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Latest advances on in situ and operando X-ray-based techniques in photoelectrocatalytic systems

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Abstract

In situ and operando X-ray techniques have emerged as powerful tools for unravelling the complex mechanisms underlying photoelectrochemical transformations. These techniques offer real-time insights into the dynamic processes occurring at the electrode-electrolyte interface during solar-driven water splitting and other PEC reactions. The present work aims to summarise the latest advances on in situ and operando X-ray absorption spectroscopy (XAS), X-ray diffraction (XRD), and X-ray photoelectron spectroscopy (XPS) for the characterisation of photoelectrocatalytic systems and materials for the generation of solar fuels and added value chemicals. This work highlights the recent advancements using these techniques for elucidating the structural, chemical, and electronic properties of photoelectrocatalytic materials and interfaces during operation. Besides, this work gives a technical guidance for performing these measurements considering the experimental requirements for each of these spectroscopies. We also give an overview of the state-of-the-art of the different synchrotron-based techniques employed for the characterisation of photoelectrocatalytic materials, focusing on the possibilities of the studied techniques, cell designs and more relevant results.

Introduction

Artificial photosynthesis technologies have been attracting an increasing attention, from fundamental understanding, to, more recently, the industry, as an alternative source of fuels and added value chemicals using ubiquitous molecules, such as CO₂ and H₂O and sunlight.¹⁻³ These technologies are achieving higher Technology Readiness Levels (TRLs) of development in the recent years, and are becoming a real alternative for the generation of solar fuels and chemicals at an industrial level. In the case of photoelectrochemical (PEC) routes, different architectures could be considered: i) full photoelectrochemical systems, where both, anode and cathode contain photoabsorber materials, ii) electrocatalytic systems coupled to photovoltaic cells (PV-EC) and iii) photoelectrochemical systems coupled to a photovoltaic cells to reduce the overpotential for the target chemical reaction.⁴ In recent years, all these technologies have increased its maturity towards industrialisation. As explained in the recent work by G. Segev *et al.*⁵ lab-scale photoelectrochemical devices currently report efficiencies over 10%, which has been traditionally considered as the minimum threshold for any industrial application. However, in order to establish the photoelectrocatalytic approach at a relevant industrial level, both from the scientific and economic point of view, there are still many technological questions to be answered and physico-chemical mechanisms to be revealed. To tackle this problem, the scientific community is currently highly interested in operando and in situ methods which allow the characterisation of the photoelectrocatalytic materials under relevant working conditions.^{6,7}



Specifically, "in-situ" characterization refers to the study of materials, usually catalysts, and reactions in their natural working environment without disrupting experimental conditions. It is performed within the reactor or reaction cell during the reaction, but not necessarily under realistic operating conditions. On the other hand, "operando" characterization refers to the study of materials and catalytic reactions under real operating conditions. Both approaches are crucial for understanding and improving catalytic systems, but are applied in slightly different contexts depending on whether a study is needed under controlled conditions (in situ) or realistic catalytic conditions (operando).⁸

Spectroscopy-based techniques are powerful tools to investigate the photoelectrode-electrolyte interface, where information on the electronic and structural nature of the active sites can be obtained, upon a careful selection of excitation and probe sources.⁹ As such, there is a growing tendency in the field to perform in situ and operando analysis using many different techniques such as Raman,^{10,11} UV-Vis,¹² Fourier-transform infrared spectroscopy (FTIR),¹³ Electrochemical Impedance Spectroscopy (EIS),¹⁴ Incident Modulated Photocurrent Spectroscopy (IMPS),¹⁵ among others, due to the key importance of understanding the materials under real working conditions. Hence, the combination of in situ and operando synchrotron techniques with photoelectrocatalytic experiments is pivotal to understand the performance of these materials, in terms of stability, efficiency, and selectivity, under working conditions and thus, establishing this technology as an important element in the green energy transition.

In terms of X-ray based spectroscopy techniques, such as XPS, XAS and XRD, the majority of in situ and operando reported PEC studies have employed synchrotron radiation. This, due to the advantages of synchrotron radiation compared with other lab scale X-ray sources: i) high brightness and energy resolution, which allows high-speed data acquisition and more accurate data, ii) high spatial resolution, which allows precise material mapping, iii) wide energy range, permitting variable energy X-ray techniques (XAS) and conducting experiments at a range of selected optimum wavelengths (XPS and XRD), iv) it is polarized, permitting specific experiments to probe the magnetic and electronic properties of materials and v) it has a precise flashing time structure at a very high frequency, that can be used to study physical and chemical processes taking place at very short time scales.¹⁶

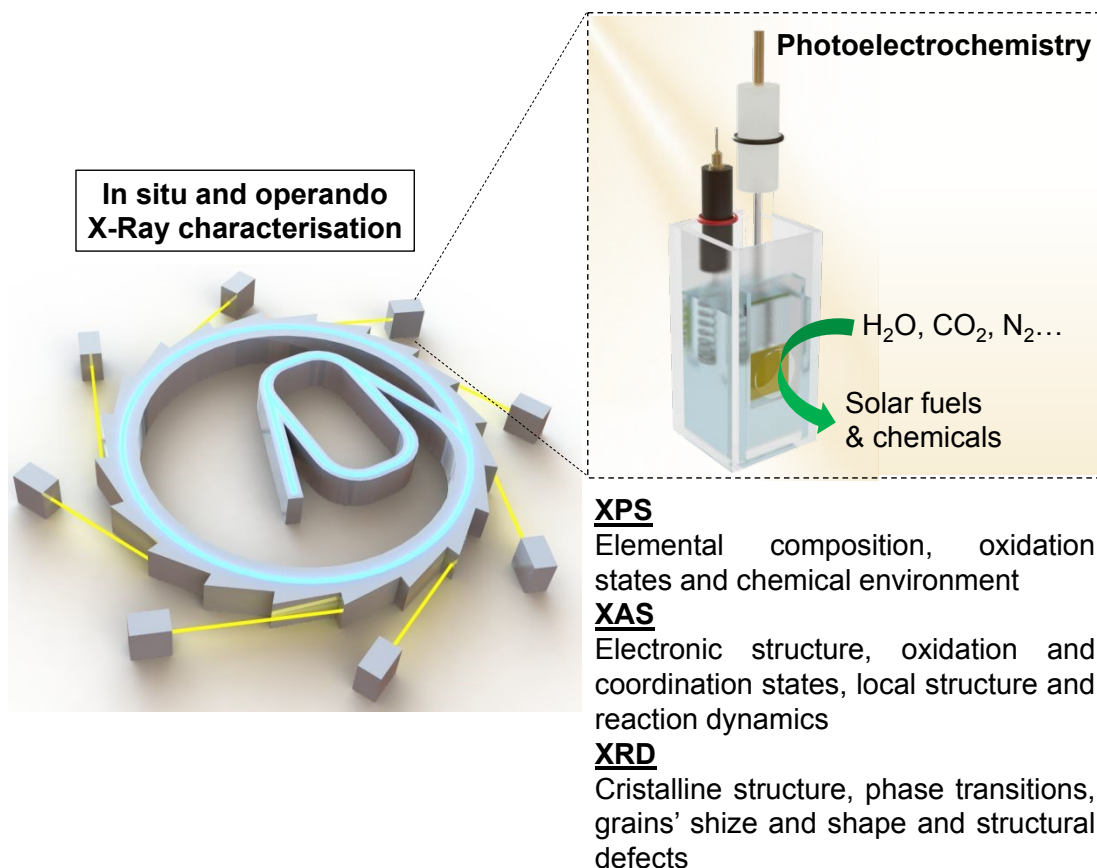
In situ and operando synchrotron techniques are more mature in the electrocatalytic (EC) field, which can be explained not only by the maturity of the field but also due to several inherent difficulties in order to study photoelectrocatalytic reactions in situ: i) although many novel oxide and chalcogenide electrocatalysts undergo bulk changes that leads to the actual catalytic species,^{17,18,19,20} the bulk of photoelectrodes are usually stable and most relevant changes are limited to their surfaces involving usually few tens of nm,^{21,22} which demands surface sensitive techniques or special analysis modes to make commonly bulk techniques surface sensitive, and both ii) the need to illuminate the sample during operation,²³ and iii) the usual employment of thick transparent conductive substrates (FTO or ITO, most frequently), instead of thin current collectors, add an extra level of complexity to the experimental set-up. Due to these challenges, only a few works have reported successful in situ or operando synchrotron-based spectroscopies in photoelectrochemical systems.

This review aims to give a general vision of the current state-of-the-art of in situ and operando X-ray-based studies, focusing mainly on X-ray Photoelectron Spectroscopy (XPS), X-ray Absorption Spectroscopy (XAS) and X-ray Diffraction Spectroscopy (XRD), in PEC systems for solar fuels generation (**Scheme 1**). Besides, this work also gives technical guidance for performing these experiments at synchrotron facilities, with the hope of increasing the activity in this field which has the potential to answer some of the main questions to take the comprehension of PEC systems to the next level.

By synthesizing and critically evaluating these analytical possibilities, this work aims to serve as a pivotal resource for guiding future research efforts and informing the design of more efficient and sustainable photoelectrocatalytic technologies. We believe that the comprehensive nature of this revision, its clear organization of existing knowledge, and its potential to advance the field,



make it well-suited for energy researchers interested in initiating or advancing their work with Article Online
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operando and in situ experiments in PEC systems.



Scheme 1. Overview of the addressed topics in the present review.

In situ and operando X-ray photoelectron spectroscopy (XPS)

X-ray photoelectron spectroscopy (XPS) is a commonly used characterisation technique for the analysis of the surface of materials. This technique, also called ESCA (Electron Spectroscopy for Chemical Analysis), provides qualitative and quantitative elemental analysis, except for hydrogen and helium, of the top 1-20 nm (depending on the sample and instrumental conditions) of a surface.²⁴ In XPS, the sample is irradiated with soft X-rays (energies lower than ~6 keV) and the kinetic energy of the emitted electrons is analysed. When an X-ray of known energy ($h\nu$), interacts with an atom, a photoelectron can be emitted via the photoelectric effect. The emitted photoelectron is the result of complete transfer of the X-ray energy to a core level electron. It simply states that the energy of the X-ray ($h\nu$) is equal to the binding energy (E_b) of the electron (how tightly it is bound to the atom/orbital to which it is attached), plus the kinetic energy (E_k) of the electron that is emitted and the spectrometer work function (Φ_{sp}), a constant value, and hence, the E_b can be calculated through the equation:

$$E_b = h\nu - E_k - \Phi_{sp} \text{ (Equation 1)}$$

Thus, valuable chemical information about the sample can be extracted because binding energies are sensitive to the chemical environment of the atom.²⁵ Hence, XPS provides unique information on the chemical composition and electronic structure of materials so that it has been traditionally used for the fundamental characterisation of semiconductors with application in the field of photoelectrochemistry. The development of near-ambient pressure (NAP) XPS has extended the applications of XPS to the performance of detailed studies of the chemical state of



photoelectrodes and the reaction intermediates under relevant or even operando conditions, which facilitate the understanding of reaction mechanisms. As a consequence, NAP XPS is nowadays commonly used for the in situ characterisation of catalytic, photocatalytic and (photo)electrocatalytic reactions. The field has evolved exponentially since the development of the new generation of NAP XPS analysers at the beginning of the century.²⁶ The last comprehensive review in 2020 estimates that there are more than 70 NAP XPS systems around the world, including laboratory and synchrotron based systems.²⁷ Most NAP XPS systems allow the dosing of gases and gas mixtures and heating in order to perform in situ characterisation of heterogeneous catalytic reactions in different gas atmospheres. These systems can be easily adapted for electrochemical systems based on solid electrolytes.²⁸ However, the measurement of electrocatalytic systems based on liquid electrolytes requires the use of especially designed electrochemical cells.²⁹ There are two main types of approaches in the literature for the operando analysis of electrochemical reactions using liquid electrolytes by XPS. One requires the use of graphene windows that separate the liquid and UHV environments and through which, X-rays and electrons can pass through, allowing XPS measurements. Despite being first established in 2011,³⁰ these cells have led to very few studies, mainly due to the fragility of these 2D windows under operation in UHV chambers.^{31,32} The other approaches are windowless and require the delicate fine control of the continuous evaporation of the electrolyte in a UHV chamber.³³ which requires a fine balance of the operating conditions but has given rise to a more significant amount of contributions in both photoelectrochemical and electrocatalytic studies.²⁸

The performance of in situ measurements of photoelectrocatalytic systems adds an additional complication to the photoelectrochemical approach, as the sample must be properly illuminated. For the illumination, there are two possibilities, direct illumination by the set-up of an UV light within the UHV chamber using electrical feedthroughs or illumination through a window outside the XPS instrument. This port can be one facing directly at the sample in the UHV chamber or the port at the photoelectron analyser that allows directing light down the axis of the spectrometer. Common UHV windows made of borosilicate allow the transmittance of most ultraviolet light down to 300/350 nm wavelengths. If the experiments require the use of more energetic UV radiation, UV Windows made of sapphire, fused silica or even MgF₂ could be used with absorbance cut-offs that go down to 200nm wavelengths.³⁴ Specifically, there are different available commercial windows for performing these experiments: i) OROFLOAT borosilicate windows (Edmund Optics) have a transmission of more than 90% between 350 and 2500 nm; (ii) TECHSPEC sapphire windows (Edmund Optics) are useful in the transmission range 330–5500 nm; (iii) fused quartz windows (Knight Optical) are a more suitable material choice for UV light; (iv) any other window materials of the correct size which satisfies the needs of the users' experiment. User-supplied light sources can also be attached to the main chamber with minimal effort. Moreover, illumination from outside the UHV chamber also requires a lens focusing system at the sample distance or a large collimated source. This later set up adds the advantage of illuminating the sample only in the area of investigation. One has also to consider that depending on the energy range and intensity, illumination can also lead to sample heating. Therefore, a way to measure temperature on the sample or a previous calibration is advisable, in order to discount the temperature effect on the studied reaction.



