett.*, 1980, **44**, 683–686.
- 48 A. V. Melo, W. E. O'Grady, G. S. Chottiner and R. W. Hoffman, *Appl. Surf. Sci.*, 1985, **21**, 160–167.
- 49 S. Yamamoto, H. Bluhm, K. Andersson, G. Ketteler, H. Ogasawara, M. Salmeron and A. Nilsson, *J. Phys.: Condens. Matter*, 2008, **20**, 184025.
- 50 T. Binninger, R. Mohamed, K. Waltar, E. Fabbri, P. Levecque, R. Kötz and T. J. Schmidt, *Sci. Rep.*, 2015, **5**, 12167.
- 51 H. Dau, C. Limberg, T. Reier, M. Risch, S. Roggan and P. Strasser, *ChemCatChem*, 2010, **2**, 724–761.
- 52 J. Yang, J. K. Cooper, F. M. Toma, K. A. Walczak, M. Favaro, J. W. Beeman, L. H. Hess, C. Wang, C. Zhu, S. Gul, J. Yano, C. Kisielowski, A. Schwartzberg and I. D. Sharp, *Nat. Mater.*, DOI: 10.1038/nmat4794.
- 53 F. M. Toma, J. K. Cooper, V. Kunzelmann, M. T. McDowell, J. Yu, D. M. Larson, N. J. Borys, C. Abelyan, J. W. Beeman, K. M. Yu, J. Yang, L. Chen, M. R. Shaner, J. Spurgeon, F. A. Houle, K. A. Persson and I. D. Sharp, *Nat. Commun.*, 2016, **7**, 12012.
- 54 M. Favaro, W. S. Drisdell, M. A. Marcus, J. M. Gregoire, E. J. Crumlin, J. A. Haber and J. Yano, *ACS Catal.*, 2017, **7**, 1248–1258.
- 55 S. Evans, *Surf. Interface Anal.*, 1991, **17**, 85–93.
- 56 J. Muñoz-Flores and A. Herrera-Gomez, *J. Electron Spectrosc. Relat. Phenom.*, 2012, **184**, 533–541.
- 57 W. Smekal, W. S. M. Werner and C. J. Powell, *Surf. Interface Anal.*, 2005, **37**, 1059–1067.
- 58 S. Techane, D. R. Baer and D. G. Castner, *Anal. Chem.*, 2011, **83**, 6704–6712.
- 59 W. S. M. Werner, M. Chudzicki, W. Smekal and C. J. Powell, *Appl. Phys. Lett.*, 2014, **104**, 233106.
- 60 W. S. M. Werner, W. Smekal, T. Hisch, J. Himmelsbach and C. J. Powell, *J. Electron Spectrosc. Relat. Phenom.*, 2013, **190**, 137–143.
- 61 H. Brune, H. Röder, C. Boragno and K. Kern, *Phys. Rev. Lett.*, 1994, **73**, 1955–1958.
- 62 K. Bromann, H. Brune, H. Röder and K. Kern, *Phys. Rev. Lett.*, 1995, **75**, 677–680.

