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## Determining the Chemical Composition of Corrosion Inhibitor/Metal Interfaces with XPS: Minimizing Post Immersion Oxidation --Manuscript Draft--

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<b>Abstract:</b>	An approach for acquiring more reliable X-ray photoelectron spectroscopy data from corrosion inhibitor/metal interfaces is described. More specifically, the focus is on metallic substrates immersed in acidic solutions containing organic corrosion inhibitors, as these systems can be particularly sensitive to oxidation following removal from solution. To minimize the likelihood of such degradation, samples are removed from solution within a glove box purged with inert gas, either N <sub>2</sub> or Ar. The glove box is directly attached to the load-lock of the ultra high vacuum X-ray photoelectron spectroscopy instrument, avoiding any exposure to the ambient laboratory atmosphere, and thus reducing the possibility of post immersion substrate oxidation. On this basis, one can be more certain that the X-ray photoelectron spectroscopy features observed are likely to be representative of the in situ submerged scenario, e.g. the oxidation state of the metal is not modified.
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**TITLE:**

**Determining the Chemical Composition of Corrosion Inhibitor/Metal Interfaces with XPS:  
Minimizing Post Immersion Oxidation**

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**KEYWORDS:**

X-ray photoelectron spectroscopy, corrosion inhibition, acidic solution, metallic substrate, oxidation, inert atmosphere.

**SHORT ABSTRACT:**

A protocol to avoid the oxidation of metallic substrates during sample transfer from an inhibited acidic solution to an X-ray photoelectron spectrometer is presented.

**LONG ABSTRACT:**

An approach for acquiring more reliable X-ray photoelectron spectroscopy data from corrosion inhibitor/metal interfaces is described. More specifically, the focus is on metallic substrates immersed in acidic solutions containing organic corrosion inhibitors, as these systems can be particularly sensitive to oxidation following removal from solution. To minimize the likelihood of such degradation, samples are removed from solution within a glove box purged with inert gas, either N<sub>2</sub> or Ar. The glove box is directly attached to the load-lock of the ultra high vacuum X-ray photoelectron spectroscopy instrument, avoiding any exposure to the ambient laboratory atmosphere, and thus reducing the possibility of post immersion substrate oxidation. On this basis, one can be more certain that the X-ray photoelectron spectroscopy features observed are likely to be representative of the *in situ* submerged scenario, e.g. the oxidation state of the metal is not modified.

**INTRODUCTION:**

Corrosion inhibitors (CIs) are substances that, when introduced into an aggressive environment, reduce the corrosion rate of a metallic material by inducing a change at the solid/liquid interface<sup>1-5</sup>. This approach to corrosion control is widely used in industry, with high performance CIs having been successfully developed for a variety of applications. There remains, however, a considerable lack of fundamental understanding of CI performance, hindering knowledge-based optimization. For example, the precise nature of interfaces formed by organic-CIs in corrosive acidic solutions is still unclear.

Given the assumption that organic-CIs inhibit acidic corrosion through formation of an adsorbed 2-D layer<sup>2</sup>, a surface sensitive technique is required to characterize these interfaces. Consequently, X-ray photoelectron spectroscopy (XPS)<sup>6</sup> has emerged as a technique of choice to probe the elemental/chemical composition of these interfaces *ex situ*<sup>7-9</sup>; XPS measurements are typically performed in, or close to, ultra high vacuum (UHV).

Various insights have been claimed, including that a surface oxide or hydroxide is present to facilitate organic-Cl binding to the metallic substrate<sup>10-12</sup>. The validity of this interface description, however, is questionable as the XPS data were acquired from samples that had been exposed to the laboratory atmosphere in between removal from inhibited solution and introduction into the UHV-XPS spectrometer. Such a procedure may result in interface oxidation, undermining conclusions about interface chemical composition. An alternative approach is required, which minimizes the potential for post immersion oxidation.

In this paper, we detail a methodology designed to allow one to acquire XPS data from organic-Cl/metal interfaces that have not undergone oxidation following emersion from acidic solution. A glove box, purged with inert gas, which is attached directly to the vacuum load-lock of the UHV-XPS instrument is employed. The utility of our approach is verified through presentation of XPS data from two organic-Cl/carbon-steel interfaces formed following addition of sufficient Cl to significantly reduce the substrate corrosion rate in aqueous 1 M hydrochloric acid (HCl) solution.

## **PROTOCOL:**

### **1. Substrate/Solution Preparation**

#### **1.1) Preparation of carbon-steel substrate**

1.1.1) Cut an approximately 2 - 3 mm thick disc from a cylindrical carbon-steel rod ( $10 \pm 2$  mm diameter), using a precision cutting machine. Employ cooling fluid during the cutting procedure to minimize mechanical damage.

1.1.2) Grind both circular faces, and the edge of the disc-shaped sample with a series of SiC papers, i.e. 600 grit, 800 grit, 1200 grit, 2400 grit, and 4000 grit.