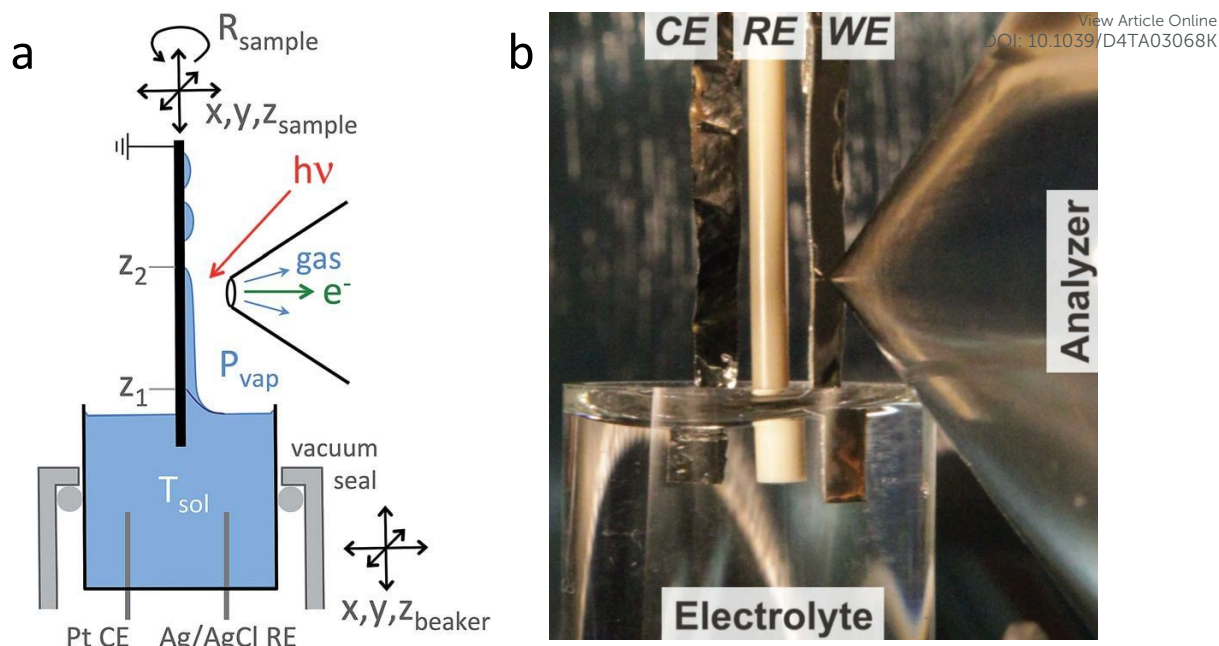
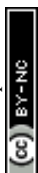


Figure 1. a) Scheme of the Dip and Pull method for the preparation of thin liquid films on electrodes. b) Picture of the Dip and Pull set up at ALS. Copyright 2015. Reproduced with permission from Springer Nature. Adapted from reference ³³.

The *dip and pull* method, shown in **Figure 1**, is probably the easier type of electrochemical cell to adapt to photoelectrocatalytic measurements and the method of choice in most photoelectrochemical studies in the literature. In particular, the SpAnTeX endstation at the BEIChem-PGM endstation in BESSY II offers a *Dip and Pull* module specially designed for photoelectrocatalytic measurements.³⁵ Other end stations that offer the Dip and Pull method such as HIPPIE in Max IV, PHOENIX I at Swiss Light Source (SLS) or BL 9.3.1 at Advanced Light Source (ALS) could be easily adapted to photoelectrochemical experiments building up on their experience on photocatalytic reaction monitoring. Most of the Dip and Pull measurements are done in Synchrotron stations using tender X-rays (2-7 keV) as photoelectrons of high energy have more probabilities of going through the thin electrolyte meniscus that is created in order to allow electrons coming from the “buried” electrode to reach the analyser. As the probabilities of emitting photoelectrons of most probed orbitals go down significantly with excitation energy, 3 KeV seems to be the sweet spot for maximising signal intensity of the buried electrodes.³³ Nevertheless, experiments can be done at lower excitation energies, e.g. the HIPPIE beamline works at a maximum excitation energy of 2.2 keV and laboratory systems have also been used successfully to probe buried liquid interfaces in battery systems for example.³⁶ Moreover, the use of Silver (Ag-K α , 2.9 KeV) or Chromium (Cr-K α , 5.4 KeV) sources can also increase the probing depth of laboratory systems, complementing the experiments done with the typical Al-K α sources (1.5 KeV). The added advantage of using synchrotron systems is that the variable energy can also be used to differentiate between surface and subsurface signals from the electrode, as surface signals will be more enhanced at lower excitation energies. Moreover, synchrotron light allows to perform Auger yield NEXAFS, which provides complementary surface sensitive XAS information.

The possibility of performing in situ and operando XPS experiments using these setups has advanced the knowledge of the surface chemistry and electronic properties of photoabsorber materials that are used as photoelectrodes under realistic conditions of illumination and chemical environment. These studies can be divided into three categories based on how close to the actual operando conditions they are performed: (1) XPS under UV-visible illumination in UHV, (2) XPS under UV-visible illumination in NAP conditions, and (3) operando XPS. In the following subsections, we discuss these contributions in terms of the information obtained and the required facilities.



(1) *XPS of photoelectrodes under in situ illumination in UHV*: Studying photoelectrodes using XPS under in situ illumination has helped in the elucidation of intrinsic light-induced changes of surface chemical state, electronic structure and the behaviour of charge carriers.³⁷ XPS has been particularly useful to study the surface photovoltage (SPV) of semiconductors used as photoelectrodes. The phenomenon of SPV arises from the distribution of photogenerated electron-hole pairs within the space charge region (SCR) of the semiconductor, providing a critical method for evaluating the minority carrier diffusion length in semiconductors. Since XPS is sensitive to changes in surface potential, this technique has been widely employed to carry out detailed SPV studies on organic and inorganic semiconducting materials.^{38–40} This method, accessible through both synchrotron radiation and laboratory setups, simplifies the analysis of SPV by following shifts in photoemission peak positions under direct illumination. Early XPS-based SPV analyses were done for Si (111) by Demuth *et al.*^{38,41} finding that at temperatures below ~ 50 K surface recombination becomes ineffective thereby allowing flat-band conditions to be achieved with mild UV irradiation. Temperature-dependent photoemission thereby provides a simple and direct method for determining band bending and barrier heights. XPS-based SPV measurement may be subject to artifacts due to charging, which motivated Suzer *et al.* to study dynamical XPS measurements for probing photoinduced voltage changes.⁴² In order to distinguish between surface photovoltage (SPV), and charging, they proposed to perform dynamical measurements while subjecting the sample to square wave pulses of ± 10.00 V amplitude at frequencies from and 10^{-3} to 10^5 Hz. Also, Ekiz *et al.* discuss a method for characterizing photovoltaic and photoconductivity effects on nanostructured surfaces through light-induced changes in XPS.⁴³ Their technique merges the chemical specificity of XPS with its sensitivity to the SPV, allowing the characterisation of photoconductivity under both static and dynamic optical excitation. They introduce a theoretical model that quantitatively describes the features of observed spectra, demonstrating its applicability on various sample systems, including solar cells and semiconductor nanoparticles. This approach underscores the potential of XPS to differentiate between surface voltage shifts due to photovoltaic or photoconductive effects, offering a pathway to predict dynamical behavior under modulated illumination.

XPS-based SPV analysis offers an advantage over traditional Kelvin probe techniques by delivering extra chemical insights, making it a valuable tool for semiconductor research.^{44,45} A study by Oropeza *et al.*⁴⁴ examines the effect of surface photoreduction on the PEC properties of CuBi_2O_4 , using XPS to study the surface reduction and its effect on the surface energetics and SPV. They report that irradiation of CuBi_2O_4 with photons ($h\nu \geq 2.7$ eV) in an inert atmosphere leads to the formation of reduced Cu on the surface. This reduction induces a downward band bending of 0.35 eV, enhancing the charge carrier transport properties, as evidenced by an increase in the characteristic SPV from 0.07 to 0.27 V. This remarkable finding constitutes the first experimental evidence of the beneficial effects of surface photoreduction on PEC properties of CuBi_2O_4 . Similarly, in their investigation of the InP interphase energetic optimization, Gao *et al.*⁴⁵ employed XPS in core and valence band regions to monitor the modulation of the surface energetics brought by the introduction of a buried p–n⁺ junction which favourably shifts the valence band edge to promote minority charge carrier's diffusion. They confirmed the enhancement of the charge carrier's diffusion length by XPS-based SPV measurements, using the binding energy shift of the P 2p and In 3d peaks upon in situ illumination, see **Figure 2a**. They observed an increase of the SPV from 0.1V to 0.35V, and confirmed that the SPV of the n⁺/p-InP depends on light intensity with the relationship:

$$SPV = \frac{kT}{e} \ln(1 - bI_0) \quad (\text{Equation 2})$$

where I_0 is the incident light flux, b is a characteristic parameter of the semiconductor, which is constant under moderate band bending ($BB \gg kT$) and e is the elementary charge. Equation 2 predicts that SPV varies linearly with $\log(I_0)$ under moderate illumination, which was confirmed for their n⁺/p-InP samples and should be taken as a general check measurement in studies of SPV.



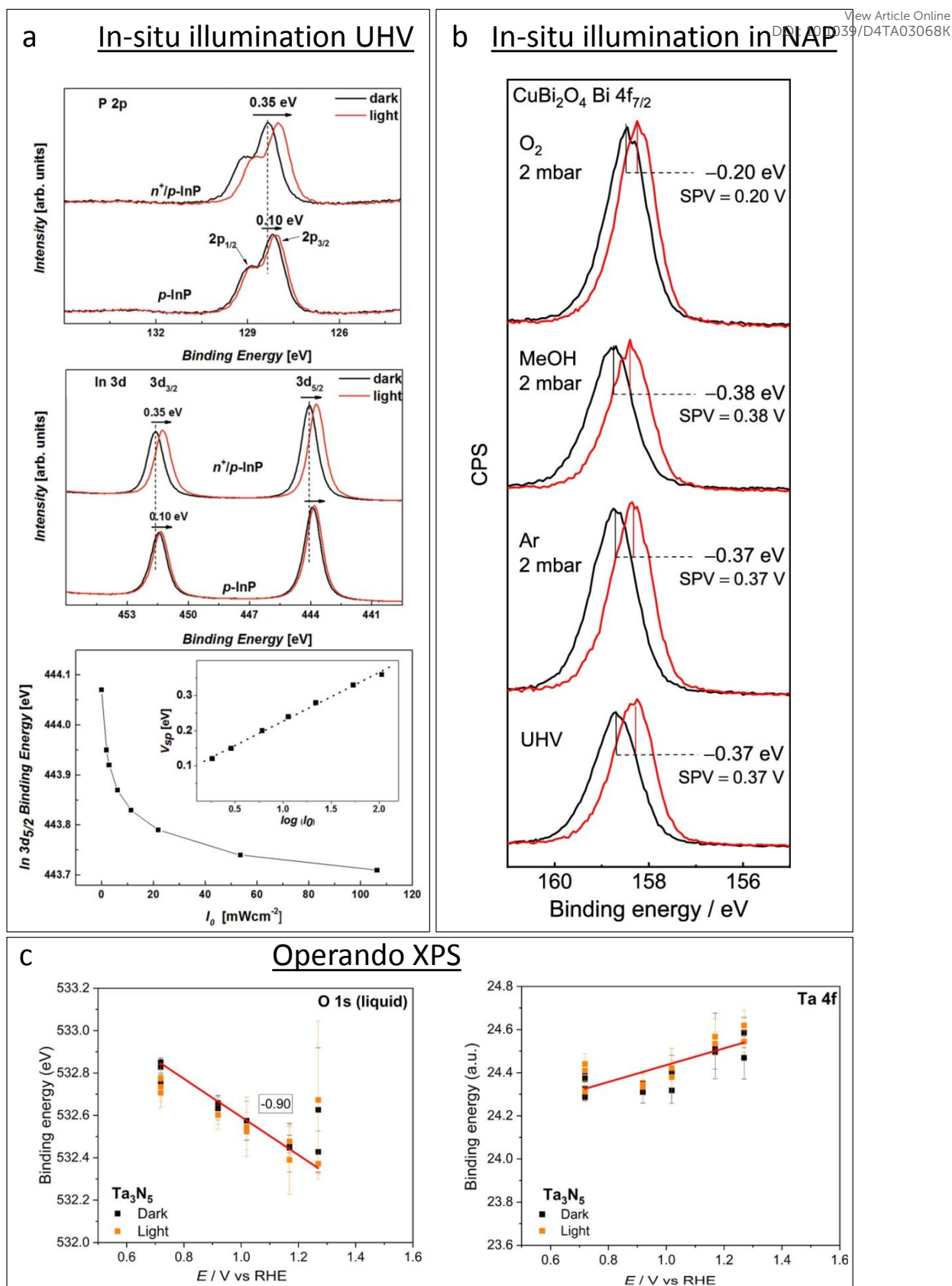
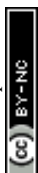


