



Applications of scanning electrochemical microscopy (SECM) for local characterization of AZ31 surface during corrosion in a buffered media

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ABSTRACT

Different modes of scanning electrochemical mapping (SECM) such as surface generation/tip collection (SG/TC), amperometry, AC-SECM and potentiometry were employed to characterize the active/passive domains, hydrogen gas (H_2) evolution and local pH on a corroding surface of AZ31 in simulated biological fluid (SBF). It was found that the main domains of H_2 evolution are associated with lower insulating properties of the surface as well as higher local pH. The near surface pH was found to be highly alkaline indicating that, even in a buffered solution such as SBF, the local pH on a corroding AZ31 surface can be significantly different to the bulk pH.

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1. Introduction

Magnesium and its alloys have gained increasing attraction in recent years as biodegradable, biocompatible materials with appropriate mechanical properties [1,2]. However, rapid degradation of magnesium in the form of localized corrosion in biological environments is a serious drawback that can significantly reduce the *in vivo* service life [3]. For example it has been shown that the stent loses its mechanical integrity and load bearing characteristics much quicker when the corrosion is highly localized in contrast to a uniformly corroding specimen [4]. Therefore, it is crucial to develop techniques to study corrosion profiles in biological systems. Also, despite the advancements in understanding the corrosion behavior of Mg and its alloys in simple chloride containing media, there is still essential need for better understanding of corrosion mechanism in physiological environments [5].

Local corrosion behavior of magnesium alloys has been studied using scanning techniques such as localized EIS [6–8], scanning vibrating electrode technique [9,10] and scanning Kelvin probe

[11,12]. Among the methods used to enable local probing of microscopic processes, scanning electrochemical microscopy (SECM) has the unique capability to recognize active/passive regions and allows for surface characterization with a resolution in the micrometer range or below [13]. Electrochemical data may be collected in three main modes; namely amperometric, potentiometric and AC modes. The amperometric and potentiometric modes have been used to study the passive layers [14,15] and organic films [16–19] on metal and defects and pitting [20–22] as well as sensing local pH on a corroding metal [23]. The more recently introduced AC mode has proven to be useful for studying the insulating/conducting domains as well as changes in electrolyte composition near a corroding metal surface [24–26]. The AC-SECM signal is most responsive to the changes in solution resistance at high frequencies, while at lower frequencies the capacitive/resistive behavior of the surface and the electrical double layer are the major contributions to the impedance [24,27].

A particular area of interest in recent years has been bio-corrosion and associated electrochemical changes on the surface of Mg alloys in buffered solutions. However, a survey of the literature returns a limited number of published studies on characterizing Mg alloy surfaces using local probing techniques. Despite the unique capabilities of SECM in this area, it has not thus far been used for studying the corrosion of Mg in such environments. In this paper SECM is utilized to characterize the local corrosion

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processes, in real time, on the surface of AZ31 Mg alloy in simulated biological fluid (SBF). The aim of this work is to explore the use of SECM in studying the corrosion of magnesium with an emphasis on its applications in biological systems.

2. Experimental

2.1. Materials and sample preparation

Simulated biological fluid (SBF) was prepared using analytical grade reagents consisting of 5.403 g/l NaCl, 0.504 g/l NaHCO₃, 0.426 g/l NaCO₃, 0.225 g/l KCl, 0.230 g/l K₂HPO₄·3H₂O, 0.311 g/l MgCl₂·6H₂O, 0.8 g/l NaOH, 0.293 g/l CaCl₂, 0.072 g/l Na₂SO₄ and 17.892 g/l HEPES (4-(2-Hydroxyethyl)piperazine-1-ethanesulfonic acid – C₈H₁₈N₂O₄S) as buffer agent. The pH was adjusted to 7.40 ± 0.05 using 1 M NaOH (Sigma) solution. Ferrocenemethanol (FcMeOH) was analytical grade from Sigma. Mg alloy AZ31 (3% Al, 1% Zn and 0.3% Mn, all in wt%) was supplied by Goodfellow in the form of a sheet 2 mm thick. Specimens (5 mm × 5 mm) were abraded using 600, 1200 and 4000 grit emery consecutively and cleaned with isopropanol before exposure to electrolyte.

2.2. Instrumentation

Scanning electrochemical microscopy (SECM) was performed using CH Instruments SECM model 920D utilizing CHI integrated software version 12.26. Pt disk ultra-micro-electrodes (UME) (10 µm and 25 µm dia), were purchased from CH Instruments Inc., and were used as working electrodes. Pt mesh and Ag/AgCl (3.0 M NaCl) were used as counter electrode (CE) and reference electrode (RE), respectively. In the manuscript all voltages are stated versus the Ag/AgCl reference electrode unless otherwise stated.

