

Sustainable application of magnesium alloy through Electrodeposition of Hydroxyapatite Coating

Motilal Lakavat^{1*} Isaac Ramalla² and G Praveen kumar²

¹Caledonian College of Engineering Muscat, P.O.Box: 2322, CPO Seeb 111, Sultanate of Oman,

²Department of Electrical Engineering, University of Petroleum and Energy Studies, Bidholi Campus, Dehradun,

Abstract: Magnesium (Mg) and its alloy are potential material for biodegradable implants which makes promising due its mechanical properties. However, corrosion of these metals alloys is too quickly results in losing mechanical integrity. The problem had risen in scientific committee to solve/reduce the corrosion rate and enhance the biocompatibility of the Mg-alloy. In order to reduce the corrosion, surface was modified with the hydroxyapatite (HAp) coating. The synthesis of HAp on Mg-alloy substrate was carried out by an electrodeposition method i.e., chemical deposition. The post treatment of deposited HAp on Mg-alloy with alkali solution which results in electrodeposited calcium hydrogenphosphate coatings into the bone-like hydroxyapatite coatings. Cyclic voltammetry and electrochemical impedance spectroscopy was studied to determine the barrier properties of the HAp coatings. Polarization test in a Hanks' solution was investigated to sense the protectiveness of the coatings, which is critical for better in-service mechanical integrity. The existence of the HAp coating and its morphology on Mg-alloy was verified using Fourier transform infrared spectroscopy (FTIR).

Keywords: Mg-alloy, hydroxyapatite coating, mechanical integrity, biocompatibility.

I. Introduction

Magnesium alloys have excellent specific strength and stiffness, exceptional dimensional stability, high damping capacity, and high recycle ability[1–4]. Based on these superior properties and a combinative requirement for reducing environmental burdens by using light weighted structures, the research and development of magnesium alloys for practical industrial application have overwhelmingly increased worldwide during the past decade. Magnesium and its alloys are becoming widely recognized as playing an increasingly important role in automotive, aircraft, and electronic consumer products[5–7].

Magnesium (Mg), by virtue of its desirable mechanical properties, non-toxicity, and degradability, is a potential candidate metallic material for degradable implants. Biodegradable implant materials in the human body can be gradually dissolved, absorbed, consumed or excreted, so there is no need for the secondary surgery to remove implants after the surgery regions have healed[8–10]. Currently, degradable implants are made of polymer materials. Owing to the low mechanical strength, polymeric implants are used only in low load-bearing applications[11,12]. Thus the development of high load-bearing degradable orthopedic implants is an attractive idea for bone fracture healing. According to a recent review by Staiger et. al, the mechanical properties of Mg are close to those of bones, and Mg is non-toxic and excess of Mg in human body can be easily excreted via urine[13].

Magnesium alloys can be divided into cast magnesium alloys and wrought magnesium ones in terms of difference in processing. Main commercial magnesium alloys include the AZ series (Mg-Al-Zn), AM series (Mg-Al-Mn), AE series (Mg-Al-RE), EZ series (Mg-REZn), ZK series (Mg-Zn-Zr), and WE series (Mg-RE-Zr). Statistically, more than 90% of the magnesium alloy structural components are produced by casting process, especially by various die-casting processes[13–15]. In the last decade, the AZ series cast magnesium alloys, especially the AZ91 alloy, have been extensively studied and used for some structural components of automobiles, aircraft, and computers, because of high specific strength and good cast ability [16]. Compared with cast magnesium alloys, wrought magnesium alloys have a more promising perspective of application, which is the reason why research and development of high performance wrought magnesium alloys, novel wrought magnesium processing technology, and producing high quality wrought magnesium alloy products are an important objective. A long time research plan of wrought magnesium alloys was put forward by the international magnesium association (IMA) in 2000[17]. In recent years, the research on wrought magnesium alloys is increasingly attracting the attention of companies and researchers[18–21].

Many methods such as plasma spraying, pulse laser melting, and physical vapor deposition cannot be used to deposit HA coating on Mg-alloy because of its low melting point and poor heat resistance. Electrochemical deposition has unique advantages due to its capability of forming uniform coating and simple setup. In addition, the deposition processing can be conducted at room temperature and the morphology of

coating can be controlled easily by varying the electrochemical potential and other parameters. In this paper we report the preparation of hydroxyapatite (HAp) coatings on magnesium Mg-alloy by electrodeposition and subsequent treatment in alkaline (NaOH) solution and by an alternative immersion method (AIM)[22–25]. The barrier properties of the coatings in the Hanks' solution were investigated using electrochemical impedance spectroscopy (EIS) and voltammetry methods. Morphology of the HAp coatings was studied using Fourier transform infrared spectroscopy (FTIR)[26–28].

