In vitro corrosion of pure magnesium and AZ91 alloy—the influence of thin electrolyte layer thickness

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Abstract

In vivo degradation predication faces a huge challenge via in vitro corrosion test due to the difficulty for mimicking the complicated microenvironment with various influencing factors. A thin electrolyte layer (TEL) cell for in vitro corrosion of pure magnesium and AZ91 alloy was presented to stimulate the in vivo corrosion in the micro-environment built by the interface of the implant and its neighboring tissue. The results demonstrated that the in vivo corrosion of pure Mg and the AZ91 alloy was suppressed under TEL condition. The AZ91 alloy was more sensitive than pure Mg to the inhibition of corrosion under a TEL thickness of less than 200 μm. The TEL thickness limited the distribution of current, and thus localized corrosion was more preferred to occur under TEL condition than in bulk solution. The TEL cell might be an appropriate approach to simulating the in vivo degradation of magnesium and its alloys.

Keywords magnesium; corrosion; thin electrolyte layer; biomaterials

Introduction

Pure magnesium [1, 2] and its alloys such as Mg-Al [3], Mg-Zn [4], Mg-Mn [5], Mg-RE [6], Mg-Ca [7] and Mg-Li [8] are promising biodegradable biomaterials for cardiovascular stents [9] and orthopedic implants [10, 11] due to their outstanding biocompatibility, biomechanical compatibility and osteoinduction [12, 13]. The screening and in-service life span predication of newly developed magnesium implants require cost-effective in vitro and in vivo tests, especially for degradation tests at an early stage in their development [14].

Biodegradation refers to corrosion occurring on the magnesium replacements as exposed to body fluid [15]. Usually, the corrosion performance of magnesium and its alloys can be assessed using a variety of in vitro corrosion methods such as immersion mass loss, hydrogen evolution and electrochemical polarization tests together with electrochemical impedance spectroscopy in bulk simulated body solutions [16]. However, the above measures cannot well-evaluate the in vivo degradation rate of magnesium due to the complicated factors including material, chemical, mechanical and biological aspects.

To date, a considerable number of in vitro degradation investigations have been made, involving the influencing factors, such as inorganic ions [3, 17], proteins [18], amino acids [19], glucose [20], corrosion medium volume [21], extraction condition [22], blood flow [23], cells [24, 25], tissues [26], pH [27] and pH buffers[28]. The degradation rate is much bigger in saline solution than in E (Earle’s solution containing calcium and magnesium salts)-MEM (Eagle’s minimum essential medium)-FBS (fetal bovine serum), which is the closest solution to human blood plasma. Protein adsorption and insoluble salt (i.e., carbonate, phosphate, Ca-P) formation retarded magnesium degradation, whereas organic compounds such as vitamins, amino acids, glucose and albumin as well as cell [29] promote the dissolution of magnesium [18–20]. Also, the degradation rate is influenced by buffering solutions, which buffered with N-2-hydroxyethylpiperazine-N’-2-ethane sulfonic acid (HEPES) and NaHCO3 leads to an increase and decrease in the degradation rate, respectively [30]. The Earle’s balanced salt solution buffered with sodium bicarbonate provides a degradation rate comparable to those observed in vivo [31].
While the in vivo degradation of magnesium implants is characterized by mass loss based on the volume loss derived from the three-dimensional (3D) reconstruction using synchrotron radiation based micro-computed tomography (μ-CT) [12]. Nevertheless, this methodology has several limitations: (i) it cannot tell the corrosion product from the new bone, thus the volume of the corrosion products cannot be removed; (ii) the average mass loss rate cannot reflect the actual degradation rate due to pitting corrosion usually occurs on magnesium alloys and (iii) the degradation of the implanted magnesium alloys was suppressed by protein and cellular/tissue encapsulation [32] due to more difficult diffusion of the species in the microenvironment. Therefore, the degradation rate of magnesium obtained from μ-CT may be different, to some extent, from the in vitro corrosion tests [12].

Moreover, the in vivo degradation rate of Mg alloys varies with different implantation positions or anatomical locations in the body [26, 33, 35]. For instance, it is higher in the femur than in the muscle (subcutaneous tissue). The deposition layer resulted from the degradation of the alloy bar, positioned in the bone marrow cavity, suffers from the removal from body fluid [26] due to the faster corrosion rate and higher pH increase under the flow condition than the static condition [23]. This finding implies that the geometrical dimensions of the electrochemical cell have a significant impact on degradation behavior of the Mg implants. However, few in vitro studies refer to the influence of the implanted positions or the geometric dimensions of the electrochemical cell on degradation rates of the Mg replacements [1, 2].