1.1.3) Polish both circular faces, and the edge of the disc-shaped sample with either diamond paste or alumina powder (3  $\mu$ m and/or 1  $\mu$ m) until a mirror finish is obtained.

1.1.4) Sonicate the sample at room temperature for approximately 10 minutes in each of the following: deionized water, acetone, ethanol, and finally deionized water again.

1.1.5) Dry all surfaces of the sample under a flow of air.

1.1.6) Store the sample in either a desiccator or vacuum desiccator until required. Ideally, finish preparing the carbon-steel substrate just prior to solution immersion.

#### **1.2) Preparation of inhibited 1 M HCl solution**

1.2.1) Add 10.2 M HCl to deionized water to produce a 1 M HCl solution. CAUTION: HCl is harmful and corrosive. Undertake this step in a fume hood, wearing suitable personal protection equipment.

1.2.2) Dissolve x g of selected organic-Cl (e.g. x = 0.300 g of 2-mercaptobenzimidazole) in 1 M HCl solution to produce a 1 M HCl solution containing x mM organic-Cl (e.g. x = 2 mM 2-mercaptobenzimidazole).

## **2. Substrate Immersion in Inhibited Acid Solution**

2.1.1) Pour a small volume (typically 25 – 50 mL) of the 1M HCl + x mM (e.g. x = 2 mM 2-mercaptobenzimidazole) organic-Cl solution into a small glass beaker.

2.1.2) Pick up the disc-shaped carbon-steel sample with either ceramic or plastic tweezers that are resistant to 1 M HCl. Only touch the edge of the sample with the tweezers.

2.1.3) Introduce the sample into the glass beaker containing the 1M HCl + x mM organic-Cl solution. Orient the sample so that cylindrical faces are in the vertical plane. Deposit the sample either directly onto the bottom of the beaker, or atop the open end of a short glass tube. Ensure that the sample is fully submerged.

## **3. Sample Transfer**

### **3.1) Glove box preparation**

3.1.1) Locate the circular sample-transfer opening in the base of the glove box onto the XPS load-lock flange mounting. Ensure that the seal between glove box and load-lock flange is formed correctly.

3.1.2) Connect the glove box to an inert gas cylinder (either N<sub>2</sub> or Ar).

3.1.3) Adhere a small square of double-sided carbon (conductive) tape onto the XPS sample bar.

3.1.4) Insert hardware required for sample transfer into XPS instrument through an open port on the glove box, i.e. sample bar with carbon tape attached, plastic/ceramic tweezers, nitrile gloves, laboratory tissues, plastic paraffin film, empty/dry wash bottle with spout, and glass beaker containing ~ 200 g of Na<sub>2</sub>CO<sub>3</sub> powder.

3.1.5) Place glass beaker containing carbon-steel sample in 1M HCl + x mM (e.g. x = 2 mM 2-mercaptobenzimidazole) organic-Cl solution inside glove box. Ensure that sample is always fully immersed during this step.

3.1.6) Seal all ports/entry points on the glove box and begin to purge with N<sub>2</sub> (or Ar).

3.1.7) Continuously purge the glove box until the sample transfer is complete.

### **3.2) From solution to XPS analysis chamber**

3.2.1) Allow the sample to remain submerged in the 1 M HCl + x mM organic-Cl solution for the desired immersion period, e.g. 4 h, as employed for the XPS data presented below.

3.2.2) Check that the relative humidity within the glove box is minimized, which is typically achieved after 60 – 90 min of initiating purging. A value of 8% relative humidity is required before proceeding with sample transfer; value prior to purging is typically 35 - 40%.

Note: Normally, there is no dedicated O<sub>2</sub> sensor within the glove box, but trial measurements with such a device indicate that the purging procedure leads to a reduction

in O<sub>2</sub> concentration by a factor of ~ 1000.

3.2.3) Introduce hands into glove box gloves, and then cover gloved hands with nitrile gloves located within the glove box. This step reduces the possibility of contamination during sample handling, as well as increases the ease of sample manipulation.

3.2.4) Remove the carbon-steel sample from the 1 M HCl + x mM (e.g. x = 2 mM 2-mercaptobenzimidazole) organic-Cl solution, using the ceramic/plastic tweezers. Only touch the edge of the sample with the tweezers.

3.2.5) **Immediately** after emersion, blow the sample dry by directing a stream of inert gas, produced by repeatedly squeezing empty/dry wash bottle located inside glove box, onto sample surfaces.

Note: Rinsing with solvents at this stage is not undertaken to minimize the potential of damaging the inhibited interface, e.g. inhibitor removal or interface oxidation.

3.2.6) Cover beaker containing 1 M HCl + x mM organic-Cl solution with plastic paraffin film.

3.2.7) Affix the sample to the small square of double-sided carbon (conductive) tape attached to XPS sample bar. Do not touch the surface to be probed by XPS.

3.2.8) Vent the XPS load-lock chamber to N<sub>2</sub>/Ar. Ensure that the associated turbo/rotary pump combination is switched off before performing this action.

3.2.9) Open the load lock flange.