II. Experimental

Cylindrical shaped electrodes having surface area 0.235 cm², in a glass tube and working electrode with Mg-alloy composition of 8.6 Al, 0.51 Zn, 0.19 Mn, 0.05 Si, 0.025 Cu, 0.004 Fe and remaining Mg. Electrodes are polished with alumina powder of 0.05 μm, in ethanol in ultrasonic bath, rinsed with distilled water, kept immersed in 1.0 M solution of NaOH at 80 °C for 1 h, and finally rinsed with distilled water. Electrodeposition of HAp was carried out at constant potential values of –2.0 V for different periods of time in a solution (100 cm³) containing 50 cm³ 0.1 mol dm⁻³ Ca(NO₃)₂ and 0.06 mol dm⁻³ NH₄H₂PO₄. The pH of the solution was adjusted with 0.5mol dm⁻³ HNO₃ to 5. The hanks solution was used to test the corrosion resistance of HAp at 37°C. All chemicals were of p.a. purity.

Hanks solution experiment was carried out in standard three-electrode cell capturing 100 cc of solution. The counter electrode was a large area platinum electrode and the reference electrode, to which all potentials are referred, was an Ag/AgCl/3M KCl (209 mV vs. SHE). Potentiometric measurements were performed at a sweep rate of 10 mV s⁻¹. Electrical open circuit potential (EOCP) was measured in the frequency range from 100 kHz to 0.1 Hz with an AC signal ±5 mV. SolartronZView® software was used to fit the Impedance data of suitable electrical equivalent circuit (EEC) model, employing the complex non-linear least squares (CNLS). The HAp deposit was characterized by FTIR. The FTIR spectra were recorded in the 4000–650 cm⁻¹ region with the scan resolution of 4 cm⁻¹ using Horizontal Attenuated Total Reflectance (HATR) method on a Perkin-Elmer spectrum one FTIR spectrometer.

III. Results and Discussion

3.1 Electrochemical deposition of hydroxyapatite

Electrodeposition of hydroxyapatite film was performed using Modification of the electrodes at room temperature in a solution (100 cm³) containing 50 mL 0.1 mol dm⁻³ Ca(NO₃)₂ and 50 ml 0.06 mol dm⁻³ NH₄H₂PO₄ which is having a pH 5 at constant potential value of -2.0 V. The electrodeposition time was chosen between 30 to 180 min. Reactions on the surface of Mg-alloy during electrodeposition are known as Step I: reduction of the hydrogen from H₂PO₄⁻ and HPO₄²⁻ and Step II: Ca²⁺ ion reacts with HPO₄²⁻ and PO₄³⁻ and form CaHPO₄ × 2H₂O and Ca₃(PO₄)₂. Brushite and β - TCP are the precursors to the more stable phase hydroxyapatite, Ca₁₀(PO₄)₆(OH)₂, HAp. Alkaline treatment (acid-base reaction) converts brushite and β-TCP film formed on the Mg-alloy surface by electrodeposition into hydroxyapatite, HAp. These results clearly show that electrodeposited coating prior to alkaline treatment is mainly consisted of calcium phosphate dihydrate and β-TCP. Alkaline treatment results in the HAp coating formation. Prior investigations have also utilized the increased solubility of brushite on metallic implants as a means of increasing the amounts of calcium and phosphate ions available in the immediate vicinity of the implant to promote increased osseointegration.

3.2 Barrier properties of the coating

Nyquist plot for various frequency dispersions (a) for an ideally capacitive behavior of plain electrodes; (b) for the in-a-pore dispersion simulated by de-Levie's standard equation (c) for the in-a-pore dispersion and by pore size distribution simulated by TLM-PSD. The impedance of most electrodes deviates from the purely capacitive behavior which is represented as a vertical line in the Nyquist plot, curve (a) in Figure 1