Usually, the conventional in vitro corrosion tests are performed in bulk solutions, while the in vivo degradation occurs in the microenvironment on one hundred micrometer scale [34], constructed by the implant and its surrounding tissue during the bone-healing process. Namely, the microenvironment can be regarded as a thin electrolyte layer (TEL), the thickness of which is lower than a few hundreds of micrometers, achieved by mechanically confining the desired distance or thickness between the surface of working electrode (WE) and a parallel waterproof wall [35].

Since the conventional or traditional in vitro corrosion in bulk cell cannot be used to well-predict in vivo corrosion rates of magnesium alloys due to the ignorance of geometric factor of electrochemical cell [12], it is thus of importance to establish a novel in vitro methodology for evaluating the corrosion resistance of magnesium and its alloys [15, 36]. The new methodologies should genuinely reflect the influence of the tissue in the microenvironment on corrosion behavior of the orthopedic magnesium implants.

Fortunately, the TEL technique has been successfully applied to investigate the atmospheric corrosion of the alloys such as Fe-Ni [37], aluminum [38], magnesium [39, 40] and copper alloys [41, 42]. It is well known for atmospheric corrosion electrochemical reaction generally occurs between micro-anoles and micro-cathodes under TEL [39]. A great change in the TEL thickness markedly affects a considerable processing such as the potential distribution, mass transport and release of metallic ions and accumulation of corrosion products [37]. Also, it is observed that the hydration of dissolved metal ions, especially the accumulation of the secondary corrosion products [41]. This particular geometry results in specific galvanic coupling and local pH variation [37]. Chen et al. [43] claimed that the anodic and cathodic processes of the corrosion of Mg-10Gd-2Y-0.4Zr alloy were both retained under TELs. The corrosion was more localized under TEL than in bulk solution, despite the cathodic process still dominated by hydrogen evolution reaction. The decrease in corrosion rate of Mg-10Gd-2Y-0.4Zr alloy with the decrease in layer thickness, which is ascribed to the inhibition of the rate of pit initiation and the decrease in probability of pit growth.

In view of the specific corrosion behavior under TEL, we design an electrochemical cell to simulate the in vivo corrosion for magnesium and its alloys based on the fact that the orthopedic implants are covered with fibrous capsule, interstitial region and muscles [32, 44] and the interstitial region is filled with body fluids between the implant and the muscle, the thickness of the interstitial region maintains at dozens to hundreds of micrometers [32]. The body fluid layer is closely related to the surface of the degradable implants [12]. Therefore, the interstitial region can be simplified as a TEL model to mimic the in vitro microenvironment. The TEL tests may be an appropriate approach to mimic the in vivo degradation of magnesium and its alloys, on the basis of the results from TEL [35, 38, 45] and no match for corrosion rates of in vivo and in vitro tests [39, 42].

The purpose of this study is to apply the TEL cell for characterizing the in vivo corrosion rate of magnesium and its alloys and to open a new window for mimicking the in vitro corrosion test of magnesium and its alloys, concerning with the influence of geometric dimensions of electrochemical cell on degradation behavior.

**Experimental**

Figure 1a schematically illustrates the microstructure of an orthopedic implant and its neighboring bone. The implant was firstly threaded into the compact bone (white part), then embedded into the bone matrix (brown part) during the in vivo tests. And the top was covered by the fibrous capsule (not given in figure), the interstitial region (blue part) and the muscle (pink part). The interstitial region with a thickness of 50–200 μm was filled with body fluids and had a physical contact with the implant surface. The muscle would further hinder the mass transfer in the body fluids. Therefore, the in vivo microenvironment of the orthopedic implant can be idealized to a TEL model (Fig. 1b). The magnesium implant was the WE. The interstitial region and the muscle were simulated by the TEL and polytetrafluoroethylene (PTFE) hydrophobic film, respectively. The specifically designed structure could diminish the influence of the surface tension of the electrolyte to maintain a thinner electrolyte thickness than the open cell [39, 41], and the PTFE hydrophobic film maintained a stable TEL on the surface of WE (Fig. 1b).

Prior to testing, commercial as-cast pure Mg (with a purity of 99.92 wt. %) and AZ91 alloy (compositions: 8.5–9.5 wt. % Al, wt. 0.45–0.9% Zn and balanced Mg) were embedded in epoxy resin as the WE and ground to 2000 grit, then inserted into a special sample stage in the electrolytic cell setup, as shown in Fig. 1. A three-electrode system was applied. The WE, saturated calomel electrode as the reference electrode and a platinum wire as the counter electrode were connected with an electrochemical workstation (PARSTAT 2273, Princeton Applied Research). All the polarization measurements were performed in Hank’s solution at a scan rate of 1 mV/s at room temperature. The area of the WE exposed to the solution was 1 cm². Triplet measurements were conducted for each group.