2.3. Experimental procedure

A SECM image in constant height mode was obtained by scanning the tip in the x-y plane and recording the tip function against tip location. Depending on the operation mode of the SECM, the tip function can be either current (in feedback and generation/collection modes), AC impedance (AC mode) or potential (potentiometry mode). A schematic representation of the SECM set-up used in this study is shown in Fig. 1. The distance between tip and substrate was adjusted to 5 µm using a digital microscope, (MEIJI Techno model MS50). A working area of 2000 µm × 2000 µm was examined in each specimen with the remaining surface masked with tape to eliminate interference. All experiments were performed at room temperature, ca. 22 °C, in solution open to air.

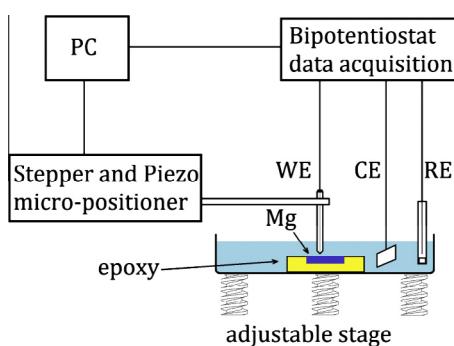


Fig. 1. Schematic representation of SECM set-up with Mg sample (mounted in epoxy) as substrate and Pt UME as working electrode (WE). In this study Pt mesh was used as counter electrode (CE) and Ag/AgCl was used as reference electrode (RE).

For the surface generated/tip collection (SG/TC) experiments a 10 µm diameter Pt UME was used and was biased at a voltage of 0.0 V. When the SECM was used in feedback mode data collection was performed using a ferrocene methanol (FcMeOH) redox mediator at a concentration of 4.0 mM prepared in SBF. The SECM probe was a 10 µm diameter Pt UME and biased at +0.4 V for oxidation of FcMeOH.

It is necessary to perform alternating current SECM (AC-SECM) in solutions with low ionic strength in order to maximize the sensitivity of the AC response [28]. In this study, SBF was diluted down by a factor of 100 and used for all AC-SECM experiments. All data collection was performed using a 25 µm diameter Pt UME. The UME diameter was increased in order to improve the signal/noise ratio. Approach curves were collected by moving the Pt UME from a distance of 75 µm above the Mg substrate down to <1 µm from the substrate surface. Approach curves were generated at the range of frequency from 75 kHz to 1 kHz in diluted SBF. The UME was held at OCP with an excitation AC signal amplitude of ±100 mV.

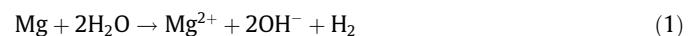
A solid-state pH sensing Iridium (Ir) UME was fabricated using a 16 µm diameter Ir wire insulated within a glass sheath (Fig. 2). The 16 µm Ir wire was produced by electrochemical etching a 75 µm diameter Ir wire (purchased from e-Filaments, LLC) at +10 V in 0.5 M H₂SO₄ for 20 min in a 3 electrode cell utilizing a Pt mesh auxiliary and Ag/AgCl reference electrodes. Iridium oxide (IrO_x) film was grown on the Ir tip by cycling the potential of the Ir UME between –0.25 V and +1.25 V at 3000 mV s^{–1} in 0.5 M H₂SO₄ electrolyte for 10,000 cycles [29] in a 3-electrode cell utilizing a Pt mesh and Ag/AgCl auxiliary and reference electrode respectively. The Ir/IrO_x UME was stored in 0.1 M NaCl overnight to avoid any influence of initial drift on the measurements [30].

3. Results and discussion

In addition to its high spatial resolution, SECM is best known for its versatility and application in different modes such as feedback mode to differentiate conducting/insulating domains or in generation/collection mode to study interaction with electroactive species on the surface. In this paper the technique is used in feedback, surface generation/tip collection (SG/TC), AC and potentiometry modes to gain information about local corrosion of AZ31 in SBF.

3.1. H₂ evolution studied in SG/TC mode

In SG/TC mode, H₂ generation from corrosion of Mg in neutral or basic solution (reaction (1) [31]) is sensed in an oxidation reaction (reactions (2) and (3)) at the UME probe potential of 0.0 V versus Ag/AgCl.