3.2.10) Transfer the sample bar into the load-lock chamber and slide it onto sample holding prong.

3.2.11) Close the load-lock flange.

3.2.12) Switch on turbo/rotary pump combination to pump down load-lock chamber.

3.2.13) Once pressure in load-lock chamber reaches at least  $\sim 5 \times 10^{-7}$  mbar, manually transfer sample to intermediate chamber using transfer arm.

3.2.14) Wait until pressure in intermediate chamber reaches  $\sim 1 \times 10^{-8}$  mbar, and then use second transfer arm to manually transfer sample onto the sample manipulator in the analysis chamber.

#### **4. Acquisition of XPS data**

4.1) Orient angle of sample, using keypad to drive sample manipulator motors, to desired photoelectron emission angle, e.g. 0° (emission along the surface normal), as employed for the XPS data presented below.

4.2) Open XPS data acquisition software by clicking on the icon located on the desktop.

Open Instrument Manual Control window.

4.3) Power up the monochromated Al K $\alpha$  X-ray source by clicking the 'ON' button in 'X-ray Gun' section. Input 10 mA and 15 kV as desired values for the anode emission and anode HT parameters, respectively. Switch on charge neutralizer, if required, by clicking 'ON' button in 'Neutralizer' section.

4.4) Select electron analyzer 'Spectrum/Hybrid' measurement mode from the 'Mode' and 'Lens' drop down menus in the 'Analyzer' section.

4.5) Input desired kinetic/binding energy ranges, pass energy(ies), step size(s), and dwell time(s) into 'Acquisition/Scan Control' section. Example input parameter values: 1200 – 0 eV binding energy range, 80 eV pass energy, 0.5 eV step size, and 0.1 s dwell time to acquire overview spectra.

4.6) Optimize sample position by adjusting position of sample manipulator, using keypad to drive sample manipulator motors, to maximize signal from selected core level, e.g. C 1s or Fe 2p.

4.7) Acquire XPS spectra by initiating data collection in XPS data acquisition software.

#### REPRESENTATIVE RESULTS:

Figure 1 shows overview, O 1s, and Fe 2p XPS data acquired from carbon-steel samples that have been immersed for 4 h in one of two different 1 M HCl + x mM organic-Cl solutions, and transferred for XPS measurement as detailed above. Analogous data from a polished sample are also displayed. The carbon-steel possessed a nominal weight% composition of C (0.08-0.13), Mn (0.30-0.50), P (0.04), S(0.05), and Fe (balance). The two organic-Cl solutions studied are: 2-mercaptobenzimidazole (MBI) and (Z)-2-2(2-(octadec-9-en-1-yl)-4,5-dihydro-1H-imidazole-1-yl)ethanamine (OMID). At the concentrations employed (MBI: 2 mM; OMID: 1mM) corrosion rate measurements<sup>13,14</sup> indicate that all of the species significantly inhibit the corrosion of carbon-steel, i.e. inhibition efficiencies ( $\eta\%$ )<sup>2</sup> > 90%. Best fits to the O 1s and Fe 2p spectral profiles are also displayed. Photoelectron peaks were modeled with Gaussian-Lorentzian (GL) line shape functions (30 % Lorentzian), with the exception of the Fe 2p level of metallic iron, where a Lorentzian asymmetric line shape with tail damping (LF) was used. The cationic Fe<sup>x+</sup> states were modeled with multiplet envelopes consisting of 3 and 4 GL functions for Fe<sup>2+</sup> and Fe<sup>3+</sup>, respectively<sup>15</sup>. A Shirley-type function<sup>16</sup> was used to describe the background of inelastically scattered electrons.

Focusing on the overview XPS data (Figure 1 (a)), the spectrum acquired from the polished sample exhibits three prominent peaks, i.e. Fe 2p, O 1s, and C 1s. These features can be assigned as follows: Fe 2p arises from the carbon-steel, O 1s derives from both an surface oxidic film and adsorbates, and the C 1s signal is due to adventitious carbon. Submersion in any of the 1 M HCl + x mM organic-Cl solutions results in significant changes to the corresponding overview spectrum. A feature assigned to the N 1s core level appears, which is consistent with surface adsorption of the inhibitors; MBI and OMID both contain N. Moreover, the O1s core level signal is significantly diminished.



As regards O 1s data from the polished substrate (Figure 1 (b)), the profile can be fitted with four components. The two components at lower binding energies (BE),  $\sim 530.0$  eV and  $\sim 531.3$  eV, can be assigned to iron oxide ( $O^{2-}$ ) and hydroxide ( $OH^-$ ) phases, respectively. The two higher binding energy components, labeled  $O^1$  (BE  $\sim 532.2$  eV) and  $O^2$  (BE  $\sim 533.3$  eV), are likely associated with adsorbed OH ( $O^1$ ) and adventitious carbon species ( $O^1$  and  $O^2$ )<sup>17</sup>. Immersion in either of the 1 M HCl + x mM organic-Cl solutions leads to the complete quenching of the  $O^{2-}$  and  $OH^-$  components. On this basis, it can be concluded that the corrosion inhibitors are adsorbed on oxide/hydroxide free surfaces. The Fe 2p spectra in Figure 1 (c) are consistent with this result, as only a metallic Fe ( $Fe^0$ ) peak is apparent on the inhibited substrates.  $Fe^{2+}$  and  $Fe^{3+}$  features are present on the polished sample, due to the surface oxide/hydroxide.