The PTFE-made tube was fixed on the frame to restrain the insulating plastic tube moving up and down along the Z axis. Three horizontal adjusting screws were placed on the bottom of the electrolytic cell to ensure planar parallelism between the top surface of the sample and the bottom of the insulating plastic tube. The plastic tube and the sample stage constituted the core of the TEL cell, as shown in Fig. 2. A pair of levers was used to control the TEL thickness by regulating the height of the plastic tube on Z axis.
When the bottom of the plastic tube was dropped down on the top surface of the sample, the dial indicator reading was set as zero and then the plastic tube was lifted, the dial indicator directly designated the TEL thickness.

The open circuit potential (OCP) measurement firstly lasted 3 min in the bulk Hank’s solution and then the TEL thickness was adjusted to a specific value, and consequently, the electrochemical experiments were conducted in triplicates for each TEL thickness.

The samples were polished and etched by natal to disclose their microstructures prior to the electrochemical tests such that to understand the influence of microstructure, particularly the secondary phase on corrosion. And then a 3D contour profiler (Zeta-200, Zeta Instruments Inc.) was employed to discern the change in surface morphology of the pure Mg sample and the AZ91 alloy before and after the electrochemical experiments.

Results and Discussion

Figure 3a and b presents the cathodic polarization curves of the pure Mg sample at various TEL thicknesses. The curves of corrosion current density, $I_{corr}$ as a function of the TEL thickness experienced two different stages. The cathodic current density under TEL is smaller than that of the bulk solution. Namely, the $I_{corr}$ of pure Mg decreased with a decrease in TEL thickness (Fig. 3b), regardless of the reversed tendency for the 150-μm and 200-μm TELs. The result indicates that the TEL leads to the inhibition of cathodic process by the hydrogen reduction, which relates to the uneven distribution of current due to the reduced effective electrode area in the TEL in comparison with the bulk solution [38].

Interestingly, the cathodic polarization curves of pure Mg almost had close Tafel slopes (Fig. 3b) under various thicknesses of TEL to the bulk solution. The OCPs of the TEL, except for the 50-μm TEL,
were higher than that of the bulk solution. This scenario means the inhibition of anodic process probably results in the rise in OCPs. Generally, the TEL prohibited the cathodic and anodic processes of pure Mg in Hank’s solution. Generally, the $I_{corr}$ of pure Mg increased with an increase in TEL thickness (Fig. 3a and b) except for the exchange of the thickness of 150 $\mu$m and 200 $\mu$m. Similarly, with decreasing TEL thickness, the $I_{corr}$ of the AZ91 alloy reduced; and the OCPs was shifted toward more noble direction (Fig. 3c and d). The OCPs for the AZ91 alloy may be related to the formation of a corrosion product layer on the surface of the AZ91 alloy and the inhibition of anodic process [39].

Furthermore, electrochemical impedance spectroscopy measurements were conducted with TEL thicknesses of 50 $\mu$m and 500 $\mu$m. Figure 4a and b provided Nyquist and Bode phase plots of pure Mg and the AZ91 alloy at thicknesses of 50 $\mu$m and 500 $\mu$m, respectively. The Nyquist plot (Fig. 4a) designates that the TEL gave rise to a higher solution resistance, the scattered data at low frequency region also means the instability of the TEL and the extremely difficult diffusion process of the H$^+$ ions under such a TEL due to the influence of the uneven distribution of current [45]. It is suggested that the current distribution would be uneven if the phase angle was less than 45$^\circ$ [46], whereas the phase angle greater than 45$^\circ$ designates the electrolyte film is continuous on the surface of the samples and uniform distribution of current.

It is clear that the current distribution was uneven under 50-$\mu$m TEL (Fig. 4b) for both pure Mg and the AZ91 alloy due to the fact that the micro-galvanic corrosion between the impurity/Mg$_2$Al$_3$ phase and the $\alpha$-Mg phase was susceptible to the uneven distribution of current. On the contrary, the current distribution was more uniform under the 500-$\mu$m TEL condition.