At higher pH values reaction (3) will dominate the hydrogen consumption at the UME. The potential at which reactions (2) and (3) take place is pH dependent as given by Eq. (4) (for H₂ at 1 atm pressure):

$$e_{\text{H}^+/\text{H}_2} = -0.059 \text{ pH} \quad (4)$$

At pH as high as 8.5, in close proximity to the Mg surface, the reduction potential of the hydrogen is –0.5 V versus SHE (~–0.7 V versus Ag/AgCl). Therefore, potentials more positive than –0.5 V SHE could theoretically be set as the UME tip potential to sense the generation

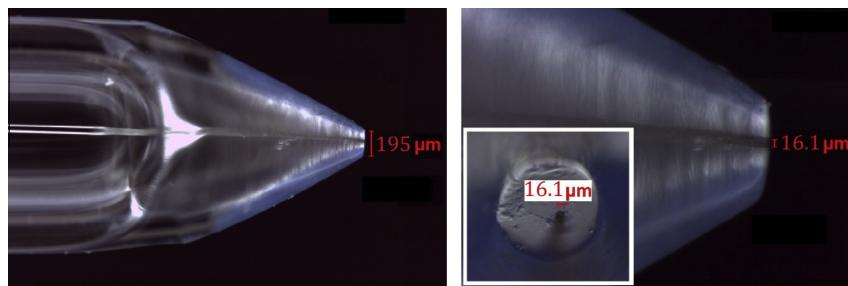


Fig. 2. Optical micrographs of the Ir UME fabricated by etching the Ir wire and insulation in glass sheath.

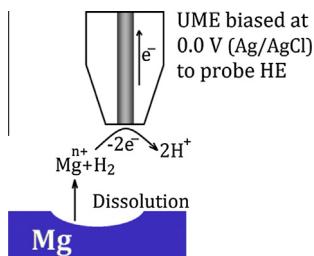


Fig. 3. Schematic principle of H_2 probing in SG/TC mode of SECM using a polarized Pt UME to oxidize H_2 over the corroding surface of Mg.

of H_2 . The schematic in Fig. 3 represents the principle of sensing H_2 at the UME biased at such potential.

Using the 10 μm UME cyclic voltammograms were recorded in SBF (Fig. 4) with and without the presence of a Mg sample. The CV recorded with the UME positioned 8 μm from the Mg surface exhibited a peak oxidation current centered at -0.3 V which is attributed to the oxidation of the hydrogen (H_2) generated at the Mg surface due to corrosion (reaction (1)). No oxidation peak is observed in this region when a CV is recorded using the UME in SBF only (no Mg sample present – SBF background) confirming that the peak recorded with the Mg sample present is due to H_2 oxidation. Also it is observed that the current was significantly reduced when the UME is positioned at long (2000 μm) distances from the

Mg substrate. This decrease is due to the lower concentration of generated H_2 at long distances from the substrate surface.

The CV of UME in fresh SBF (Fig. 4 – SBF background) shows a significant increase of negative (reduction) current at potentials more negative than -0.1 V which is related to reduction of dissolved O_2 in the SBF. Under standard conditions, at potentials negative of -0.2 V the reduction of O_2 ($\text{O}_2 + 2\text{H}_2\text{O} + 4\text{e}^- \leftrightarrow 4\text{OH}^-$) occurs in solution open to air [32]. Mapping the O_2 has been used previously in corrosion studies of metals [32,33], however it is known that O_2 plays a minor role in corrosion of magnesium [31] and therefore should not be considered in SG/TC mode for studying Mg corrosion. In order to avoid any potential interference from O_2 reduction, a UME potential more negative than -0.1 V was not considered in this work. Although the CVs recorded in the presence of Mg specimen (Fig. 4) does not show negative (reduction) current in the range of -0.1 V to -0.7 V , it is thought that the O_2 reduction current is masked by the strong positive oxidation current of H_2 . Therefore, in order to eliminate any contribution of dissolved O_2 to the tip current, a potential range of -0.1 to $+0.2\text{ V}$ was considered appropriate for mapping the H_2 evolution (solid box in Fig. 4). At potentials more positive of $+0.2\text{ V}$ the current generated from the oxidation of generated H_2 decreases sharply. Further investigation showed good resolution and sensitivity towards H_2 oxidation at the UME probe positioned close to the Mg surface when the UME was biased at 0.0 V and therefore the SECM experiments in SG/TC mode were performed at this potential.