O 1s and Fe 2p core level XPS spectra from two carbon-steel samples immersed in 1 M HCl + 2 mM MBI are compared in Figure 2. One sample was transferred using a fully  $N_2$ -purged glove box, whilst the other was removed from solution into a partially  $N_2$ -purged glove box, i.e.  $O_2$  concentration was significantly higher than target value. For the latter sample, it is apparent that post immersion oxidation has occurred, i.e.  $Fe^{2+/3+}$  and  $O^{2-}/OH^-$  features are present.

#### FIGURE LEGENDS:

**Figure 1. XPS spectra from polished and inhibited carbon-steel samples.** (a) Overview, (b) O 1s, and (c) Fe 2p XPS spectra. Data presented in each panel were acquired from carbon-steel samples that had been immersed for 4 h in one of two different 1 M HCl + x mM organic-Cl solutions, i.e. 2 mM MBI, and 1 mM OMID. Spectra were all acquired at a photoelectron emission angle ( $\theta_E$ ) of  $0^\circ$  (emission along the surface normal). For (b) and (c) the best fits (light blue markers) to the experimental data (solid black lines) are also displayed, achieved with a combination of GL (broken red lines), LF (broken red lines), and Shirley-type (broken grey lines) functions. Peak labels are explained in the main text.

**Figure 2. Impact of post immersion oxidation on XPS spectra.** (a) O 1s and (b) Fe 2p XPS spectra. Data were acquired from carbon steel samples that had been immersed in 1 M HCl + 2 mM MBI solutions ( $\eta\% = 99\%$ ) for 4 h. Spectra were acquired at  $\theta_E = 0^\circ$ . In each panel, the lower (upper) spectrum is from a sample transferred through a fully (partially)  $N_2$ -purged glove box. Peak labels are explained in the main text. (Modified version of Fig. 5 in Ref. 9.)

#### DISCUSSION:

XPS spectra displayed in Figures 1 and 2 clearly demonstrate that the inert atmosphere employed during sample transfer is essential to avoid post immersion oxidation of these carbon-steel/organic-Cl interfaces. On this basis, results from other similar XPS studies (e.g. <sup>18,19</sup>), which involved exposure of the inhibited substrate to the ambient laboratory atmosphere, should be critically re-evaluated, as the interface chemistry may have been modified through oxidation. It should be noted that there is no reason to assume that all organic-Cl adsorb onto oxide/hydroxide free surfaces in acidic solution. In some cases, such phases may indeed facilitate the organic-Cl surface binding. Distinguishing this scenario from post immersion oxidation is not so straightforward. One possible solution is to acquire XPS data from a *reference* inhibited interface (e.g. 1 M HCl + 2 mM MBI) in

tandem with the sample of interest to check that post immersion oxidation is not an issue.

To ensure a successful outcome to the sample transfer procedure, it is essential that the glove box is fully purged with inert gas ( $N_2/Ar$ ) i.e. the  $O_2$  concentration in the glove box is minimized. Care should be taken to verify that all seals on the glove box ports/entry points are correctly formed, including the seal between the glove box and XPS load-lock flange. Ideally, an *in situ* sensor should be used to directly monitor the  $O_2$  concentration, although it is not essential, as demonstrated by our own work. As indicated in step 3.3.2, we normally use a relative humidity sensor as a guide for when to undertake a sample transfer.

One further potential issue with the glove box environment is the presence of volatile solution components, which can contaminate the sample surface following removal from solution and prior to insertion into the load-lock. For example, the presence of 1 M HCl solution in the glove box leads to evolution of HCl vapor, which can react with carbon-steel samples leading to adventitious Cl signal in the XPS data. To minimize the likelihood of such contamination being significant, small volumes of HCl solution should be employed and sample transfer should be completed as quickly as possible. On this basis, as indicated in the Protocol, typically only one beaker/sample is inserted into the glove box for sample transfer at any one time. In addition, one should minimize the surface area of the HCl solution, as well cover the beaker following sample immersion.  $Na_2CO_3$  powder (step 3.1.4) is inserted into the glove box in an attempt to control the amount of HCl vapor. In addition, it can be used to clean up any acidic solution spillages.

Besides careful control of the glove box environment, sample handling is also critical to the integrity of acquired XPS spectra. One should not acquire XPS data from any surface that has come into contact with any solid object, e.g. tweezers or glove. Furthermore, upon removal of a sample from solution, it should be immediately blown dry with inert gas (step 3.2.5). This procedure is undertaken to prevent evaporation and subsequent physical deposition of solution components onto the sample surface, which may lead to misinterpretation of data. As an additional precaution, one can also consider replacing the nitrile gloves (step 3.2.3) with a fresh pair for transferring the sample bar to the load-lock chamber, i.e. prior to step 3.1.10.