Generally, the TEL inhibited, to some extent, the corrosion of both pure Mg and the AZ91 alloy, as shown in Fig. 5. When the thickness was not greater than 100 $\mu$m, the $I_{corr}$ of the pure Mg sample was higher than that of the AZ91 alloy. This scenario is due to the fact that more severe pitting corrosion occurred on the surface of the pure Mg samples rather than the AZ91 alloy (Fig. 6), on which no remarked change happened.

Moreover, the hydrogen reduction current density of pure Mg had not marked change when the TEL thickness is within 200 $\mu$m. But for the AZ91 alloy, there was a rapid increase in $I_{corr}$ at the electrolyte layer thickness of 150 $\mu$m. When the TEL thickness was greater than 200 $\mu$m, the $I_{corr}$ of the AZ91 alloy increased slowly with the increasing TEL thickness. That is, there is a linear relationship between the corrosion current density and the TEL thickness for the AZ91 alloy as the TEL thickness is not less than 200 $\mu$m. The scenario is attributed to the larger volume fraction and higher potential of the second phases (β phases). The big potential discrepancy between the β phase and the $\alpha$-Mg matrix led to

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**Figure 3.** cathodic polarization curves of (a, b) pure Mg and (c, d) AZ91 alloy under TEL condition with various thicknesses of electrolyte layer. The TEL with a thickness of 1000 $\mu$m is regarded as the bulk solution [37].
micro-galvanic corrosion between \(a\) and \(b\) phases in the AZ91 alloy. And the TEL caused the rapid formation of Mg(OH)\(_2\) corrosion product due to the fact that cathodic process was predominately controlled by diffusion of H\(^+\) ions, which was restricted by TEL. Thus, the corrosion of the AZ91 alloy was inhibited in comparison to the pure Mg sample.

Anodic reaction:

\[
\text{Mg} \rightarrow \text{Mg}^{2+} + 2e^- \quad (1)
\]

Cathodic reaction:

\[
2\text{H}_2\text{O} + 2e^- \rightarrow 2\text{OH}^- + \text{H}_2 \uparrow \quad (2)
\]

Total reaction:

\[
\text{Mg} + 2\text{H}_2\text{O} \rightarrow \text{Mg(OH)}_2 + \text{H}_2 \uparrow \quad (3)
\]

In this study, the corrosion current density measurement was only conducted from the cathodic branch of polarization curves to avoid the experimental errors caused by anodic polarization. Taking into consideration of the diffusion process, the cathodic current density \(I_c\) can be expressed by following equation [38, 47]:

\[
|I_c| = \frac{I_{\text{corr}} \exp \left( -\frac{\Delta E}{\beta_c} \right)}{1 - \frac{I_{\text{corr}}}{I_L} \exp \left( -\frac{\Delta E}{\beta_c} \right)} \quad (4)
\]

where \(I_{\text{corr}}\) is the corrosion current density, \(I_L\) is the diffusion limiting current density, \(\Delta E\) is the overpotential and \(\beta_c\) is the cathodic Tafel slope. According to Eq. (4), the cathodic current density \(I_c\) was determined by the diffusion limiting current density.

For a one-dimensional diffusion system, the theoretical diffusion limiting current density can be calculated according to the Nernst–Fick equation [39]:

\[
I_L = \frac{nFD_{\text{H}^+}}{\delta} \quad (5)
\]

where \(n\) is the number of the electrons involved in the hydrogen reduction reaction, \(F\) is the Faraday constant, \(D_{\text{H}^+}\) is the diffusion coefficient, \([\text{H}^+]\) is the concentration of H\(^+\) in the TEL and the \(\delta\) is the thickness of the diffusion layer. It can be seen from Eq. (5) that \(I_L\) is dependent on the value of both \([\text{H}^+]\) and \(\delta\), since the other factors are constants in same system. Generally, \(\delta\) is in the range of 10^\(-100\) \(\mu\)m [47]. Therefore, the influential factor to \(I_L\) should be discussed in two situations of TEL model:

1. When the TEL thickness \(d > 200\) \(\mu\)m (\(d > 2\delta\)), the diffusion layer of both corroded metal electrode and PTFE film are intact, the TEL model can be regarded as a one-dimensional diffusion system and the Nernst–Fick equation is suitable for this situation. Therefore, \(I_L\) is mainly dominated by \([\text{H}^+]\). Considering surface alkalinization resulted from dissolution of magnesium, the cathodic reaction produces hydroxyl and thus the local pH increases at cathodic sites [39]. So an increase in \([\text{H}^+]\) results in a lower \(I_L\). That is, a higher \(I_c\) for the same material according to Eq. (4), which fit for cathodic polarization curves in Region II.