Hydrogen collection is one of the most trusted methods of measuring the corrosion rate of magnesium and its alloys in aqueous media [34,35]. The technique produces quantitative values of corrosion rate based on cumulative hydrogen evolved due to the corrosion of magnesium (see reaction (1)). Using SECM in SG/TC mode presents an alternative way of studying the corrosion rate employing a local approach. Due to the complexity of Mg corrosion mechanism, bulk electrochemical techniques, such as potentiodynamic polarization, have been found to produce erroneous results in many cases by not taking into account the chemical dissolution of Mg [36]. The SG/TC mode of SECM allows for instantaneous measurement of the H_2 evolution as a direct measure of corrosion rate and has a great potential as a qualitative method for comparative studies. There has also been ongoing debates in the literature over the source of hydrogen evolution during the corrosion of magnesium [37–40]. It is believed that the SG/TC mode of SECM for probing H_2 evolution introduces new possibilities to gain better insights into the degradation mechanism of magnesium.

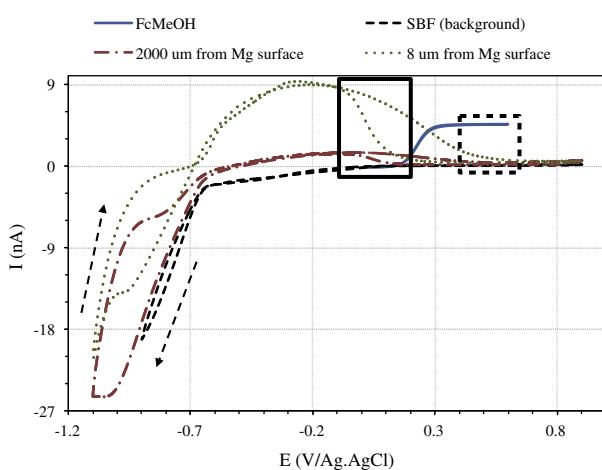


Fig. 4. Cyclic voltammograms of a 10 μm Pt UME in SBF showing the redox behavior of the redox mediator FcMeOH compared with the probe current generated by H_2 oxidation. Solid line box represents the potential window in which H_2 evolution can be measured using SG/TC mode whilst the dash box represents the potential window in which the FcMeOH redox mediator can be detected in feedback mode. SBF background was obtained with fresh SBF and no Mg present in the solution. The concentration of FcMeOH in SBF was 4.0 mM.

3.2. Conducting/insulating domains studied in feedback mode

Data collection in feedback mode involves the use of a redox mediator that changes its oxidation state in contact with the appropriately biased UME tip and substrate. In the vicinity of a suitably biased conductive or electrochemically active surface, the mediator returns to the original oxidation state resulting in

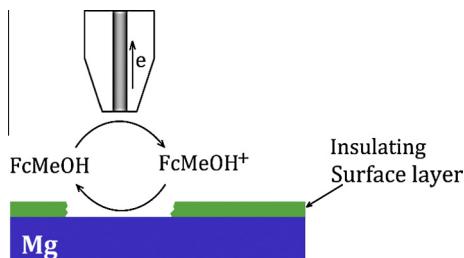


Fig. 5. Schematic representation of SECM probe in feedback mode of operation showing redox activity of ferrocenemethanol (FcMeOH) in contact with Mg substrate and cathodically biased Pt UME.

higher concentration of detectable mediator species at the UME tip and higher feedback current. This effect is called the “positive feedback” effect. Conversely, in the vicinity of an insulating surface, the diffusion of mediator from the bulk solution is hindered by the insulating material on the substrate around the UME tip, resulting in a decrease in feedback current. This effect is called “negative feedback” effect. **Fig. 5** illustrates the principle of positive feedback effect using ferrocenemethanol (FcMeOH) as the redox mediator in contact with the Mg surface.

In the bulk solution, because of the small dimension of the microelectrode, mass transfer is controlled solely by hemispherical diffusion of the redox mediator towards the UME tip. At a set distance sufficiently far from the Mg surface (i.e., in the bulk solution) the steady-state current measured at the UME (I_{∞}), is proportional to bulk concentration of redox mediator (C^{∞}), diffusion coefficient of the redox mediator (D), and number of electrons involved in the redox reaction, (n), according to Eq. (5) [41] where a is the UME radius and F is the Faraday constant.