Figure 2. Progress of in situ XPS studies from illumination in UHV operando XPS. a) From top to bottom: XPS in the P 2p and In 3d regions of p-InP and $n^+/p\text{-InP}$ in dark (black solid lines) and under illumination with an LED $h\nu_2 = 1.98$ eV (red solid lines), and In 3d_{5/2} binding energy shift for $n^+/p\text{-InP}$ as a function of the incident light flux (I_0). The inset shows the surface photovoltage

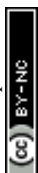


(SPV) as a function of $\log(I_0)$. Copyright 2016. Reproduced with permission from Wiley. Adapted from reference.⁴⁵ b) XPS of a CuBi_2O_4 photoelectrode in the Bi $4f_{7/2}$ region, showing the XPS-based SPV measurement (spectra in black lines is in dark, spectra in red lines are in light with a 405 nm laser) in UHV and in the presence of Ar, methanol vapour, and oxygen. Copyright 2019. Reproduced with permission from American Chemical Society. Adapted from reference.⁴⁴ c) Binding energy of the Ta $4f_{7/2}$ component and liquid contribution in the O 1s spectra during operando AP-XPS measurements for Ta_3N_5 photoelectrode. Copyright 2023. Reproduced with permission from American Chemical Society. Adapted from reference.⁴⁶

(2) XPS of photoelectrodes under in situ illumination in NAP conditions: Then introduction of near-ambient pressure XPS has allowed the study of adsorption processes of water, CO_2 and other reactant gases on the surface of photoelectrodes, providing key insights into dissociation mechanisms which are key to understanding the overall artificial photosynthesis process. Although most of NAP XPS studies has been carried out in dark conditions, important aspects on the catalytic performance of the photoelectrochemical processes have been addressed. Research has been focused on two major topics: (i) molecular dissociation (water or CO_2) process on semiconductor photoabsorbers or the co-catalyst/semiconductor interface, and (ii) effect of probe molecules on the chemistry and electronic structure of semiconductor photoabsorbers.

Zhang and Ptasinska conducted a series of studies of the water dissociation on model Ga-based photoelectrodes exposing GaP (111) and GaN (0001) to water vapour up to 5 mbar and temperatures from 298 to 773K.^{47–49} They found enhanced surface hydroxylation and oxidation of gallium on GaP⁴⁸ as water pressure increased, with significant changes at temperatures above 673 K, leading to the formation of non-stoichiometric gallium hydroxide and oxidation of phosphorus atoms. For GaN (0001), water dissociation at room temperature led to the formation of hydroxyl groups and a decrease in intrinsic upward band bending, enhancing its photocathodic properties. However, increased temperatures strengthened the chemical interaction, forming more oxides and hydroxides, which increased upward band bending and negatively affected GaN's photocathodic properties.⁴⁹ Additionally, their studies on Pt/GaN (0001) revealed that water dissociates readily at the Pt/GaN interface, unlike on pure Pt surfaces.⁴⁷ This interface showed significant upward band bending, promoting the accumulation of photogenerated holes and enhancing oxidative dissociative adsorption of water, leading to the formation of hydroxyl groups and surface oxides bound to Pt, which are active catalytic sites for the hydrogen evolution reaction (HER).^{50,51} Jackman et al. studied water adsorption on anatase TiO_2 (101) using NAP-XPS,⁵² finding that water adsorbs in a mixed molecular and dissociated state on stoichiometric surfaces. Sputtered surfaces exhibited similar characteristics, indicating subsurface vacancies do not alter water interaction. The significant amount of adsorbed molecular water on TiO_2 suggests limited chemical interaction compared to more reactive surfaces like GaP, InP, and GaN, where a more dissociative process occurs.^{47–49}

The chemical and electronic interactions between photoelectrodes and adsorbates have also been an important topic of research using NAP-XPS. An early study by Salmeron *et al.* showed that the adsorption of O_2 (with a O_2 pressure of 1 Torr) on the TiO_2 surface leads to a shift in the Ti 2p and O 1s core level binding energies towards lower binding energy (by 0.4 eV), suggesting that the adsorption process involves the transfer of electrons from TiO_2 to O_2 , resulting in such an upwards band bending.⁵³ Mansfeldova *et al.* studied more specifically the effect of the environment on the surface work function (WF) of TiO_2 photoelectrode.⁵⁴ Their NAP-XPS data showed a significant decrease of the surface WF of TiO_2 (by about 1 eV) upon water vapour exposure at 0.5 mbar, indicating a strong upwards shift of the conduction band, which correlates with its photoanode behaviour in liquid electrolytes. The effect of the environment on the SPV of photoelectrodes is also accessible by NAP-XPS. Using in situ illumination under NAP-XPS conditions, Oropeza *et al.* studied the effect of key electron donors and acceptors in the SPV of CuBi_2O_4 photocathodes, finding that whereas electron acceptors like O_2 causes the reduction of the SPV, electron donors like methanol favours and increased SPV, see **Figure 2b**. Since XPS can simultaneously monitor the surface chemistry of the sample, the authors could correlate the



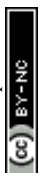
diminution of the SPV with the re-oxidation of reduced Cu surface states, which led to the conclusion that those reduced Cu surface states are key achieving good photoelectrochemical properties of CuBi_2O_4 .

(3) *XPS of photoelectrodes under operando conditions*: Research on operando XPS of photoelectrodes has been particularly important in the context of artificial photosynthesis because it has been critical for understanding the dynamic chemistry occurring at the surface/interface of photoelectrodes during the PEC processes like water splitting and CO_2 reduction. Recent work has been focused on two main topics: i) studies of the energetics at the photoelectrode/electrolyte interface and ii) the generation of reaction intermediates at the surface of the photoelectrodes.

For instance, the research conducted by Lichterman *et al.*⁵⁵ provides a fundamental perspective on the operational characteristics of semiconductor/liquid junctions, in more realistic conditions than previous gas-phase-based NAP-XPS studies. Using the dip-and-pull technique described above, they demonstrated how operando ambient-pressure XPS can be used to directly characterize the energetics of semiconductor/liquid junctions, focusing on the dynamics of semiconductor accumulation, depletion, and Fermi level pinning by defect states. In their work, the observed shifts in binding energy with respect to the applied potential directly revealed the ohmic junction nature at the liquid interface on metallized samples, and the rectifying junction behaviour at the liquid interface with semiconductor photoabsorbers. Furthermore, their findings on the behavior of metallized semiconductor/film contacts under varying conditions contribute to clarifying the mechanisms of charge transfers in photoelectrochemical cells. They highlighted that all parameters that can be directly observed by operando XPS are crucial variables for the design and operation of semiconductor–liquid junction devices for the PEC water splitting and CO_2 reduction. In a similar dip-and-pull approach, and introducing in situ illumination, Dahl *et al.*,⁴⁶ studied the semiconductor/electrolyte interfacial energetics of a tantalum nitride (Ta_3N_5) film, with and without a nickel oxide (NiO_x) protective layer, were explored under applied bias in both dark and illumination conditions. **Figure 2c** shows the variation of the binding energy of Ta 4f from the sample and O 1s from the electrolyte as a function of the applied potential in dark conditions and under illumination. As expected, the O 1s peak position of the electrolyte shifts constantly with the applied potential; however, the Ta 4f peak position is independent of the applied potential both in dark conditions and under illumination. The authors interpreted it as a direct evidence of Fermi-level pinning in Ta_3N_5 at the Ta_3N_5 /electrolyte that effectively prevents band bending toward the Ta_3N_5 /electrolyte. This means that there is no potential drop in the SCR that can promote the separation of photogenerated holes and electrons, which leads to poor PEC performance of bare Ta_3N_5 . However, this work highlighted that surface passivation through the NiO_x layer could significantly reduce the recombination of photogenerated carriers, leading to increased surface band bending of Ta_3N_5 , and better PEC performances. They did not find any clear oxidation of the Ta_3N_5 , which correlated with the significantly larger photocurrents and stability of the NiO_x -coated materials. These types of operando XPS studies of the interface energetics provide key elements for the development of efficient and durable systems for water photoelectrolysis.

On the other hand, Favaro *et al.*⁵⁶ focused on bismuth vanadate (BiVO_4) electrodes in contact with an aqueous potassium phosphate (KPi) solution at open circuit potential under both dark and light conditions, observing that visible light illumination triggers the formation of bismuth phosphate on the surface of the BiVO_4 photoanode. They suggested that this chemical modification leads to surface passivation that impacts the distribution of ions within the thin electrolyte layer due to the repulsive interaction between the negatively charged surface under illumination and the phosphate ions in solution, which is observed as an increase in their photoelectron signals. Interestingly, they found that such light-induced changes at the BiVO_4 /KPi electrolyte interface are fully reversible upon returning to dark conditions. Their findings emphasize the significance of in situ methods to understand the impact of operational conditions on the performance of photoanodes.

These studies collectively illustrate the critical role of operando XPS techniques in advancing the field of photoelectrochemistry. By providing direct insights into the interfacial energetics and chemical states of photoelectrodes under operational conditions, researchers can better



understand and mitigate the challenges of photoelectrode degradation and inefficiency. Such advancements not only contribute to the fundamental knowledge of photoelectrochemical mechanisms but also pave the way for the development of more robust and efficient solar fuel production systems. The current availability of several synchrotron stations capable of performing operando photoelectrochemical studies with robust setups is the optimal platform to establish routine analysis of photoelectrochemical systems and quickly advance on these ambitious scientific objectives. The possibility of adapting the new AP XPS instruments such as *Polaris* in *DESY*⁵⁷ and *3Sbar* in *ALBA* (in construction phase), capable of performing XPS analysis at pressures of a few bars using tender X-ray sources could provide more stable environments to establish routine analysis of photoelectrochemical systems. Moreover, in combination with the recently developed stroboscopic technique in *MAX IV*,⁵⁸ it could lead to the performance of time resolve experiments in the microsecond range and allow the investigation of charge transfer dynamics and reaction mechanics in the microsecond time scale, exploiting this way the maximum potential of operando XPS experiments of photoelectrochemical systems.