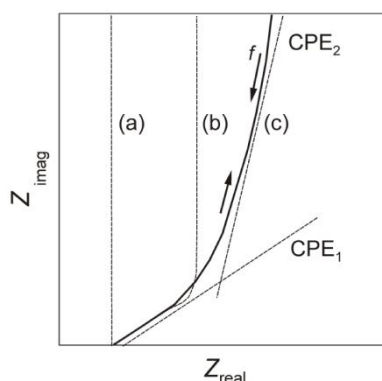


Figure 1 Nyquist plot of various frequencies to compare the ideal behavior

3.3 Electrochemical impedance spectroscopy

From the figure 1, it is describe that CPE_1 at high frequencies and with an ideal capacitance at low frequencies while curve (c) can be approximated with the CPE_1 at high frequencies and roughly with the CPE_2 at low frequencies. Frequency dispersion leads to an inclined line whose phase angle is equal to $(\pi/2)n$. The empirical distributed element was used to describe the non-ideality due to frequency dispersions by the constant phase element (CPE). Its impedance is equal to $Z_{CPE} = [Q-(j\omega)^n]^{-1}$, where Q is the frequency independent parameter, ω is the angular frequency, and n is the CPE power. When $n = 1$, Q represents the pure capacitance, while for $n \neq 1$ the system shows behavior that has been attributed to the surface heterogeneity, or to the continuously distributed time constants for charge transfer reactions.

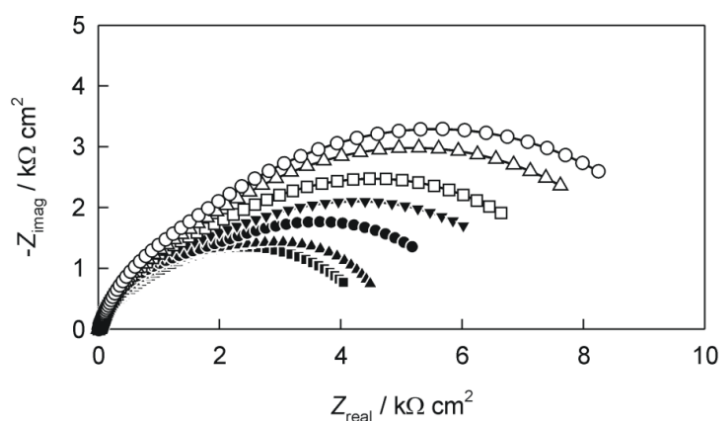


Figure 2. Nyquist and Bode plots for the impedance data for uncoated Mg-alloy

Nyquist and Bode plots for the impedance data recorded with uncoated Mg-alloy (with a native surface layer) and HAp coated Mg-alloy in the Hanks' solution (37°C) at E_{OCP} . Electrodeposition was performed at -2.0 V for denoted time periods. Corrosion resistance of Mg-alloy unmodified and modified with a hydroxyapatite (HAp) film in the Hanks' solution was investigated by EIS. Figure 2 shows the Nyquist and Bode plots of impedance spectra of unmodified and with HAp film modified Mg-alloy recorded in Hanks' solution at open circuit potential after 30 min of stabilization. The Nyquist plot for unmodified Mg-alloy show flattened capacitive semicircle with the center below the real axis.

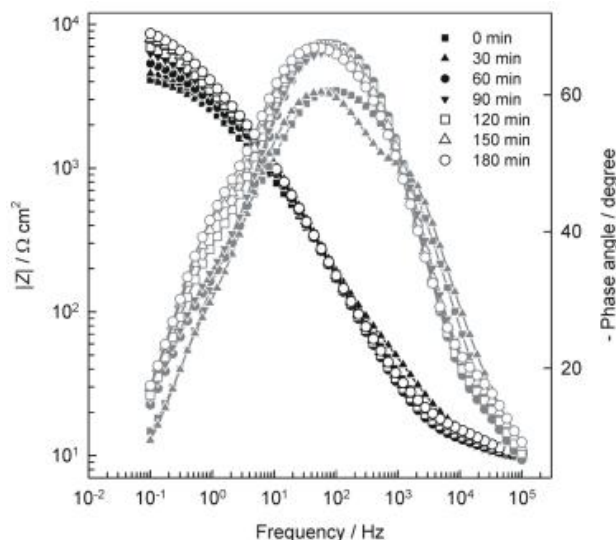


Figure 3. Nyquist and Bode plots for the impedance data of Mg-alloy

The overall impedance of modified Mg-alloy was plotted with Nyquist for coated with hydroxyapatite was significant compared to unmodified Mg-alloy. Electrodeposition time of modified Mg-alloy was significant as shown using capacitive semicircle increase. The presence of hydroxyapatite was observed in low frequency region of bode plot. The value of $\log |Z|$ is significant at which dominates the polarization resistance. EIS data recorded for the unmodified Mg-alloy in Hanks' solution was modelled by the electrical equivalent circuit (EEC) with two time constants.