$$I_{\infty} = 4nFDC^{\infty}a \quad (5)$$

The redox mediator, in order to be effective for SECM feedback mode imaging, must have fast redox kinetics and display minimal interaction with the active substances on the Mg surface to minimize interference of the corrosion processes. In studying the corrosion of Mg, it is important to make sure that the redox kinetics of the mediator is not influenced by the high pH and Mg^{2+} dissolution in close vicinity of the surface. In the feedback mode imaging of Mg corrosion, it is also crucial to avoid using UME bias potentials at which H_2 is oxidized. These conditions will ensure that only the feedback current produced by the mediator is recorded. A dashed box in **Fig. 4** identifies the potential window at which the redox mediator (FcMeOH) is oxidized at the UME with no interference from H_2 evolution (region designated by the solid line box in **Fig. 4**). There was no change in the bulk steady-state current (I_{∞}) recorded in solutions of SBF containing 4 mM FcMeOH at various pH (adjusted using NaOH) and Mg^{2+} concentrations (adjusted using $MgCl_2$) indicating that this mediator is not influenced by changing pH or Mg^{2+} concentration. Ferrocene undergoes a one-electron oxidation at potentials positive to +0.2 V, as shown by the positive current in **Fig. 4**. This reversible reaction serves as an electron transfer mediator for collecting the feedback mode data on the Mg surface. In order to eliminate interference from H_2 oxidation, potentials more positive than +0.4 V were used for feedback imaging.

Fig. 6a and **b** shows the H_2 map and conducting/insulating domains on the surface of the same AZ31 sample after 1 h immersion in SBF acquired using SG/TC and feedback modes, respectively. The upperscale colors (magenta and blue¹) on **Fig. 6a** indicate the areas with high H_2 evolution rate whereas the lower scale colors (red and yellow) represent the relatively passive areas with low H_2

evolution. In **Fig. 6b**, the upper scale colors indicate the areas with higher feedback current (i.e., higher conductivity regions) versus the lower scale colors that represent areas with lower feedback current (i.e., lower conductive regions). These lower conductive regions are most likely due to the formation of an insulating surface layer. Comparing **Fig. 6a** and **b**, relatively higher activity/conductivity is observed at (but not limited to) the main domains of high H_2 evolution. This result is consistent with the micro-galvanic corrosion mechanism in buffered media [42] and chloride containing solutions [35] caused by intermetallic particles. It has been shown that intermetallic particles and secondary phases, when isolated, may increase the corrosion rate of Mg by forming micro-galvanic corrosion cells [43,44].

In the context of corrosion of Mg, the feedback mode of SECM can be particularly useful in obtaining information about the formation and breakdown of passive surface layers. It is well known that corrosion inhibition of Mg alloys to a large extent depends on the formation and stability of the formed passive layer. Here it is postulated that the local formation of a surface layer when immersed in SBF slows down the corrosion rate and therefore reduces the generation of H_2 .

3.3. Capacitive/resistive characteristics studied in AC mode

AC-SECM imaging comprises of recording impedance (imaginary and real) and phase shift via counter electrode (CE) as a response to the AC signal applied at the UME at a set frequency and excitation amplitude. **Fig. 7** illustrates the electrical model for SECM data acquisition in AC mode at conducting and insulating domains of the Mg surface [45].

Prior to performing AC-SECM in diluted SBF each Mg sample was immersed in standard SBF (undiluted) solution for 1 h. Approach curves were obtained by moving the UME towards the Mg surface whilst holding the UME at its open circuit potential and measuring the resulting impedance. An AC potential (± 100 mV) was applied onto the OCP and each approach curves were obtained at a set frequency to determine the response of the surface film as a function of the frequency (**Fig. 8**). Normalized impedance was calculated using Eq. (6):

$$\text{normalized impedance} = Z_L/Z_{\infty} \quad (6)$$

where Z_L is the impedance at distance L from the substrate and Z_{∞} is the bulk impedance at 75 μ m distance from the Mg substrate. Normalized tip-substrate distance was calculated as;

$$\text{normalized tip} - \text{substrate distance} = L/D \quad (7)$$

where L is the actual tip-substrate distance in μ m and D is the tip diameter.

Fig. 8 shows the normalized approach curves in which a positive feedback (i.e., increasing normalized AC impedance as the UME approaches the Mg surface) was observed at lower frequencies ($f \leq 15$ kHz) and negative feedback (i.e., decreasing normalized AC impedance as the UME approaches the Mg surface) was observed at higher frequencies ($f > 42$ kHz). In general, a positive feedback effect is observed when the frequency of the applied AC signal to the UME is smaller than the reciprocal of the time constant for the interfacial impedance ($f < \frac{1}{2\pi RC}$), whereas negative feedback is observed for higher frequencies [26]. As a surface film grows on the bare Mg surface, the time constant associated with the surface film increases, resulting in a lower frequency at which the switch from positive to negative feedback occurs [26]. Thus, the frequency at which a distinct variation between bare Mg surface and Mg surface covered with a film is observed will decrease. Approach curves shown in **Fig. 8** suggest that adequate sensitivity of AC signal to insulating properties of the Mg surface layer is obtained at low frequencies, namely 1 kHz.