In situ and operando X-ray absorption spectroscopy (XAS)

Synchrotron **X-ray absorption spectroscopy (XAS)** has been proven as a highly effective tool for materials characterisation. Specifically, in XAS, the absorption of energy, carried by a photon, promotes an electron from a core-level to an empty orbital, monitoring the unoccupied electronic states, and hence, probing the local chemical environment (4–5 Å), providing information regarding the specific chemical bonds, oxidation states, spin states, site symmetry, and coordination environment.⁵⁹ In a typical XAS experiment, the monochromatic incident X-ray beam scans the vicinity of the absorption edge of an element, usually requiring a synchrotron X-ray source (tunable energy, highly monochromatic and high intensity), even there are currently available lab-scale XAS equipments.⁶⁰ Specifically, there are two ways of performing XAS experiments: i) X-ray absorption spectra are recorded by measuring the ratio of the transmitted and incident X-ray beam intensity, known as transmission mode, or ii) the yield of secondary particles, fluorescence photons or emitted electrons, as a function of photon energy varied across the absorption edge of an element (total fluorescence yield, TFY, and total electron yield, TEY, respectively).⁶¹ In transmission XAS, X-rays undergo attenuation while passing through the matter and intensity drops exponentially with the travelled distance, which is given by:

$$I_t = I_0 e^{-\mu x} \quad (\text{Equation 3})$$

where I_t and I_0 are the transmitted and the initial X-ray beam intensities, respectively, μ is the absorption coefficient and x is the thickness of the sample.⁶² On the other hand, in TFY, also known as X-ray fluorescence (XRF), the emitted fluorescence X-rays after the interaction of the incident beam with the matter, are measured as a function of the incident X-ray photon energy (E), and the absorption coefficient is:

$$\mu(E) \propto \frac{I_{fluo}}{I_0} \quad (\text{Equation 4})$$

Finally, in analogy to XRF, the TEY measurements are based on the proportionality between the X-ray absorption coefficient and the emitted Auger electrons during the core decay:

$$\mu(E) \propto \frac{I_{TEY}}{I_0} \quad (\text{Equation 5})$$

Specifically, this experimental mode consists of measuring the total necessary applied current to compensate the photoinduced drain current in the sample.⁶³ Hence, every XAS spectrum represents the X-ray absorption coefficient, $\mu(E)$, of a system as a function of the incident X-ray photon energy. Besides, different regions can be distinguished in a typical XAS spectrum: i) the pre-edge region, determined by the transition to the lowest unoccupied states, ii) the X-ray absorption near-edge structure (XANES), also called NEXAFS from near-edge X-ray absorption fine structure, dominated by core transitions to quasi bound states (multiple scattering resonances) for photoelectrons with kinetic energy in the range from 10 to 150 eV above the



chemical potential and iii) the extended X-ray absorption fine structure (EXAFS) region, associated with a weak scattering cross-section with neighbour atoms.

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X-ray Absorption Spectroscopy (XAS) can be categorized into two measurement methods: hard-XAS, which employs high-energy X-rays to study bulk properties and heavy elements through core-level transitions, and soft-XAS, which utilizes lower-energy X-rays, offering surface sensitivity and probing valence band transitions for studying surface phenomena and organic material. Specifically, hard-XAS experiments are usually conducted in the transmission mode, which provides bulk-sensitive analysis and does not suffer from self-absorption effects.⁵⁹ However, due to the attenuation length of less than one μm , soft-XAS is usually measured through fluorescence and electron yield modes, having different probe depths.⁶⁴ TFY technique shows an estimated probing depth of a few hundred nanometres, while TEY measurements is more surface-sensitive, with a probing depth of approximately a few nanometers.⁶⁵ Additionally, depending on the nature of the sample, different detection modes can work better. For example, sometimes, transmission XAS experiments can suffer from sample damage, caused by inhomogeneous concentrations or densities in the sample. To avoid this problem, transmission experiments require careful sample preparation and relatively concentrated samples, ideally in a light matrix. On the contrary, fluorescence yield mode is more suitable for diluted samples or systems with heavy environments.⁶⁶

In situ XAS measurements have been demonstrated to be essential for advancing our fundamental understanding of photoelectrocatalysis and accelerating the development of efficient catalytic materials for sustainable energy technologies. This spectroscopy allows monitoring changes in the chemical and electronic states of the catalyst in real time as they undergo photoelectrocatalytic reactions. This real-time monitoring provides information on reaction mechanisms, intermediate species, and reaction kinetics that are difficult to obtain using ex-situ techniques. Besides, XANES and EXAFS, can provide information about the local coordination environment and electronic structure of catalytic materials. This information helps in identifying the active sites responsible for catalytic activity, facilitating the design of more efficient catalysts.

In general, the experimental requirements for performing in situ and operando (hard)-XAS measurements are easier than those for XPS mainly due to the vacuum requirements, which make the development of in situ electrochemical cells much more complicated. Due to this reason, a considerable higher number of studies have reported in situ and operando (hard)-XAS measurements in electrodes for solar fuels generation compared to in situ and operando XPS.^{67,68} However, still limited studies have reported in situ XAS on photoelectrodes, due to X-ray absorption measurements with photoelectrochemical cells require complex engineering to ensure compatibility and functionality of both systems simultaneously, compared to the in situ study of electrocatalysts.^{69,70} This involves designing specialized cells that can accommodate both X-ray transparency and photoelectrochemical functionality. Furthermore, most relevant chemical and electronic changes in photoelectrochemistry take place on the surface of the electrode, and not in the bulk, further complicating the collection of reliable data that provides true and reliable information. Thus, due to the bulk nature of the XAS technique compared with the shallow nature of the changes under PEC conditions in the photoelectrodes, there are mainly two ways to perform in situ experiments: i) to fabricate really thin films of the photoelectrode under study, trying to maximize the surface-to-volume ratio or ii) to investigate the induced changes in a co-catalyst by the effect of the photogenerated carriers in the absorbing material. Both type of studies has been considered in the present review.

Among the characterisation of photoelectrodes for solar fuels generation by in situ and operando XAS, much effort has been developed in the oxidation reactions, mainly in the oxygen evolution reaction (OER) and hence in characterising n-type semiconductors. However, there are already some recent works that report in situ XAS in p-type photocathodes for the hydrogen evolution reaction (HER), and the challenging CO_2 reduction reaction (CO2RR), as well as N_2 fixation (N2RR) to ammonia (NH_3).

Regarding photocathodes analysis, in 2020 Chen *et al.*⁷¹ investigated through in situ XAS bimetallic Ni based p-Si micropillar photocathodes for solar-driven HER. The authors incorporated



Ni/Mo, Ni/Co, and Ni/Fe to the silicon electrodes and analysed the effect of the incorporation of Mo, Co and Fe on the photoelectrochemical properties, finding out that the most active is the Ni/Mo combination (**Figure 3a**). To investigate the oxidation state and local atomic environment under photoelectrochemical conditions, they performed in situ XANES measurements in the liquid medium. A custom reaction cell designed for this in situ measurement was used, but the authors do not provide details or schematics about its design. After introducing NiMo/p-Si MPAs in the alkaline electrolyte, no noticeable change was observed in the Ni K-edge spectra (**Figure 3b**). However, under cathodic conditions, above 0 V, they found out that Ni XANES edge shifted to a higher energy indicating an increase in the Ni valence state (**Figure 3c**). Interestingly, when the same potential is applied to the NiCo/p-Si and NiFe/p-Si MPAs, negative changes in the absorption energy can be observed, indicating that the reduction of Ni was caused by the cathodic potential during HER (**Figure 3d and 3e**).

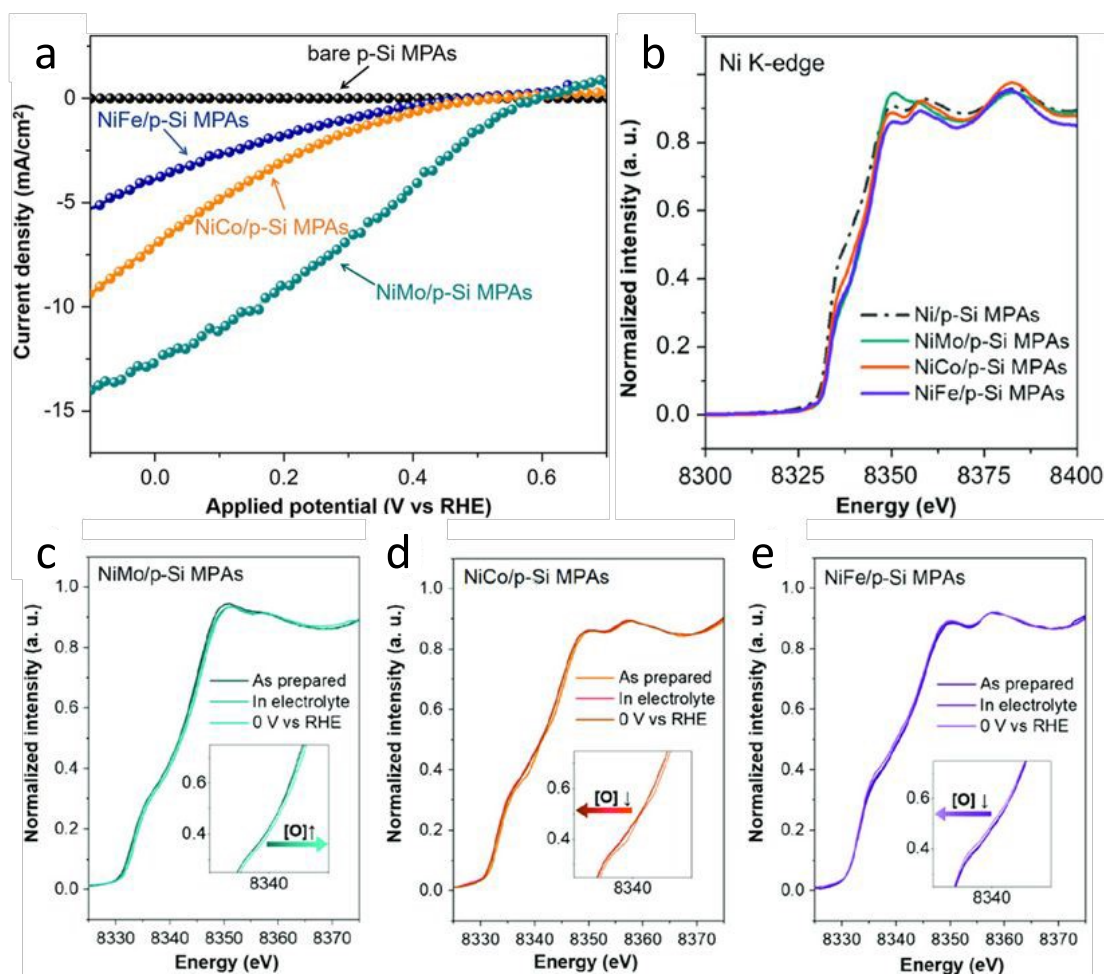


Figure 3. a) Photoelectrochemical characterisation of the Ni-based alloys/p-Si MPAs photocathodes in 1 M KOH under illumination. b) Normalized Ni K-edge XANES spectra of the p-Si MPAs/alloys photocathodes. c-d) Ni K-edge of in situ X-ray absorption spectroscopy measurements of the Ni-based alloys/p-Si MPAs during the photoelectrochemical hydrogen production reaction. Copyright 2020. Reproduced with permission from Wiley. Adapted from reference ⁷¹.

Furthermore, by in situ EXAFS, the authors analysed the local atomic environment. Once the HER starts, the peak intensity of the Ni–O scattering path at 1.4 Å in NiMo/p-Si sample significantly increased. This effect it is not observed in the rest of the electrodes. The scattering paths of Ni–Ni were mostly unchanged in all cases. Besides, the intensity of the Ni–O peak in NiCo/p-Si MPAs decreased, indicating that the nickel oxides were changed into the metallic state during the reaction. These in situ-formed Ni-species might be considered as the active intermediates for the high activity of NiMo/p-Si MPAs. Other researchers correlated the HER



activity with the NiO introduction on the Ni surface before.⁷² According to these results, the authors suggest that the growth of the high-valence Ni (hydr)oxides during HER could favour the photoelectrochemical properties in alloy-based photocathodes.

On the other hand, Wang et al. investigated silicon-based photocathodes for photoelectrochemical (PEC) nitrogen reduction to ammonia, aiming to emulate natural reaction processes involving multiple protonation and electron transfer steps. They designed a two-cell electrochemical system to enhance PEC nitrogen reduction reaction (NRR) efficiency. This system included a stainless-steel mesh cathode modified with encapsulated Fe₃C nanoparticles and amorphous carbon. The two cells, physically and electrically separated, allowed the N₂ feed gas to pass through the porous electrode of the first cell (C2) before reaching the working photocathode in the second cell (C1). In 2021, they reported significant improvements using this setup on a Au/SiO₂/Si photocathode, achieving an NH₃ yield rate of 22.0 μg·cm⁻²·h⁻¹ and a faradic efficiency (FE) of 23.7% at -0.2 V vs. RHE. Without the stainless-steel-based cathode, the yield rate and FE were markedly lower, at 6.6 μg·cm⁻²·h⁻¹ and 7.2%, respectively.⁷³ First, XPS was used to track changes in the oxidation states of the Au/SiO₂/Si photocathode before and after illumination. The survey scans showed similar characteristic peaks for Au 4f, Si 2p, and O 1s under both conditions. High-resolution XPS spectra revealed a decrease in the Au¹⁺ to Au⁰ ratio under illumination, and a higher Si⁴⁺ to Si⁰ ratio compared to dark conditions.