Finally, given the effectiveness of the approach outlined here, we expect it to be applied to other topics in corrosion (i.e. in addition to corrosion inhibition), where acquisition of XPS data from air-sensitive interfaces would add to understanding. Moreover, such an approach should be considered in other fields where XPS measurements are undertaken from air-sensitive interfaces formed in a fluid environment. Clearly, this procedure is not restricted to XPS, but could also be applied to any other UHV-based measurement from a surface that has previously been immersed in a liquid, e.g. scanning probe microscopy.

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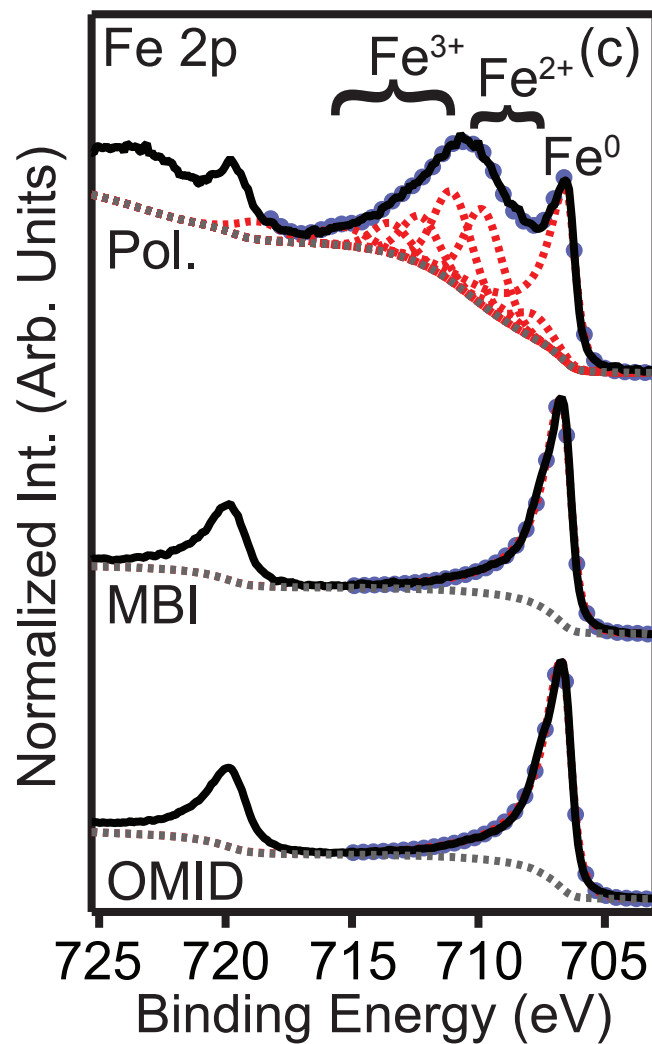
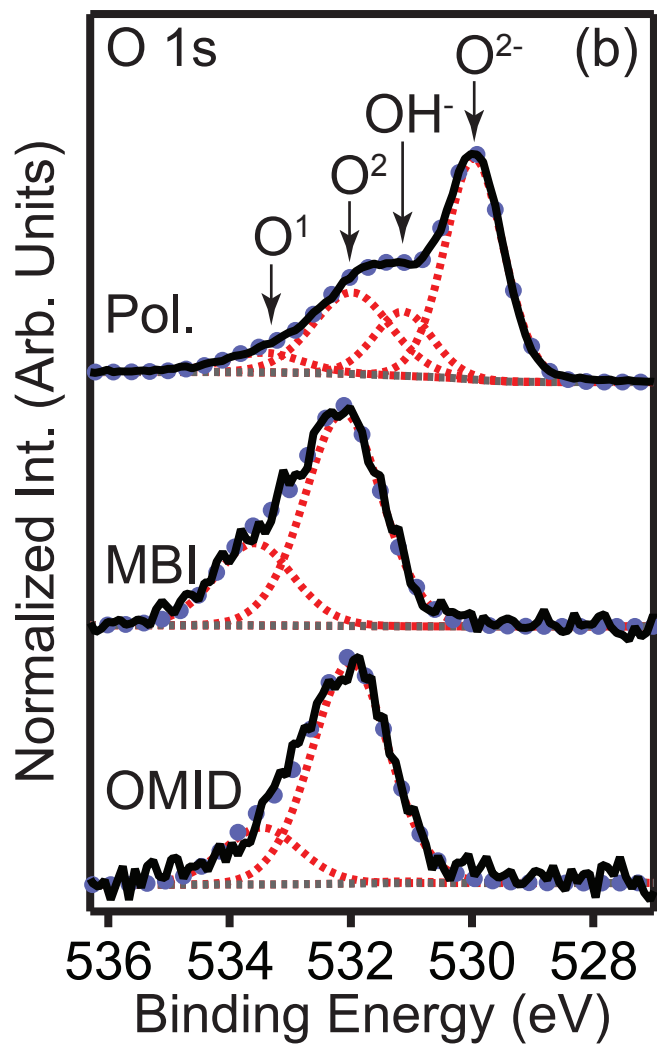
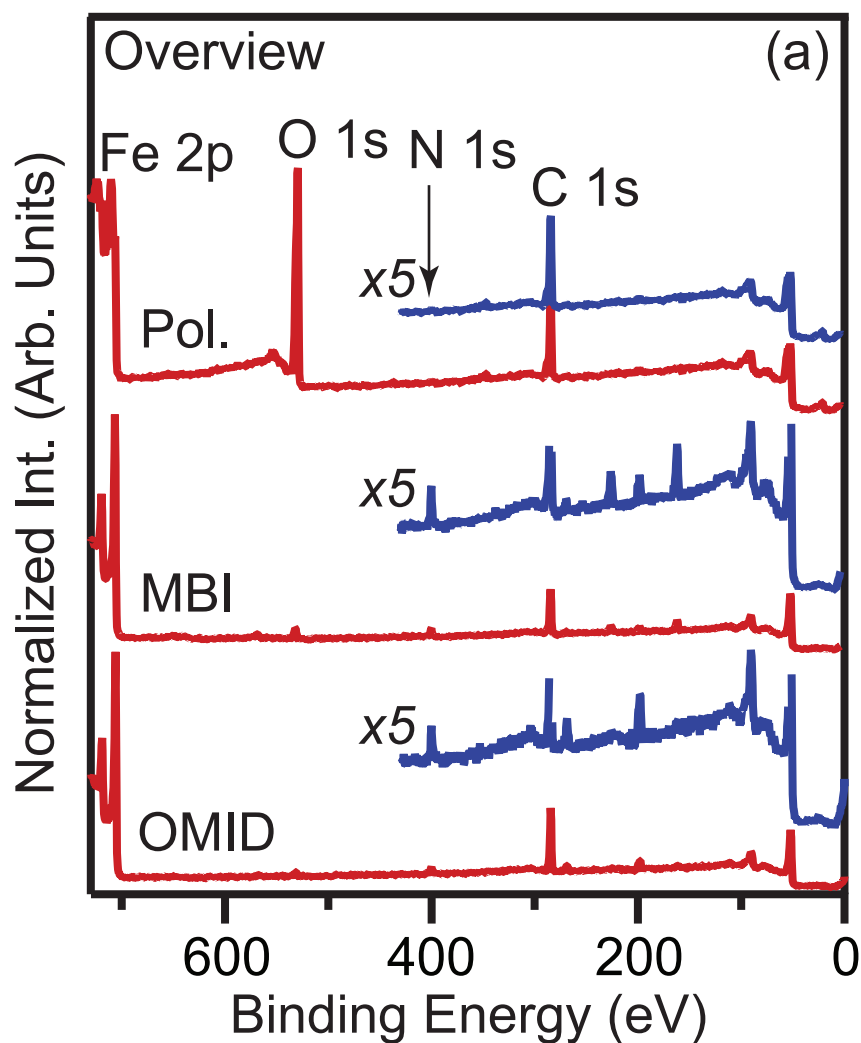
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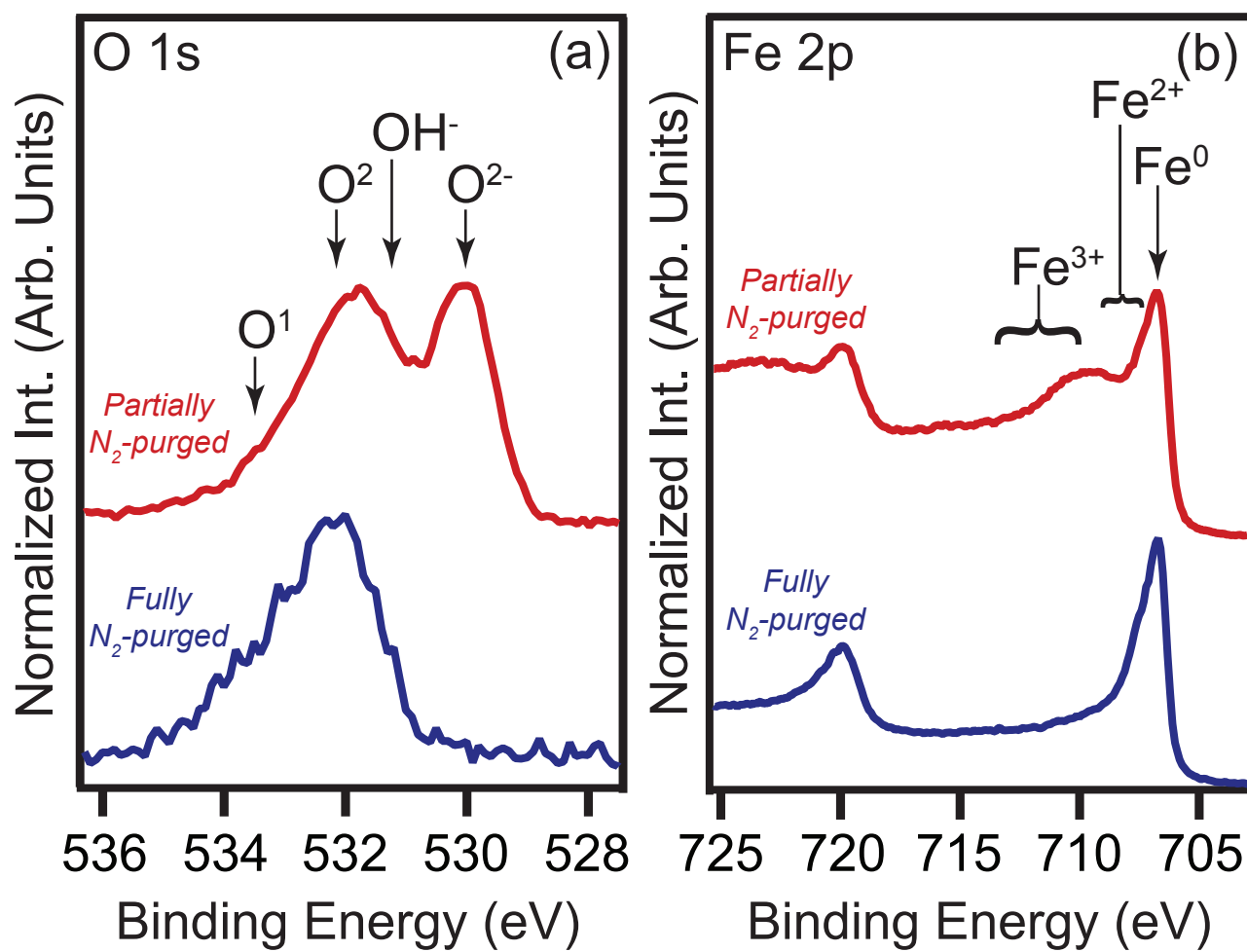
The authors declare that they have no competing financial interests.