¹ For interpretation of color in Figs. 6 and 9, the reader is referred to the web version of this article.

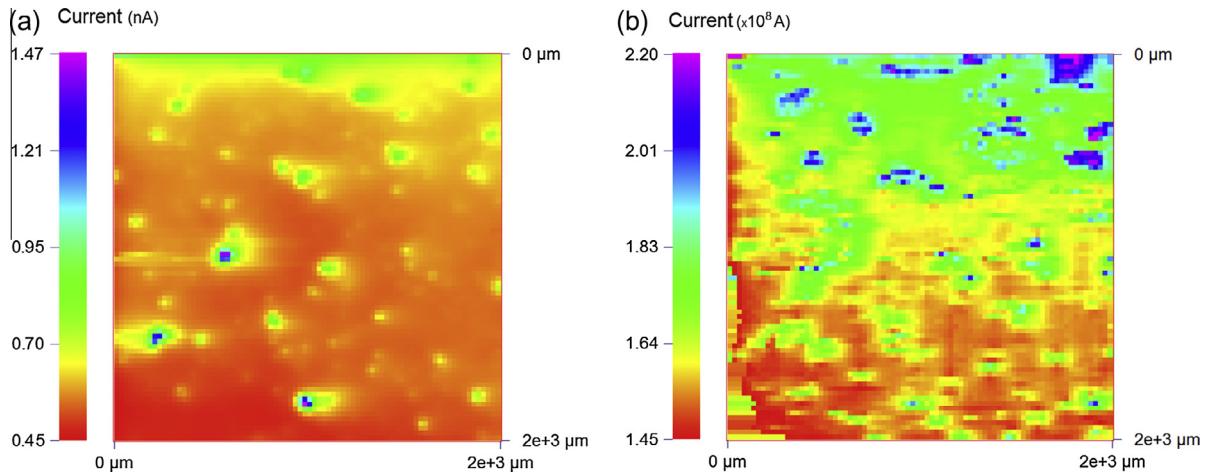


Fig. 6. SECM images of AZ31 surface in SG/TC (a) and feedback (b) modes acquired via sensing the H_2 evolution and FcMeOH (4 mM) regeneration. The tip potential in feedback mode was 0.55 V (versus Ag/AgCl) and in SG/TC mode was 0.0 V (versus Ag/AgCl). Both images were acquired on AZ31 surface after 1 h immersion in SBF with 10 μm Pt UME.

As indicated by the approach curves (Fig. 8), good resolution with high sensitivity towards the surface layer was observed at 1 kHz with 100 mV amplitude of excitation voltage. A similar frequency has been used elsewhere for imaging pitting corrosion [46,47]. Fig. 9a shows the impedance map from AC-SECM taken from the same area that was scanned in SG/TC and feedback modes (Fig. 6a and b respectively). Upper scale colors in Fig. 9a (magenta and blue) represent the relatively low conductivity areas versus lower scale colors (red and yellow) that indicate the more conductive areas. Fig. 9b shows the optical micrograph of the sample after the AC-SECM experiment. It is notable, by comparing Figs. 9a and 6a, that the surface layer is significantly more conducting at most sites of active H_2 evolution. Furthermore, from Fig. 9a we observe that very low conductive domains exist in close proximity to some active H_2 evolution sites.

The increase of conductivity at the domains of higher H_2 evolution (comparing Figs. 9a and 6a) could be attributed to the increase of ionic strength within the gap between the sample and the UME due to dissolved Mg^{2+} metal ions and/or the removal of the surface film from the Mg sample thus exposing the underlying metal. The porosity of the surface film results in a positive AC-feedback effect at both corroding and covered areas [26]. These domains are distinguishable by their difference in the strength of positive feedback signal indicating that even the areas covered by a resistive surface film are not perfectly insulated (i.e. the surface film is porous) allowing the AC signal to pass through.