2. When the thickness of TEL \(d < 200\) \(\mu\)m (\(d < 2\delta\)), the opposite effects of \([\text{H}^+]\) and \(\delta\) will result in a swing in the cathodic polarization for pure Mg. However, the micro-galvanic corrosion between the \(a\)-matrix and the \(b\) phase should be taken into consideration for the AZ91 alloy and it could not be regarded as a simple one-dimensional diffusion system. The \(I_{\text{corr}}\) is mainly dominated by the uneven distribution of current, which explains the reason that the AZ91 alloy is more sensitive to the TEL thickness in Region I.

Figure 4. (a) nyquist plots and (b) bode phase plots of pure Mg and AZ91 alloy in region I (50 \(\mu\)m) and region II (500 \(\mu\)m).

Figure 5. plots of corrosion current density of pure magnesium and AZ91 alloy as a function of TEL thickness.

In vitro corrosion of pure magnesium and AZ91 alloy
Figure 6a and b shows the three-dimensional (3D) images of the microstructures of pure Mg and the AZ91 alloy, respectively. The microstructure for pure Mg was characterized by α-Mg phase, while for the AZ91 alloy both Mg17Al12 (β) phase and the α-Mg phase. The Mg17Al12 distributed at grain boundaries (GBs) were evidently distinguished by the height difference between the α-Mg matrix and the β phase (Fig. 6b). When exposed to the electrolyte, the Mg17Al12 phase with a relatively high potential thus acted as the cathode with respect to the adjacent α-Mg phase as the anode. It is clear from Fig. 6b that a relatively rougher surface was formed on the AZ91 alloy due to the potential difference between the α-Mg and the β phases after etching. Figure 6c and d discloses the corrosion morphology of the pure Mg sample and the AZ91 alloy after TEL tests (with a thickness of 50 μm), respectively. Corrosion pits were randomly distributed on the surface of the pure Mg samples due to the presence of impurities enriched in GBs, the potential of which is higher than that of the α-Mg phase. Interestingly, it is found that the height between the α-Mg and β phases increased from 3.54 μm to 5.20 μm for the AZ91 alloy before and after the TEL tests due to the dissolution of the α-Mg matrix. The Mg(OH)2 corrosion products of the AZ91 alloy precipitated on the boundary of Mg17Al12 phase due to the dissolution of the α-Mg matrix surrounding the β phases. Furthermore, Fig. 6e and f illustrates the corrosion morphology of the pure Mg sample and the AZ91 alloy after TEL tests with a thickness of 500 μm. The corrosion product was uniformly covered on the pure Mg sample and the number of the corrosion pits was much less in Fig. 6e than that in Fig. 6c, implying for the pure Mg sample uniform corrosion occurred more preferentially under thicker TEL (500 μm) than thinner TEL (50 μm). The stacking of the corrosion product reduces the height between the α-Mg and β phases of AZ91 alloy in Fig. 6f from 5.2 μm at a thickness of 500 μm to 3.04 μm at a thickness of 50 μm. These phenomena describe that the corrosion behavior of both pure Mg and AZ91 alloy tends to be localization under TEL condition especially when the thickness of TEL is lower than a certain value. Namely, localized corrosion is easier to occur under TEL condition than in the bulk solution.

Conclusions

A TEL cell has been built to mimic the in vivo microenvironment and utilized to evaluate the in vitro corrosion rate of biomedical magnesium alloys. It is concluded as following:

1. The TEL can inhibit the in vivo corrosion of pure Mg and the AZ91 alloy. When the thickness of TEL less than 200 μm, the AZ91 alloy is more sensitive than pure Mg to the inhibition of corrosion; however, there is no significant change in corrosion current density for pure Mg.
2. The TEL thickness affects the distribution of current. Under the TEL condition (i.e. 50 μm thick), corrosion resistance for the pure Mg and the AZ91 alloy is predominately controlled by the uneven distribution of current. On the contrary, in the bulk solution (i.e. 500 μm thick), corrosion resistance for the pure Mg and the AZ91 alloy is predominately controlled by the uniform distribution of current.

3. The 3D images demonstrate that localized corrosion is more preferential to occur under the TEL condition than in the bulk solution.

4. The in vitro corrosion performed on the TEL cell may reflect the slight influence of tissue microenvironment. The TEL device might be a universal in vitro method for biodegradable Mg implants. However, it should be noted that the in vitro circumstance was a complex system including inorganic ions, corrosion medium volume, pH, blood flow, proteins and tissues. The TEL was still a micro-scale model, which needs to consider with other factors on comprehensive degradation.

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Conflict of interest statement. The authors declare that they have no competing financial interests.

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