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Name of Material/ Equipment	Company	Catalog Number
Mild steel rod, 1m x 10 mm diameter	RS Components	770-412
Silicon Carbide Grinding papers	Spectrographic Limited	T13316, T13317, T13318, T13156, T13153
Polishing Cloth	Spectrographic Limited	62113
Monocrystalline Diamond compound	Spectrographic Limited	G22003
OmegaPol TWIN 250mm Metallurgical Polisher	Spectrographic Limited	n/a
BRILLANT 220 - Wet Abrasive Cut-Off Machine	ATM GmbH Advanced Materialography	n/a
Ultrasonic Bath	NICKEL-ELECTRO LTD.	SW3H
Vacuum Desiccator	DURAN	24 782 57
Low form beaker 25mL	Fisher Scientific	FB33170
N2 gas cylinders, 230 bar, size W, UN1066	BOC	n/a
Regulator for N2 gas cylinders	Freshford Ltd.	MS-10B-N2
Nitrogen Purge Glove Box	Terra Universal, Inc	n/a
Dual Purge System	Terra Universal, Inc	1606-61
NitroWatch System with sensor	Terra Universal, Inc	9500-00A, 9500-02A
Big Digit Hygro-Thermometer	FLIR Commercial Systems, Inc. Extech Instruments Division	445703
Kratos Axis Ultra (XPS spectrometer)	Kratos Analytical Ltd	n/a

**Comments/Description**





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Determining the chemical composition of corrosion inhibitor/metal interfaces..

Author(s):

M. Walczak, P. Morales-Gil, T. Belashehr, K. Kousar, P. Arellanes Lozada, R. Lindsay

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Institution:	University of Manchester		
Article Title:	Determining the chemical composition of corrosion inhibitor/metal interfaces..		
Signature:	R Lindsay	Date:	23/6/16

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August 27, 2016

JoVE

**Re: Manuscript Resubmission**

Dear Editor,

On behalf of all the authors, I would like to thank the reviewers for their careful consideration of our manuscript, entitled “*Determining the Chemical Composition of Corrosion Inhibitor/Metal Interfaces with XPS: Minimizing Post Immersion Oxidation*”. We have addressed their comments, along with the editorial comments (see below for our detailed responses). Alterations to the text are highlighted by red, bold text in the revised version of the manuscript (with track changes also on).

Sincerely,

Rob Lindsay

## Editorial Comments/Responses:

1. Formatting: Please remove references to the video.

The reference to the video has been removed (line 103).

2. Grammar: -Please use American English throughout the manuscript, including figures. For instance, “normalised” should be “normalized”.

We ensured ‘Language’ was set to American English in Word to check for suitable spelling. Also, have changed ‘Normalised’ to Normalized in figure labels.

3. Line 91 – “a highly surface sensitive spectroscopy” – delete “a” and “highly”; if you mean “highly sensitive, “surface” needs to be moved.