When the AC mode results are combined with SG/TC mode data, it reveals that the Mg is not covered by a surface film in the domains of high H_2 evolution where anodic dissolution of Mg takes

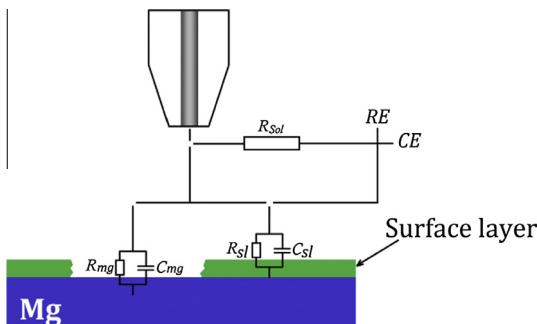


Fig. 7. Schematic of the electrical model of AC mode data acquisition at conducting and insulating regions of an Mg surface.

place [48]. The feedback mode results identified these areas with increased activity and enhanced electron transfer properties. This suggests that anodic dissolution of Mg is either accompanied by H_2 evolution or it takes place at very close vicinity to isolated cathodic areas, presumably Al–Mg intermetallics. Preferential anodic dissolution of Mg against Al–Mg intermetallics has been shown elsewhere [49].

Here the AC mode of SECM is shown to be a useful method for studying resistive properties of the surface layer that forms on the surface of Mg during corrosion. While the low, 1 kHz, frequency scan distinguish between domains of lower conductivity and those of higher conductivity, more information about capacitive properties of the surface layer at a certain area may be achieved through AC approach curves (Fig. 8) over a range of frequencies. The main advantage of the AC mode over the other feedback modes is the elimination of the redox mediator that in some cases may interfere with the natural corrosion process or may react with active/passive domains on the surface. Although not attempted here, AC mode SECM has shown potential for studying the surface profile *in situ* at high frequency ranges [27,50].

3.4. pH sensing in potentiometric mode

Potentiometric mode is a two electrode configuration of SECM in which the open circuit potential of UME tip is measured against

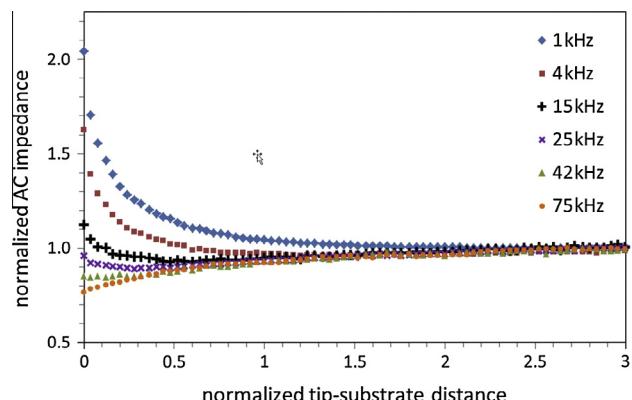


Fig. 8. Normalized approach curves at the range of frequency from 75 kHz to 1 kHz towards the surface of AZ31 in diluted SBF. AZ31 was at open circuit potential. The probe was a 25 μm diameter Pt UME held at OCP with excitation signal amplitude of ± 100 mV.

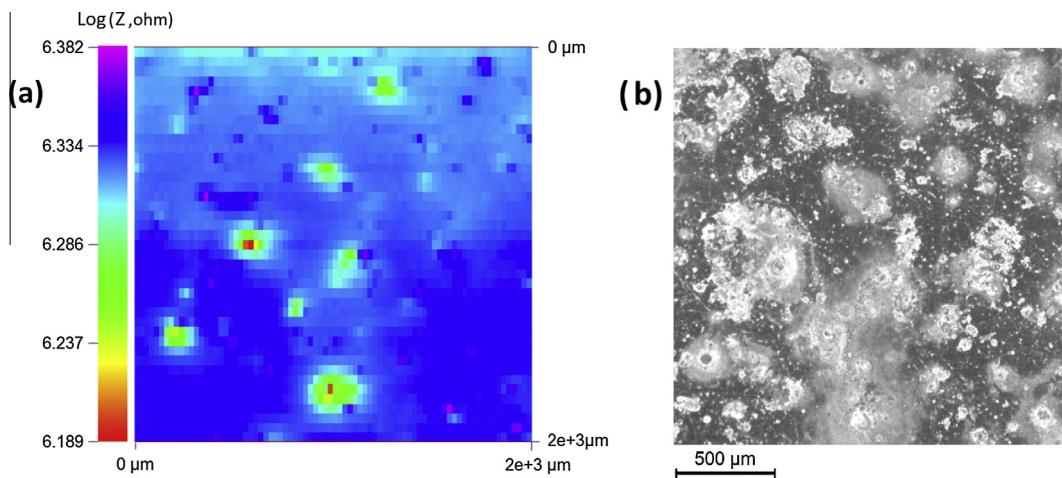


Fig. 9. AC-SECM image (a) and optical micrograph (b) of AZ31 after 1 h immersion in SBF. AC mode experiments were performed in diluted SBF using a 25 μm Pt UME with excitation signal amplitude of 100 mV at 1 kHz.

the reference electrode as a function of tip location and can be used to map the pH near the surface of a sample surface. In order to do this a calibration curve of OCP versus solution pH must be prepared. Therefore, SBF solutions of varying pH were prepared using fresh PBS and 0.1 M HCl and 0.1 M NaOH. These solutions were used to generate a calibration curve (Fig. 10a). A similar method was used to sense pH on Al-Zn in un-buffered media by Lowe et al. [30]. The possible interference of H_2 was ruled out by measuring OCP in SBF solutions with and without dissolved H_2 (from a corroding Mg sample) without any variation in OCP (at a fixed pH) observed.