In the EEC, constant phase element CPE_1 attributed to the double layer capacitance, and R_1 to the charge transfer resistance and CPE_2 and R_2 attributed to the resistance of the film and capacitance of the ions travel through the film. In the case of electrodes modified with hydroxyapatite at low frequency region of the impedance spectra an additional time constant appears. It is attributed to the diffusion processes and described by the Warburg diffusion element W_s . The total impedance, Z of the investigated electrochemical system is the sum of the ohmic resistance and the impedance of the electrochemical interface ($Z_{el} + Z_{HF} + Z_{LF}$) and is described by the transfer function:

$$Z(j\omega) = R_{el} + \left\{ Q_1(j\omega)^{n_1} + \left\{ R_1 + \left[Q_2(j\omega)^{n_2} + (R_2 + Z_W)^{-1} \right]^{-1} \right\}^{-1} \right\}^{-1}$$

The sum of R_1 and R_2 represent the polarization, resistance and a diffusional resistance (R_1-R_2). Electrodeposition of hydroxyapatite affects the corrosion behavior which was observed during the experiment, i.e., polarization resistance of the Mg-alloy in Hanks' solution. The value of polarization resistance increases with an increase of the electrodeposition time. Electrochemical double layer capacitance decreases with increases of the electrodeposition time.

Therefore, the EEC includes also the so-called "O" circuit element ("porous bounded Warburg"). The term "O" represents a dimensional diffusion through a layer of finite thickness or the (penetrability) penetration depth of the AC signal across pores of finite length. The impedance parameters of uncoated Mg-alloy and Mg-alloy samples coated with HAp coatings, obtained in Hanks' solution (37°C) at EOCp. Electrodeposition was performed at -2.0 V for denoted time periods.

3.4 Polarization measurements

Polarization curves measurements were recorded with uncoated and HAp coated alloy specimens, using a scan rate of 10 mV s^{-1} , are shown in Figure 3. It is observed that polarization curve for all specimens exhibits a linear Tafel region with slope values ranging from -235 mV dec^{-1} to -140 mV dec^{-1} at cathodic branch of cell. This is similar observation as in case of hydrogen evolution reaction, which occurs at electrodes covered with a surface film. The values obtained agree well with those reported for Mg-alloys that were unmodified and modified with SAMs of fatty acids or phosphonate SAMs, and immersed in near neutral aqueous solutions containing chloride ions. The hydrogen dissolution at anodic region i.e., simultaneous combination of both anodic dissolution and localized corrosion has been reported for reason of complicated anodic polarization behavior of unprotected Mg-alloy. Figure 4, shows at different times the deposition of HAp at cathodic and anodic electrodes.

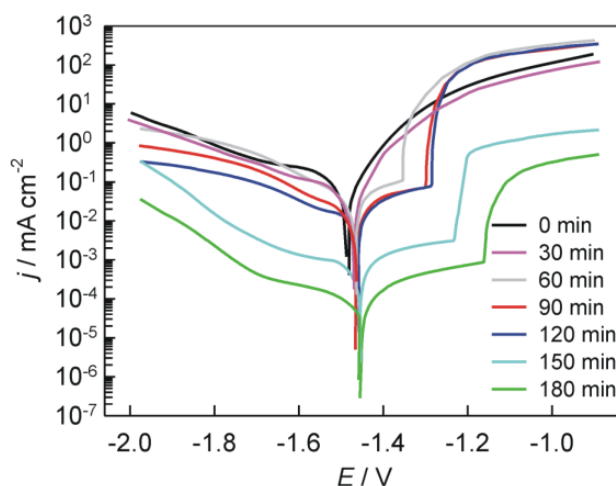


Figure 4 Tafel plots recorded with uncoated Mg-alloy

Tafel plots recorded with uncoated Mg-alloy (with a native surface layer) and HAp coated Mg-alloy in the Hanks' solution at 37°C; $v = 10 \text{ mV s}^{-1}$. It can be observed from the Figure 4, that Electrodeposition was performed at -2.0 V for denoted time periods. The rate determining for corrosion process of all specimens are indicated around E_{corr} by the cathodic and anodic portions of the polarization curves. Extrapolation of the linear part of the cathodic currents to the corrosion potential was used for a rough estimation of the corrosion current densities (j_{corr}). The numerical values of the corrosion kinetic parameters b_c , b_a , j_{corr} and E_{corr} , obtained by fitting experimental data, are shown in figure 4. The numerical values of the corrosion current densities determined for untreated alloy specimens is $41.2 \mu\text{A cm}^{-2}$ and for coating HAp films formed on Mg-alloy for 180 min at -2 V is $0.1 \mu\text{A cm}^{-2}$.