Additionally, operando XAS was used to explore the chemical environment under working conditions. The XANES spectra at the Au L₃ edge (**Figure 4a**) indicated the oxidation state of Au was related to the intensity of the white line (WL) peak at ~11,920 eV. The Au L₃ edge XANES difference spectra (**Figure 4b**) revealed a WL peak at ~11,920 eV, which intensified with increasing Au oxidation state. This peak intensity varied with different photoelectrode potentials and the presence of the modified stainless-steel cathode, correlating with changes in PEC NRR efficiency. The system with the modified stainless-steel cathode maintained at -0.2 V vs. RHE exhibited the highest oxidation state of Au, suggesting an increase in vacancies at the Au 5d valence level. Besides, Fourier transform k₃-weighted EXAFS spectra (**Figure 4c**) under reaction conditions showed contributions from Au-Au bonds and a new peak at ~1.3 Å, attributed to Au-N coordination, which was not present in less efficient PEC NRR reactions. The single-cell system, lacking the modified stainless-steel cathode, showed a lower Au oxidation state and reduced HER performance. The study suggested that the modified stainless-steel cathode first activates N₂ molecules at the optimal potential, forming active nitrogen species that adsorb onto the surface of Au NPs, enhancing covalent bonding and improving PEC NRR performance on the Au/SiO₂/Si photocathode.

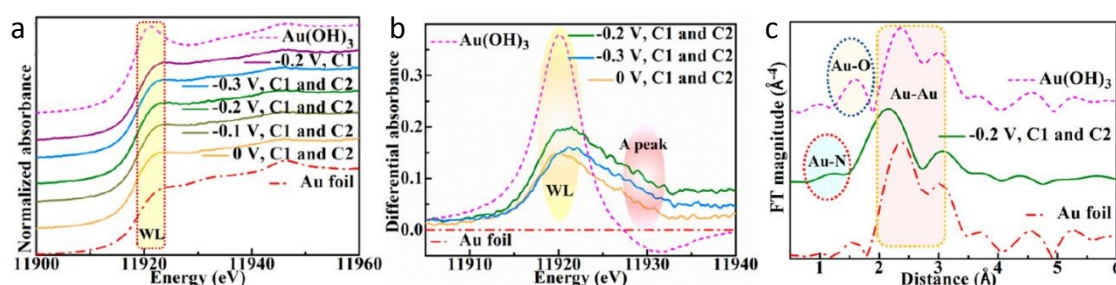


Figure 4. Operando XAS of Au/SiO₂/Si photocathode in the NRR PEC process. a) Au L₃-edge XANES spectra with and without the excitation of a modified stainless-steel cathode at each given photoelectrode potential during PEC NRR. b) Comparison between the Au L₃-edge XANES difference spectra of the photocathodes with a modified stainless-steel cathode at different photoelectrode potentials. The Au foil as the baseline was chosen to obtain the various difference spectra. (c) Fourier transform of the experimental EXAFS spectra of the photocathode with modified stainless-steel cathode at -0.2 V vs RHE, Au foil, and Au (OH)₃. Copyright 2021. Reproduced with permission from American Chemical Society. Adapted from reference ⁷³.



In 2023, the same group investigated the role of the synergy between electron localization and alloying Au, Co and Pd (ACP) in determining the photoelectrochemical conversion of nitrogen to ammonia in the multilayered AuCoPd-CoO_x/SiO₂/Si photocathodes.⁷⁵ On the one hand, XPS analysis showed that while Au and Pd maintained their peak positions under illumination, their average oxidation levels decreased. The peak positions associated with Co were unaffected, but the Co^{x+}/Si⁴⁺ to Co⁰/Si⁰ ratio increased, differing from Au and Pd trends. These tests were pseudo in-situ since no potential was applied. On the other hand, as in their previous work, The XANES spectra of the Au L₃ edge indicated that ACP samples had spectra similar to Au foil and different from Au(OH)₃. The ACP sample performed better than the Au-only sample. At the highest yield potential (-0.3 V vs. RHE), the ACP sample showed the highest Au oxidation state, indicating maximum formation of hybridized chemical bonds during PEC NRR. EXAFS spectra revealed consistent Au–Au bonds across all bias potentials. However, the intensity of the ~2.0 Å peak, corresponding to Au–Pd bonds, increased as the applied potential decreased, peaking at -0.3 V vs. RHE. This increase reflected the enhanced formation of the AuPd alloy due to greater reduction of Ag⁺/Pd sites under illumination.

In 2021, Sawangphruk *et al.* analysed a cobalt-based photocathode by in situ XAS.⁷⁶ Specifically, the authors investigated a cobalt hydroxide nanosheets as photoelectrode and the effect of silver particles on the optical properties and conductivity. This material is usually used as electrode material for various energy storage and conversion applications,⁷⁷ and recently has been reported as photoactive material.^{78,79} Photon energy can cause cobalt hydroxides to generate charge carriers. This process can generate additional photocurrent, leading to higher performance when Co(OH)₂ is used as the electrode. However, their intrinsic electronic and electrical conductivities are very poor and the introduction of AgNPs can be a solution to overcome these problems. In this work the authors claim that under dark conditions, AgNP-doped α-Co(OH)₂ shows higher capacitance than undoped β-Co(OH)₂, and the authors attributed this to its higher electrical conductivity and therefore a faster charge transfer. However, under illumination, while an improvement in the charge storage performance of undoped Co(OH)₂ is observed, the opposite phenomenon occurs with the addition of Ag. The excessively narrow band gap and the competitive surface of the resonance process of the plasmon occur in parallel on the AgNP surfaces, leading to less charge capacity performance. In this sense, the change in the oxidation state of Co during the electrochemical process is crucial evidence to investigate the charging mechanism of electrode materials. To gain a deep understanding of the local structural change in Co and its pseudocapacitive behavior when the photoelectrodes are charged and discharged, the authors carried out in situ XAS experiments by analyzing the XANES spectra of the Co-K edge. To do that, the authors used an electrochemical cell developed for a previous work.⁸⁰ Then, they analysed the Co-K edge XANES spectra of α-Co(OH)₂ Ag doped with the AgNP photoactive electrode under light and dark conditions at different potentials. The average oxidation state of Co in Ag–Co(OH)₂ is +2.58 at an energy edge of 7719.89 eV. This result refers to mixed valences of Co²⁺/Co³⁺. The initial oxidation state of Co in α-Co(OH)₂ Ag doped in 1 M NaOH is found to be +2.46 eV. Under -0.4 to 0.1 V versus SCE bias potential, the edge positions change significantly to higher energy for both conditions, indicating an increasing oxidation state of Co during the charging process. Under dark conditions with the fully charged state of Ag-doped α-Co(OH)₂, the oxidation state of Co increases to +3.08, while under light illumination was +3.14. The higher oxidation number of the Ag-Co(OH)₂ electrode under illumination was attributed to the photovoltaic effect, which generates the photoelectrons. However, the oxidation state of Co in Ag-Co(OH)₂ under dark conditions is +3.0, while under light irradiation it decreases to +2.92. The conclusion they reached was that when the Ag-doped α-Co(OH)₂ photoelectrode is discharged, the photoelectrons can recombine with the holes generated in the CB, while the photoexcited electrons under dark conditions were transferred to the external system.

In the case of photoanode materials, hematite, Fe₂O₃, and bismuth vanadate, BiVO₄, stand out as the most studied candidates for PEC oxidation reactions and are the most studied semiconductors with in situ and operando synchrotron XAS. Specifically, hematite (α-Fe₂O₃) photoanodes have attracted an increasing attention as photoanode material for photoelectrochemical applications thanks to its adequate bandgap, abundance and stability.⁸¹ Although progress has been made to achieve highly efficient hematite photoelectrodes for the



OER,⁸² understanding the nature of recombination and charge transfer processes using X-ray based techniques is essential.⁸³ In this sense, XAS measurements have been used to correlate the presence of defects in hematite photoanodes with changes in their photoelectrochemical behaviour. *Ex-situ* FT-EXAFS of the Fe K-edge revealed that the peaks intensity corresponding to whether the Fe–O or Fe–Fe bonds can be reduced when oxygen vacancies are present.⁸⁴ Mesa *et al.*, have recently reported in situ XAS showing that the Fe–Fe bond signal, in the Fe K-edge FT-EXAFS data, reduces when applying a positive potential (1.5 V vs RHE compared to open circuit potential conditions) in oxygen vacancies-containing hematite photoelectrodes (**Figure 5a**).⁸⁵ The application of an oxidative potential resulted in a slight shift of the absorption edge in the XANES measurement towards positive energies, which was tentatively associated with the Fe²⁺/Fe³⁺ oxidation, given that the presence of oxygen vacancies in the hematite structure would generate Fe²⁺ atoms. Likewise, for M-doped α -Fe₂O₃ photoanodes (where M = Nb or Ta), an increased X-ray absorption of the Fe L-edge, upon illumination, was observed for the doped hematite photoelectrodes.⁸⁶ This increased absorption was assigned to an enhanced electron transfer in the Fe 3d conduction band of hematite giving rise to the observed improved photocurrent.

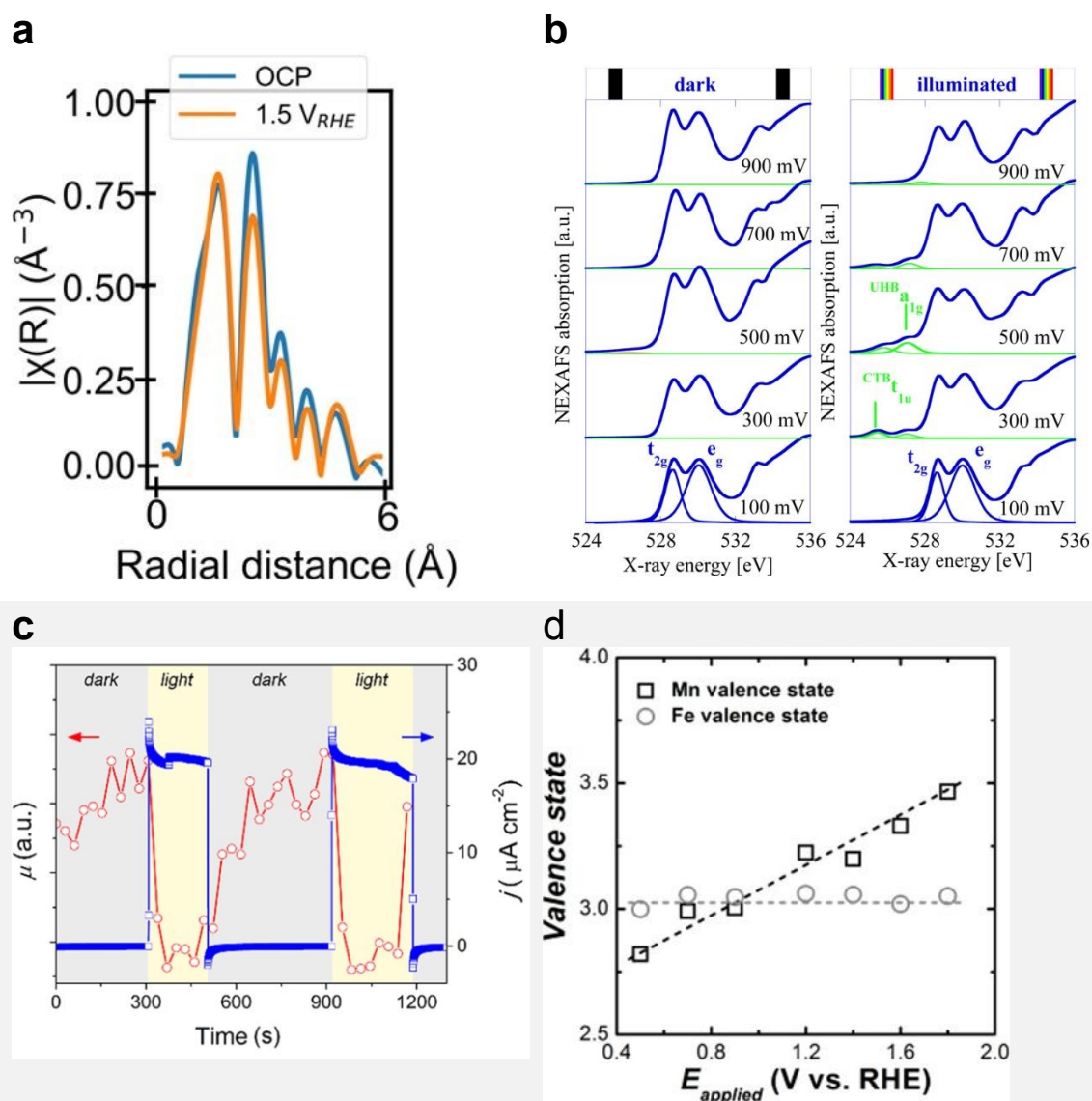
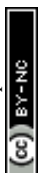


Figure 5. a) R-space plot of the Fe K-edge XANES spectra of the hematite (α -Fe₂O₃) photoanode in open circuit potential (i.e., no applied potential, blue trace), applying 1.5 V vs RHE (orange trace). Copyright 2024. Reproduced with permission from Springer Nature.