In order to measure local pH (Fig. 10b), a H_2 map was first collected using the SG/TC mode (Section 3.1). The H_2 map was then used as a guide to find the domains with different levels of H_2 evolution. The Ir/IrO_x UME was then moved precisely above the points of interest and OCP was measured against Ag/AgCl at approximately 30 μm distance from the Mg surface. The calibration curve (Fig. 10a) was used to convert the OCP values to pH. The pH of bulk buffered solution was measured and remained at 7.6 during these experiments. As illustrated in Fig. 10b, the near surface pH is significantly higher than that of bulk; with local pH at some sites of H_2 evolution reaching as high as 11.9.

The near surface pH plays an essential role in determining the formation and stability of the surface layer on Mg. It is well known that most magnesium hydroxide and calcium phosphate layers

that form on the surface of Mg are stable at high pH and tend to break down at low or neutral pH [5,51]. It is believed that near surface pH measured by Ir/IrO_x UME in this study can provide valuable information that helps better understand the formation of a surface layer of Mg alloys in physiological media.

4. Conclusion

Four modes of SECM operation (i.e. SG/TC, feedback, AC and potentiometry) were used for *in situ* characterization of Mg surface corrosion in a biological buffered solution. Combining the variables measured in different modes of SECM provides new insight into the complex corrosion mechanism of Mg in a buffered media. The H_2 probing performed using SG/TC provides a direct measure of the spontaneous corrosion rate on a local scale. H_2 evolution is sensed in an electro-oxidation reaction at the SECM probe producing a current that increases by increasing local concentration of H_2 . Using the feedback mode it was found that the Mg surface is more active (in the context of electron transfer) at the domains of high H_2 evolution. The AC mode of SECM shows promise in probing the areas that are covered with an insulating surface film, with high sensitivity at low AC frequencies. Complementary to the SG/TC and feedback modes, the AC mode results revealed that Mg is not covered by a surface film in the domains of high H_2 evolution where hypothetically anodic dissolution of Mg takes place. This

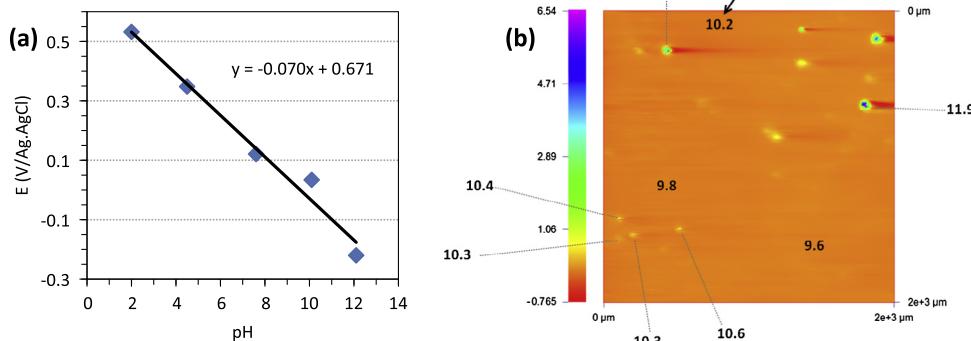


Fig. 10. Calibration curve correlating Ir/IrO_x potential to the pH of solution (a) and the local (single point) pH values shown on a H_2 map (b). Ir/IrO_x UME was positioned precisely above the points of interest and OCP potential was recorded against Ag/AgCl. Individual pH values were converted from OCP values using the calibration graph.

implies that anodic dissolution of Mg either coincides with H₂ evolution or it takes place at very close proximity to isolated cathodic areas and is also in-line with the highly alkaline pH at these domains. The near surface pH was found to be alkaline despite the SBF bulk solution being buffered to 7.4. This has significant impact on the mechanisms involved in formation of surface passive layer as well as the cytocompatibility of magnesium. The regions of highest pH correlated with the major domains of H₂ evolution and lower surface resistivity.

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