3.5 Characterization of HAp coating

The Mg-alloy coated with electrodeposited HAp coating was characterized by FTIR spectra. Electrodeposition was performed at -2.0 V for 3 h. Figure 5 shows a FTIR spectrum of Mg-alloy substrate coated by electrodeposited HAp film that was subsequently alkali treated. A strong ν_3 PO_4^{3-} vibration band, associated with the internal modes of PO_4^{3-} (P-O asymmetric stretching vibrations), are observed inside the frequency range from ca. 1000 cm^{-1} to 1100 cm^{-1} at 1009 cm^{-1} and 1088 cm^{-1} . The ν_3 asymmetric P-O stretching mode is triply degenerate in the free phosphate ion, but this mode is resolved into at least two distinct peaks in crystalline hydroxyapatite.

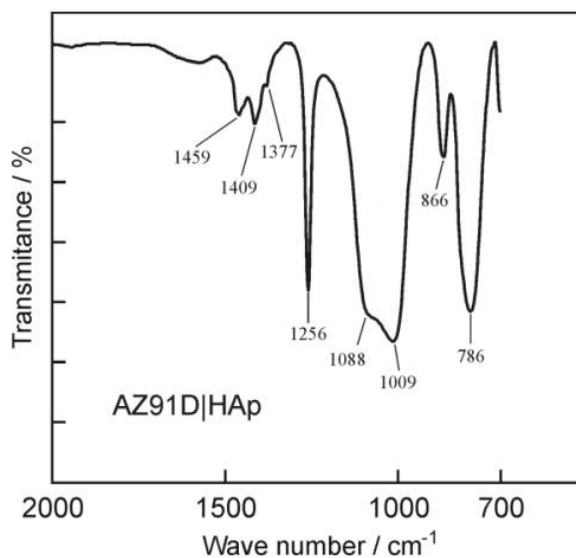


Figure 5. FTIR spectra of Mg-alloy coated with electrodeposited HAp

Hydroxyapatite was observed from two ν_3 bands become increasingly resolved the crystallinity from the figure 5. In our infrared spectrum, the two ν_3 bands are fine-resolved, which indicates that the coating is crystalline. The peaks at 1459 , 1409 and 1337 cm^{-1} are due to the presence of carbonate ions in the hydroxyapatite structure. It is well known that carbonate, generated from reaction of atmospheric CO_2 with aqueous solutions, readily substitutes for phosphate ions (B-type incorporation; i.e., CO_3^{2-} for PO_4^{3-} substitutions) in the crystal structure of hydroxyapatite. The introduction of these different ions was beneficial to the biocompatibility because of natural bone tissue also contained Mg^{2+} , Na^+ , HPO_4^{2-} and CO_3^{2-} which gives an additional advantage because it leads to an enhanced osseointegration.

3.6 Chemical deposition of hydroxyapatite

When it comes to light metals, such as Al and Mg that have surfaces with mesoporous morphology, the alternative immersion method (AIM) can be used to efficiently deposit synthetic HAp. AIM method is a strikingly simple method to apply HAp coatings to a surface schematically outlined in Figure 5. In the present approach, the specimens were exposed to 20 cycles of alternating immersion in saturated $\text{Ca}(\text{OH})_2$ and $0.02\text{ M NH}_4\text{H}_2\text{PO}_4$. The holding time in each solution was 1 min at room temperature. Between each soaking step and after each cycle, the samples were rinsed in ultrapure water for 1 min. Finally, the samples were dried by nitrogen gas. After coating with HAp by AIM barrier properties of coating on the Mg-alloy surface were investigated by EIS. As the shape of spectra is identical to those of fitted with the same EEC.

IV. Conclusion

Electrodeposition method was used to modify/increase the bioactivity and biocompatibility the Mg-alloy surface with hydroxyapatite film. Brushite film and film of a β -TCP was deposited on Mg-alloy surface and converted into hydroxyapatite (HAp) via an acid-base reaction. Corrosion was tested in Hanks' solution and found that Mg-alloy modified with hydroxyapatite has a higher corrosion resistance in comparison to the unmodified Mg-alloy. The protecting efficiency of HAp increases with the increasing deposition time to obtain a thicker and a more compact coating. By providing higher polarization resistance values in comparison to the surface film by immersion method, electrodeposition method was superior. FTIR was implemented to identify the chemical groups of HAp coating on Mg-alloy confirmed the presence of the thick uniform layer of hydroxyapatite. Hydroxyapatite-coated Mg-alloy alloy has good potential for use in medicine, because the HAp films has a good bioresorption.

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