Adapted from reference ⁸⁵. b) O 1s NEXAFS spectra recorded at bias from 100 to 900 mV under dark (left) and light (right) conditions. Copyright 2012. Reproduced with permission from American Chemical Society. Adapted from reference ⁸⁷. c) X-ray absorption coefficient (μ) at fixed energy (8350 eV) and photocurrent density at a fixed potential (1.3V) acquired simultaneously, switching the light on and off. Copyright 2020. Reproduced with permission from American Chemical Society. Adapted from reference ⁸⁸. d) Valence states of Mn and Fe on α -Fe₂O₃/Mn₃O_{4+ δ} photoanodes during photoelectrochemical OER. Copyright 2018. Reproduced with permission from American Chemical Society. Adapted from reference ⁸⁹.

On the other hand, both O 1s NEXAFS and O K-edge measurements have been used to investigate the electronic structure of the photogenerated electrons and holes in hematite photoanodes.⁹⁰ For example, *ex-situ* time-resolved O K-edge XAS was used to identify the transient (~0.3 ps) formation of O 2p hole states upon photoexcitation.⁹¹ Interestingly, Braun *et al.*, identified two transitions in the pre-edge region of the in situ O 1s NEXAFS as a function of applied potential under light and dark conditions. These two transitions appear to be separated by 1.5 eV and were assigned to photogenerated holes of both Fe 3d and O 2p character (**Figure 5b**).⁸⁷ Although changes in the O 1s NEXAFS due to changes in the filling state of oxygen vacancies have not been ruled out,⁹² evidence of the electronic and chemical nature of photogenerated holes, in particular those at the surface, is still lacking.

Interfacing photoelectrodes with electrocatalysts to enhance charge transfer to the electrolyte is a highly used engineering strategy in PEC applications. Thus, studying and understanding the interface is crucial for producing highly efficient photoelectrode architectures. For example, Naldoni *et al.*, studied the charge transfer at the interface between hematite and a highly active OER electrocatalyst, NiO_x by means of fixed-energy XAS in both, potentiostatic and galvanostatic conditions.⁸⁸ In this work, the authors probed the Ni K-edge to probe the hole transfer from the hematite photoanode to the NiO_x catalyst layer prior OER. Interestingly, under photoelectrochemical water oxidation conditions, the differential XANES spectra (light – dark) coincided with the differential XANES spectra of γ -NiOOH – Ni(OH)₂ standards. Given that, in NiO_x electrocatalysts, the active species towards OER are reported to be NiOOH, Naldoni and *et al.*, assigned their differential XANES spectra to the accumulated OER active species formed after a hole transfer process from hematite. Additionally, the authors performed time-resolved (in the s timescale) X-ray absorption of a fixed energy (8350 eV), thus probing the X-ray absorption coefficient of the NiO_x during OER (**Figure 5c**). The observed reduction in the X-ray absorption coefficient, measured at 1.3 V, was assigned to the transformation of Ni(OH)₂ into NiOOH, which is recovered upon turning off the light.⁸⁸ A similar system, but with a NiFeOOH electrocatalyst was studied by Ismail *et al.*, reporting, through a linear combination analysis, that a γ -FeOOH phase is formed at the α -Fe₂O₃/NiFeOOH interface under illumination and a positive potential (**Figure 6**).⁹³

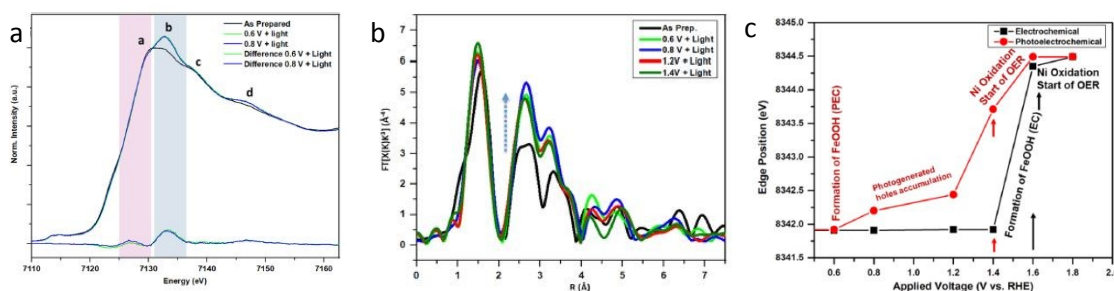
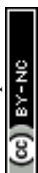


Figure 6. Operando XAS of iron in the α -Fe₂O₃/Ni_{0.8}Fe_{0.2}OOH photoanode. a) Iron K-edge XANES at different applied potentials under illumination. b) FT-EXAFS of Fe at different applied potentials under illumination. c) Plot of the α -Fe₂O₃/Ni_{0.8}Fe_{0.2}OOH photoanode at different applied potentials in the dark (EC) and under illumination (PEC), explaining the OER mechanism. Copyright 2021. Reproduced with permission from American Chemical Society. Adapted from reference ⁹³.



The increase of the PEC water oxidation rate upon the deposition of co-catalysts, based on metals such as Co and Mn, on $\alpha\text{-Fe}_2\text{O}_3$ photoanodes have also been investigated by in situ XAS means. For example, Lange *et al.* measured in situ the Co K-edge on $\alpha\text{-Fe}_2\text{O}_3/\text{CoB}_i$ (cobalt borate) photoanodes.⁹⁴ In this work, the authors, through the use of Co standards with precise oxidation states, calculated a slightly higher Co oxidation state when photodeposited on hematite than when grown on FTO. This was assigned to a built-in potential at the interface $\alpha\text{-Fe}_2\text{O}_3/\text{CoB}_i$. Additionally, in situ measurements revealed that, the CoB_i oxidises by accepting holes from the hematite and the authors suggest that a fraction of them oxidise water. Likewise, Liu *et al.*, reported an increase in the oxidation state of Mn in $\alpha\text{-Fe}_2\text{O}_3/\text{Mn}_3\text{O}_{4+\delta}$ photoanodes during photoelectrochemical OER.⁸⁹ However, in this case, the authors did not observe any change in the oxidation state of Fe and suggested the transfer of holes from hematite to the $\text{Mn}_3\text{O}_{4+\delta}$ catalyst (**Figure 5d**). It is possible that the authors did not observe any changes in the absorption of the Fe K-edge given the thickness of the reported photoelectrode.

In general, it is apparent from the reported studies that the oxidation of the metal centers in co-catalysts overlayers can be readily monitored by in situ XAS, most likely due to the low thickness exhibited by such co-catalyst's layers. Once hole transfer takes place to thin layers of co-catalysts, metal centers oxidise, and a change in the X-ray absorption edge is more straightforward to observe and quantify. As such, to further determine the nature of photogenerated holes, in bare hematite photoelectrodes, that are active for OER, it is key to use thin $\alpha\text{-Fe}_2\text{O}_3$ photoanodes.

Moving to BiVO_4 , Lifei Xi *et al.*⁹⁵ reported the study of a MnPi catalyst layer over a BiVO_4 photoanode by in situ soft X-ray absorption spectroscopy (XAS) at the Mn L-edge upon varying the applied potentials and the illumination conditions (**Figure 7**). As explained before, in situ NAP-XPS is the most suitable approach for PEC conditions, but it can be challenging to obtain reproducible spectra due to the varying thickness of the electrolyte layer and possible pressure changes inside the experimental chamber.³³ Because of that, the authors of this work focused on soft-XAS, because while hard-XAS probes the transition metal K-edge, with soft XAS the metal L-edge is probed, being more favourable for deciphering mixtures of oxidation states due to the narrower natural line widths (Mn K-edge, 1.12 eV; Mn L-edge, 0.32 eV).⁹⁶ To perform these experiments, the authors developed an in situ and operando cell.^{97,98} In these works, the authors reported an innovative design for an operando soft X-ray transmission and fluorescence cell that allows rapid membrane replacement and simpler operation compared to earlier designs. Specifically, the cell consists of three chambers (a vacuum chamber and two helium-filled chambers) separated by silicon nitride (Si_3N_4) membranes. This cell allows for the confinement of a liquid or electrolyte between these membranes and supports the application of both, electrochemical techniques and visible light. It includes a gold-coated membrane for conductivity, photodiodes for detecting transmission and fluorescence signals, and LEDs for visible light application. The helium pressure can be regulated to adjust the liquid layer thickness, and the cell can be precisely positioned using a translational and angular stage, facilitating in situ studies of solid/liquid interfaces with X-ray absorption spectroscopy. Specifically, Mn L-edge spectra under different applied potentials and light conditions were studied. The authors found that the photogenerated holes transferred into the MnPi layer and to the electrolyte interface was enhanced by a band bending in the semiconductor under illumination which is related to an increased potential by a built-in photopotential. They also found that the photogenerated holes could oxidize the MnPi co-catalyst layer even at open circuit potential conditions. Then, when increasing the applied bias, the electronic and crystalline structures of the MnO_x layer were changing to a birnessite-type layer structure associated with the electron migration from the MnPi film into the BiVO_4 photoanode. The formation of birnessite must be accompanied by the incorporation of K^+ between the MnO_6 layers, leading, for example, to a $\text{KMn}_8\text{O}_{16}$ phase. These experiments were able to conclude that Mn^{3+} species, instead of Mn^{4+} , in birnessite were the active sites for the OER.



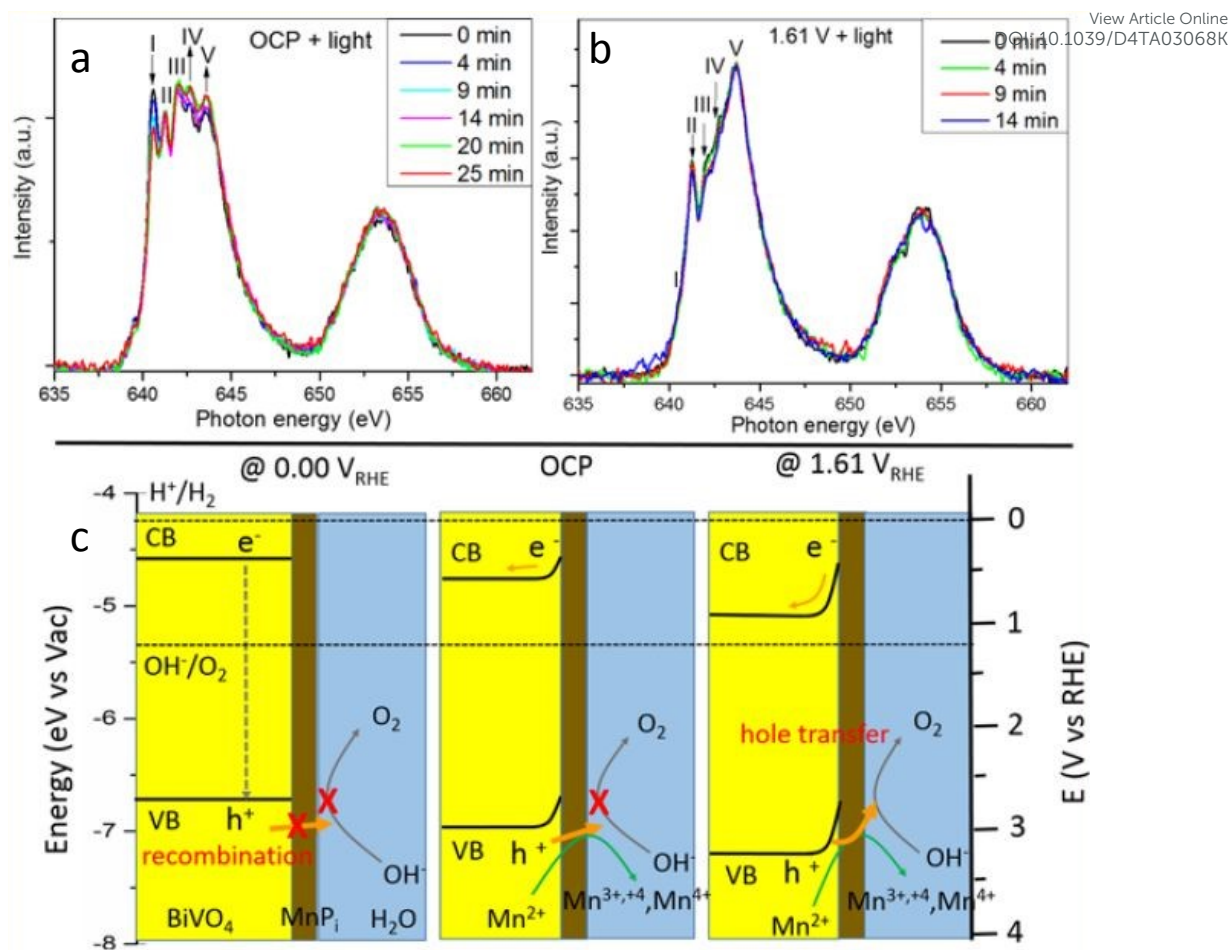


Figure 7. Temporal evolution of XAS spectra of a MnPi-modified BiVO₄/Au/Si₃N₄ photoanode in 0.1 M KPi buffer solution with KNO₃ added at a) OCP and b) 1.61 V under light. c) Schematic illustration of the photogenerated carrier transfer from the BiVO₄ photoanode to the MnPi film at 0.00 V, OCP condition, and 1.61 V vs RHE. Copyright 2017. Reproduced with permission from American Chemical Society. Adapted from reference ⁹⁵.