The intention is to indicate a spectroscopy where the vast majority of the detected signal comes from the surface region (top few nanometers). On this basis, have changed text to ‘*a surface sensitive technique*’. This is a commonly accepted description of XPS, and so we believe it is appropriate to use it here.

4. 1.2.1 – “fume cupboard” – should be “fume hood”.

We have replaced ‘fume cupboard’ with ‘fume hood’.

5. 3.2.13, 3.2.14 – How is this performed? Manually or via software?

These operations are carried out manually. Have modified 3.2.13 and 3.2.14 to reflect this approach.

6. 4.1, 4.6, 4.7 – How is this done? Manually or via software?

For 4.1 and 4.6, a keypad is used to drive the sample manipulator motors. The text has been edited to reflect this procedure.

For 4.7, the XPS data acquisition software is used to acquire the XPS spectra. Have modified text to reflect this action more clearly.

7. Branding: 4.2 – Vision Manager

We have removed reference to Vision Manager from 4.2.

8. Discussion: Please discuss any future applications of the protocol.

A final paragraph has been added to the Discussion to address this issue, i.e.

Finally, given the effectiveness of the approach outlined here, we expect it to be applied to other topics in corrosion (i.e. in addition to corrosion inhibition), where acquisition of XPS data from air-sensitive interfaces would add to understanding. Moreover, such an approach should be considered in other fields where XPS measurements are undertaken from air-sensitive interfaces formed in a fluid environment. Clearly, this procedure is not restricted to XPS, but could also be applied to any other UHV-based measurement from a surface that has previously been immersed in a liquid, e.g. scanning probe microscopy.

9. For figures and tables that have been published before, please include phrases such as “Re-print with permission from (reference#)” or “Modified from..” etc. And please send a copy of the re-print permission for JoVE’s record keeping purposes.

Figure 2 is a significantly modified version of Figure 5 in Ref. 9. Have added text to figure caption to acknowledge this point.

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DOIs are included for all references where a DOI is available. Could not find DOIs for Refs. 1, 2 and 6.

## Reviewers' Comments/Responses:

### Reviewer #1:

1. Page 3 line 118-119, authors wrote: »Employ inhibited cooling fluid during the cutting procedure to mitigate any corrosion«. This is not due to corrosion, but not to induce additional mechanical damage.

We agree with the author that this is the purpose of the cooling fluid, and have modified the text accordingly.

2. -Page 3, line 128: why in acetone. Is that really necessary? Any reference for that?

We have no specific reference for this procedure. It is our usual procedure to ensure, as far as possible, that the substrate surface is contaminant free. We have not systematically explored other solvent combinations, but are satisfied that our procedure is fit for purpose. We note that, from empirical experience, the acetone is not strictly necessary.

3. -What is the composition of carbon steel?

The following sentence has been added to the Representative Results section (lines 270-271) to state the composition of the carbon-steel:

The carbon-steel possessed a nominal weight% composition of C (0.08-0.13), Mn (0.30-0.50), P (0.04), S(0.05), and Fe (balance).

### Reviewer #2:

1. I am quite impressed with the success of the method in avoiding oxide build up during sample transfer. This results, I think, from maintaining a reacting surface right up to the point that a vacuum is drawn. This probably differs from earlier attempts to maintain a representative reactive surface where too much time is spent washing the sample in other solvents. I wonder if the authors agree. Perhaps there could be some discussion of this in the ms? To see that if this hypothesis is so, then it would be a good idea to minimize the sample area since acid vapors would be drawn into the pump and would eventually degrade the oil.

Our initial reason for avoiding post-immersion washing of the substrate was to preserve the inhibitor coverage and structure as far as possible. We were concerned that solvent washing was more likely to remove adsorbed corrosion inhibitor. However, we do agree with the reviewer that oxidation may also occur during washing, especially if the solvent(s) used have not been de-aerated. We

have not tested this hypothesis, and so have no experimental evidence. To ensure that readers are aware of our thinking behind our specific procedure we have added a Note to 3.2.5:

**Note: Rinsing with solvents at this stage is not undertaken to minimize the potential of damaging the inhibited interface, e.g. inhibitor removal or interface oxidation.**

We also agree with the reviewer that acid vapors may reduce the life of the load lock pumps. As the reviewer suggests, a smaller sample may be reduce such damage. However, if it is too small sample handling becomes increasingly challenging, especially as the glove-box gloves are being used.

2. I know that the research results are secondary in this ms. However, it would be nice to see detailed S 2p and Cl 2p spectra to determine the nature of the bonding of these species to the metal.

As the reviewer implies, the purpose of the current paper is to detail the experimental procedure, so that others may employ it, and to show some representative results to demonstrate it utility. In our opinion, this goal has been achieved without presenting the S 2p/Cl 2p spectra. These spectra for MBI have been presented previously (see Ref. 9, including Supplementary Data). As regards OMID, we are preparing a manuscript to discuss this system, and prefer to not present/comment further on these data.