A couple of years later, some of the team members of the previous example, Lifei Xi and coworkers,⁹⁹ studied a NiBi co-catalyst over a BiVO₄ photoanode by in situ soft and hard-XAS. The authors successfully unravel the electronic structure of NiBi-modified BiVO₄ photoanodes under working conditions (**Figure 8a**) and understand the OER catalytic cycle and the formation of active sites (**Figure 8b**). The authors showed that Ni was in 2+ state in the electrodeposited amorphous NiBi film and it was readily oxidized up to 4+ state under OER conditions. This work reported for the first time the direct observation of Ni⁴⁺ species during light-driven water oxidation. They also observed that the oxidation of Ni was accompanied by an O K-edge prepeak at 529.17 eV. Specifically, the formation of Ni⁴⁺ was associated with the formation of electron deficient oxygen sites. These sites acted as electrophilic centers leading to the formation of a stable precursor state prior to O–O bond formation. Finally, cyclic voltammetry–XAS results supported that the formation of Ni⁴⁺ happened before the formation of partly electron deficient oxygen sites.



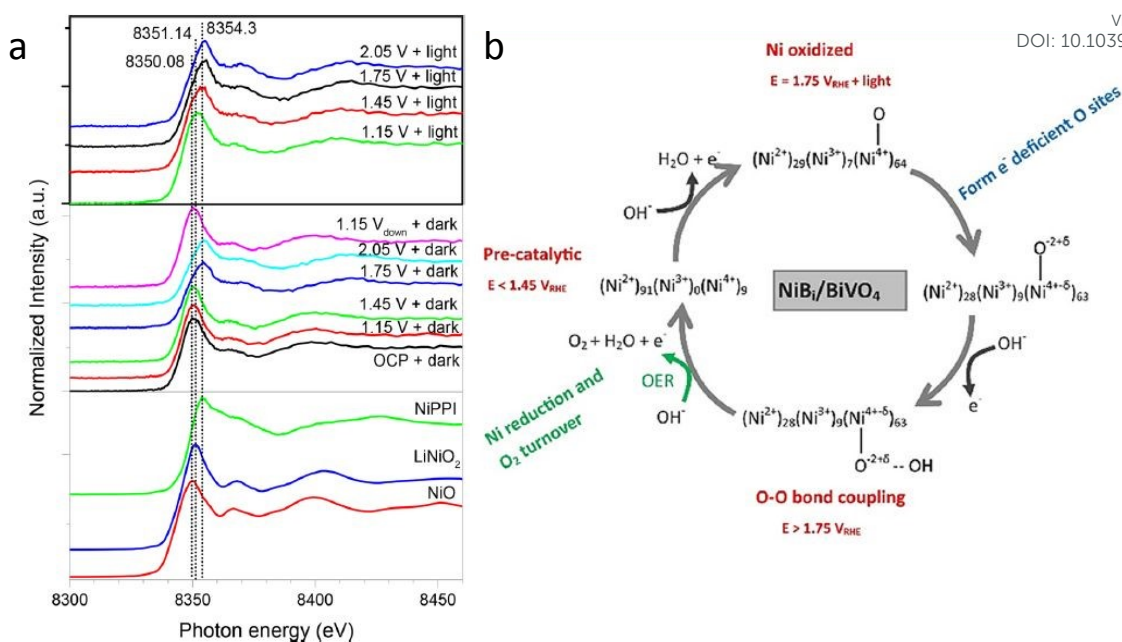
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Figure 8. a) In situ Ni K-edge spectra of the NiBi/BiVO₄ photoanode at different potentials under illumination. b) Schematic showing the NiBi co-catalyst change observed in situ XAS measurements of the NiBi/BiVO₄ photoanode. Copyright 2019. Reproduced with permission from American Chemical Society. Adapted from reference ⁹⁹.

All the above-explained works have demonstrated how in situ-XAS measurements are a powerful tool to determine the catalytic mechanisms (even catalytic cycles) in photoelectrodes, achieving a true understanding of these materials under relevant working conditions. It is also crucial to develop a correct design for the photoelectrochemical cells which allows the performance of the in situ and operando experiments with all the above-mentioned requirements. In general, the photoelectrochemical cells found in the reported studies are custom-made, either 3D printed or machined to work in both fluorescence and transmission modes. **Figure 9** displays different PEC cell designs and configurations. Firstly, two examples for in situ XAS measurements in fluorescence mode are shown in **Figures 9a** and **9b**. **Figure 9a** shows a (photo)electrochemical cell with an X-ray and visible light transparent window, *i.e.*, Mylar, and the sample illumination is from the photoelectrode/electrolyte interface. In this type of photoelectrochemical cell, the thickness of electrolyte that the X-rays have to travel through is between 0.1 to 1 mm and the sample is illuminated from the same direction of the X-rays irradiation, both to the front of the sample.^{73,100,101} Similarly, **Figure 9b** shows a cell configuration where the photoelectrode is directly prepared on a X-ray transparent material such as Si₃N₄, thus the X-rays irradiation and fluorescence is performed through the back of the photoelectrode. In this case the electrolyte thickness plays a less relevant role and the illumination is achieved from the front of the photoelectrode.^{87,102} Alternatively, when the photoelectrode is deposited on a non-X-ray transparent material, such as FTO, the photoelectrode can be placed facing the X-ray transparent window, where Kapton, Si₃N₄ and Mylar have been used at maximum 1 mm distance. In this configuration, the X-ray irradiation comes from the front (*i.e.*, electrode/electrolyte interface) and the visible excitation is performed through a mica/quartz window at the back of the photoelectrode.^{88,93} On the other hand, for measurements of soft X-ray XAS in transmission mode are presented in **Figures 9c** and **9d**. **Figure 9c** shows the schematic illustration of a tight (photo)electrochemical cell able to support measurements in vacuum by having integrated the illumination system, by LEDs, in the electrochemical chamber.⁹⁸ Likewise, **Figure 9d** presents a cell configuration where the catalyst can be deposited on a X-ray transparent, Au-coated Si₃N₄ membrane.



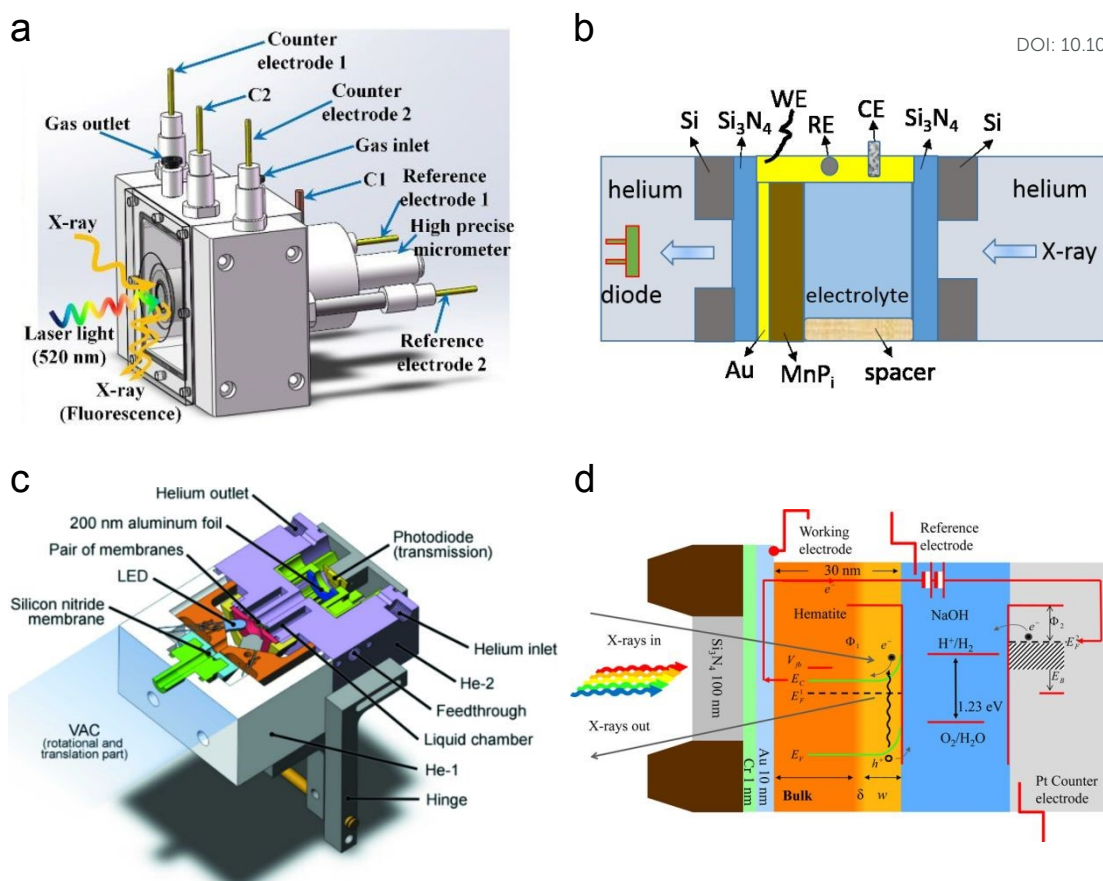


Figure 9. Different cell configurations for in situ/operando XAS experiments. a) Copyright 2021. Reproduced with permission from American Chemical Society. Adapted from reference ⁷³. b) Copyright 2017. Reproduced with permission from American Chemical Society. Adapted from reference ⁹⁷. c) Copyright 2016. Reproduced with permission of IUCr. Adapted from reference ⁹⁸. d) Copyright 2012. Reproduced with permission from American Chemical Society. Adapted from reference ⁸⁷.

In situ and operando X-ray diffraction (XRD) and X-ray scattering (XRS)

Diffraction happens with the scattering of incident X-rays by a certain material at different angles. Scattered X-rays are intense in some directions, as a consequence of constructive interferences, and deficient in other ones, due to destructive interferences. The constructive interference can be explained by Bragg's law, given by:

$$2d_{hkl} \sin(\theta) = n\lambda \quad (\text{Equation 6})$$

where, h_{kl} are the Miller indices for identifying the crystallographic planes, d_{hkl} represents the distance between parallel planes, which is a function of the unit cell characteristics (shape and size), θ is half of the angle between the incident and scattered beams, n is an integer corresponding to the plane from which the beam is scattered, and λ is the wavelength of the incident light.¹⁰³ An XRD spectrum represents the X-Ray diffraction intensity at different scattering angles, where the structure and size of the analysed material produce a certain XRD pattern. During the last decades, different diffraction-based techniques have been developed for different applications.

X-ray diffraction (XRD) is a powerful technique to study the atomic crystal structure of bulk materials. This technique is based on the scattering of hard X-rays (photon energies of 5–25 keV) by the crystal lattice of the sample, which in turn provides information about the structure of nanoparticles, surfaces, interfaces and defects, and even average structural arrangement within



liquids and at solid/liquid interfaces.¹⁰⁴ In situ XRD has been widely used to correlate X-ray diffraction and electrochemistry. Due to the high energy nature of the employed X-rays for XRD, easily able to pass through the electrolyte, performing successful in situ and operando experiments is easier compared to XAS and XPS. To be more surface-sensitive, Grazing Incidence X-ray Diffraction (GIXRD) is employed. In this configuration, the incident X-rays beam is directed onto a sample at a very low angle of incidence (usually $< 1^\circ$), interacting only with the top few nanometers of the material.¹⁰⁵ However, the examples reported so far for PEC applications are very scarce, especially for solar fuels generation. Kang *et al.*¹⁰⁶ used in situ XRD, together with pole figures and HR-TEM, to investigate the formation process and the epitaxial properties of a Bi-WO₃ photoelectrode for PEC water oxidation. In situ XRD patterns revealed the process of self-assembly-oriented construction during the temperature-programmed calcination of the film, finding the preferential formation of monoclinic WO₃ nanocrystals at high temperature ($T \geq 470^\circ\text{C}$) with dominant orientation along the (002) direction. Very recently, Wang *et al.*¹⁰⁷ performed quasi in situ GIXRD to study the phase structure and chemical composition of a PdCu/TiO₂/Si photocathode surface before and after Li-mediated PEC nitrogen reduction reaction (NRR). In GIXRD measurements, the incident angle was close to the critical angle for total reflection to increase the X-ray scattering signal.¹⁰⁴ By using this technique, the authors showed the appearance of a strong peak associated with Li₂N₂O₃ as the intermediate from the oxidation of Li or Li₃N under ambient air. Thus, these results revealed the formation of Li₃N from Li and N₂ during the PEC NRR, and also showed a higher stability of Li₃N and Li₂N₂O₃ in air than Li.¹⁰⁷

X-ray scattering (XRS) is an all-atom technique that allows to resolve structural parameters across the distance range from 10^{-11} m to 10^{-6} m.¹⁰⁸ Whereas XRD describes the scattering of X-rays by a crystal lattice, other X-ray scattering techniques investigate the atomic, nano, and micro scale structure of less ordered matter, providing structural information of nanoparticles, defects, surface interfaces, and even the average structural arrangement within liquids or at liquid interfaces.¹⁰⁴

Small-angle X-ray scattering (SAXS) can be used to study morphological and structural changes in photoelectrodes and electrolytes, since it allows to determine inhomogeneities ranging from the atomic up to nanometer length scales.^{104,109} Thus, it allows to probe local atomic structures and the formation of new phases on the photoelectrode surface.¹⁰⁹ More specifically, grazing incidence small-angle X-ray scattering (GISAXS) technique is used for studying the nano- and mesoscale evolution of surface structure in the range between one to hundred nanometers. This is a powerful technique for time-resolved operando studies during interface reactions, since GISAXS images can be obtained within seconds. Besides, this technique is insensitive to the crystal structure and it can be performed on any sufficiently planar interface, including amorphous and liquid substrates.¹⁰⁴

Koziej *et al.*¹⁰⁹ developed an operando grazing-incidence X-ray scattering approach to investigate the photocorrosion mechanisms and the evolution of structural modifications of a CuBi₂O₄ photoelectrode under illumination and applied bias. A scheme of the employed setup is shown in **Figure 10a**. In this study, the total scattering (TS) measurements at grazing-incidence angle were collected by using a high-energy X-ray probe and a large-area detector. Simultaneously, they also used a second large-area detector to collect the GISAXS signal, which was positioned at a different sample-to-detector distance than TS to acquire both signals at the same time. Specifically, by means of simultaneous and SAXS measurements they found that CuBi₂O₄ electrodes quickly underwent a cathodic photocorrosion only few minutes after beginning of the chronoamperometry, and a decrease of around one order of magnitude in the photocurrents. The authors ascribed these results to the formation of a segregating metallic Bi phase on the surface of the electrode, shown in **Figure 10b**, which significantly reduced the CuBi₂O₄ surface area in direct contact with the electrolyte. Besides, they observed an additional degradation processes at slower timescales, associated with the formation of metallic Cu and subsequent dissolution in the electrolyte as a result of anionic photocorrosion and acidic dissolution. Both Cu²⁺ and Bi³⁺ ions released in solution interacted with the phosphate species from the electrolyte forming an insulating BiPO₄ layer on the surface of the electrode. This proposed mechanism is schematised in **Figure 10c**. On view of these results, they suggested to avoid the use of a phosphate buffer to



prevent the unfavourable formation of BiPO_4 . They also suggested the employment of a HER co-catalyst to avoid the reduction of CuBi_2O_4 to metallic Bi and Cu by quickly transferring the photogenerated electrons to the liquid phase.

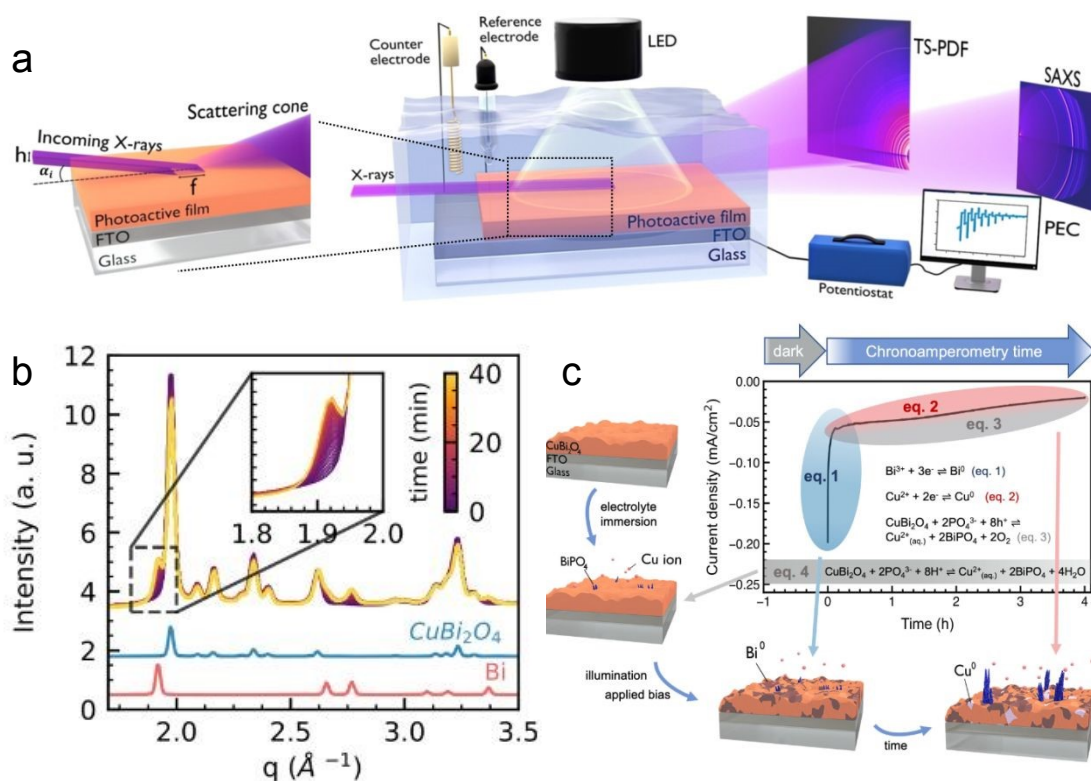


Figure 10. a) Schematic illustration of the multi-modal experimental set up: PEC properties of a photoactive thin film under illumination and immersed in an electrolyte solution are probed by use of a potentiostat in a standard 3-electrode configuration. Two large-area X-ray detectors permit the simultaneous measurement of both TS and SAXS signals from the film surface in a grazing incidence geometry. b) Time-resolved plot of the operando X-ray scattering signal collected from the surface of a CuBi_2O_4 electrode under illumination and applied constant bias (0.5 V vs RHE). In the inset, a zoom into the 1.8–2.0 \AA^{-1} region highlights the formation of metallic Bi with time. c) Schematic summary of the multiple reaction pathways and relative time scales determining the degradation of CuBi_2O_4 photocathodes before and during the chronoamperometric test. Copyright 2023. Reproduced with permission from Wiley. Adapted from reference ¹⁰⁹.

Wide-Angle X-ray Scattering (WAXS) can provide signals from the catalyst at high diffraction angles, without any influence from the background or the PEC cell, and with higher temporal resolution than SAXS.¹⁰⁴ even we were not able to find any work using in situ or operando WAXS in a PEC system, in 2020, Strasser *et al.*¹¹⁰ were able to combine electrochemical OER measurements with operando WAXS, XAS data and theoretical calculations (ab initio molecular dynamic simulations and DFT) to track structural transformations during the activation of NiFe and CoFe LDH catalysts. These authors provided the first direct atomic-scale evidence for the oxidation of LDH from their as-prepared precursor state (α -phase) to the catalytically active states (deprotonated γ -phase) at a potential below the onset of the OER. The active γ -phases showed around 8% contractions of the lattice spacing, in both the in-plane lattice constant and the interlayer distance induced by the oxidation of the cations and the dynamics of the intercalated ions. This work demonstrates the high potential of WAXS for the understanding of the structural changes of electrocatalytic and surely photoelectrocatalytic materials under working conditions.



As it has been extensively detailed along this manuscript, (photo)electrochemical cell configurations and the information provided by XPS, XAS and XRD techniques vary significantly. Summarising, XPS analyses surface chemistry by detecting electrons emitted from a material, revealing elemental composition and chemical states of the top few nm. On the other hand, XAS, including both XANES and EXAFS, examines how a material absorbs X-rays to provide insights into local atomic structures, coordination environments, and oxidation states. Finally, XRD uses diffracted X-rays to determine the crystallographic structure, phase identification, and other structural properties of a material. Then, each technique serves specific purposes, where XPS focuses on surface analysis, XAS on local atomic environments and overall elemental composition and XRD on crystal structure. Hence, due to these technical differences, the PEC cells for in situ and operando characterisation need to be designed precisely. **Table 1** summarises the most relevant technical aspects of the analysed samples in the present study.

	XPS	XAS		XRD	XRS
		Soft-XAS	Hard-XAS		
Principle	Photoemission	Absorption		Diffraction	Scattering
Techniques	UHV-XPS NAP-XPS	Total Electron Yield Total Fluorescence Yield		Conventional XRD GIXRD	SAWS WAXS
Acquisition Depth	1-10 nm	TEY: ~10-20 nm TFY: ~100nm – 100µm	TEY: ~10-20 nm TFY: ~mm	1-100 nm	1-100 nm
Detection limit	0.1-1.0% at.	10-100 g/g	10-100 µg/g	Depends on crystallinity of the sample	Depends on crystallinity and long range order of the sample
Advantages	Surface sensitive data acquisition dynamics (~seconds) Quantitative information	Easier in-situ cell configuration than for XPS.	Ease of in-situ and operando cell configuration Possibility of obtaining surface sensitive data (GIXRD, GISAWS)		
Disadvantages	In-situ cell complexity (pressure conditions, membranes, etc.)	Complexity of surface sensitive measurements More qualitative than quantitative Data acquisition (minutes to hours)		More qualitative than quantitative Data acquisition (tens of minutes to hours)	
		Cell complexity in comparison to XRD and hard XAS	Difficult to decouple surface and bulk chemical changes	Only for crystalline samples	Complex data analysis (WAXS) SAWS (only for crystalline samples)

Table 1. Most relevant technical aspects of XPS, XAS and XRD techniques.

Apart from the above-detailed classic X-ray-based techniques (XPS, XAS and XRD), there are other configurations that are currently being developed at synchrotron facilities. Recently, several instruments allow to perform different techniques in the same camera, known as *multi-technique* or *multi-modal approach*, such as NOTOS beamline in ALBA synchrotron, which allows to perform XRD and XAS. Moreover, 3Sbar beamline in ALBA, allows to perform XPS and SAXS. On the other hand, in Max IV, XPS and IR are possible to be carried out. Besides, NAP XPS in synchrotron, also allows to perform Auger yield XAS, which is surface sensitive. The main advantage of this multi-technique approach, summarized in the previous examples, relies on getting information from the exact same point in the sample and under exactly the same conditions. Moreover, there are other techniques such as tomography, for example available at Faxtor beamline in ALBA, that give a 3D image of the sample and could be well adapted to in situ and operando experiments. All these spectroscopies have demonstrated an enormous potential



in the characterization of materials in general, and PEC systems in particular, under working conditions. Article Online
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The future of XPS, XAS, and XRD techniques for in situ and operando characterisation of photoelectrodes is highly promising, where significant advancements are expected. These advances include the development of ambient pressure and time-resolved XPS for realistic and dynamic surface studies, improved resolution and sensitivity in XAS through next-generation synchrotron sources as well as the integration of XAS with neutron scattering for comprehensive structural insights. Additionally, real-time and rapid XRD measurements, along with X-ray tomography, will enhance understanding of structural changes and phase distribution in photoelectrodes. Additionally, the synergy of multi-modal techniques, as detailed above, and the use of artificial intelligence for data analysis will provide a more comprehensive characterization, driving the discovery and optimization of new materials. Therefore, the enhanced operational conditions during measurements will lead to more precise and relevant studies, ultimately contributing to the development of more efficient and durable photoelectrocatalytic materials.

Conclusions

The present work summarizes the most relevant in situ and operando X-ray studies in PEC systems, demonstrating the strength and potential of these spectroscopies for the characterisation of PEC materials under working conditions. Specifically, this work is focused in operando and in situ XPS, XAS and XRD techniques, trying to remark the recent progress using these techniques for elucidating the structural, chemical, compositional and electronic properties of photoelectrocatalytic materials and interfaces. Moreover, this work gives a technical guidance for adequately carrying out these complex experiments, focusing on the possibilities of the analysed techniques, cell designs and more relevant results.

It is apparent that many of the above discussed X-ray-based techniques have benefited from the developments and upgrades of different synchrotron beamlines. Accordingly, we believe that the success of the in situ and operando characterisation is currently mainly limited by the PEC cell engineering, where film and electrolyte thickness, as well as photoelectrode disposition, play a key role. In this context, making the cell designs publicly available will greatly benefit the field and will promote a deep understanding of PEC processes aiming to design more efficient and stable photoelectrodes for solar-driven fuels production and added-value photoelectrosynthesised compounds.

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Data Availability Statement

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