In-vitro Biodegradation of Micro-Arc Oxidation on AZ31 Hybridized with Dopamine Compared to Phytic Acid

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Abstract
Hybrid coatings composed of the inorganic film; Micro-Arc Oxidized (MAO) and organic film; Dopamine (DOP) or Phytic Acid (PA) was successfully deposited on AZ31. The DOP@MAO@AZ31 film reduced the corrosion current 3-times the MAO film while, the PA@MAO@AZ31 presented a slight improvement. Here, the film morphology is the master key for the corrosion resistance enhancement in Simulated Body Fluid (SBF). The Dopamine film supported the MAO inner layer and blocked the porous structure. The Phytic acid transformed the porous MAO layer into a cracked structure to facilitate corrosive ions attack. All experimental outcomes were verified by Scanning Electron Microscope (SEM), Fourier Transform Infrared Spectroscopy (FTIR) and X-ray Diffraction (XRD). The in-vitro corrosion performance was estimated utilizing electrochemical measurements.

Keywords: Micro-arc oxidation; Corrosion resistance; Hybrid coating; Mg

Graphical abstract

(Figure 1)

Introduction
More and more attention has been paid to magnesium alloys as anew generation of medical implant materials [1]. Mg alloys present physical and mechanical properties compatible with those of human bone however, the major obstacle for their use is the undesirable high corrosion rate [1]. This problem encourages the search for a protective coating to expand the use of Mg alloys. Currently, surface modification such as plasma electrolytic process (PEO) [2]
or familiar as Micro-Arc Oxidation (MAO) is an effective method for improving their corrosion resistance [3-5].

Micro arc oxidation is sparked-anodization method where a high-voltage is used for the surface treatment of magnesium and its alloys. It consists of three layers; outer porous layer, the middle-fairly compact layer and the inner-compact layer sticking to the surfaces [6,7]. Thus, MAO process usually deposits a micro-nano scale porous bio ceramic layer on substrates [8] that allows a corrosive medium to move quickly into the substrate. Coupled with another technique [9] is a suitable solution for more protection such as dipping coating. Taking account of the fact that immersion the MAO films in the appropriate additive as post-treatment is valuable in inhibiting the corrosive ions.

Dopamine (DA) is a common adhesive material that serves as an antibacterial film for biomedical applications [8]. It is 3, 4-dihydroxyphenethylamine which is a natural hormone with a fundamental role in the brain and body. It is made of the catecholamine and phenethyl amine families consequently consists of a catechol structure (a benzene ring with two hydroxyl side groups) with one amine group attached via an ethyl chain. This structure qualify it to chelate with surface as a protective coating with bioactivity characters [10,11].

Phytic Acid (PA) is a non-toxic organic compound with 12 hydroxyl groups to provide a powerful capability of chelating with Mg substrate. When PA hybridized with Silane compound on MAO film, an excellent barrier effect was obtained [12]. The purpose of the present work is to compare the hybrid coatings of DOP@MAO@AZ31 and PA@MAO@AZ31 to the single MAO@AZ31coating under the same conditions of electrolytes.

**Experimental Procedure**

**MAO coating preparation**

AZ31 magnesium was provided from Xinxiang Jiuli magnesium Co., Ltd. (Xinxiang, China) with nominal compositions wt.%: 2.5-3.5 Al, 0.6-1.3 Zn, 0.1 Si, 0.04 Ca, 0.2-0.8 Mn, 0.05 Cu and balanced Mg. The samples were cut with a thickness of 1.5mm and a diameter of 30mm then, ground with silicon carbide sandpaper up to 2000 grit. Those samples were ultrasonically cleaned 5-10 s in alcohol and distilled water to let dry in air. Then, immersed in Na$_2$SiO$_3$, 9H$_2$O 0.04M and Na$_3$PO$_4$, 12H$_2$O 0.04M for 3 minutes with 200V using 1kW AC power supply with a duty cycle of 10%.to get MAO@AZ31. Those samples were dried and immersed in Phytic acid (CH$_3$CH$_2$COOH) 0.25M 4h, 60°C to yield PA@MAO@AZ31 or in dopamine solution (C$_8$H$_{11}$NO$_2$) 0.25M 12h, pH 9 to get DOP@MAO@AZ31. Finally, those specimens were washed with distilled water three times and dried in the air prior to characterization.

**Coating characterization**

The optical microscope and scanning electron microscopy (SEM, JSM-7401F JEOL, Japan) were utilized to observe the surface morphologies of the prepared samples. The electron beam of SEM is 0.8nm at 15kV under a pressure of 4.45-10.4Pa. A very thin layer of gold must be sputtered to the samples before the SEM observations. A goniometer (DSA100, Kru€uss, Hamburg, Germany) at 25°C and 60% relative humidity was utilized to record the water contact angle for the coated samples in static drop mode. For each sample, at least five measurements on random surface sites were tested.

Fourier transform infrared spectroscopy (FTIR, Nicolet 5700, Thermo Electron Corporation, MA, USA) was utilized to characterize the functional-groups within the coatings (organic film). The scanning rates from 500 to 4000cm$^{-1}$, then the data are converted into absorbance spectra. For further more exposure (inorganic film), the phases structures of the samples were characterized by X-ray diffraction (XRD, Philips X'Pert) using a CuK radiation with a glancing angle of 2°. The XRD data were obtained over 20 range of 20-40° at a step size of 0.25°.

**Electrochemical corrosion behavior**

Potentiostat (IM6, Zahner, Germany) was used to test potentiodynamic polarization (PDP) and Electrochemical Impedance Spectroscopy (EIS). The corrosive electrolyte was the fresh-prepared Simulated Body Fluid (SBF) with similar ionic composition to that of human blood plasma. It contains 0.8g/l NaCl, 0.4g/l KCl, 0.14g/l CaCl$_2$, 0.35g/l NaHCO$_3$, 1.0g/l C$_4$H$_8$O$_3$.glucose, 0.2g/l MgSO$_4$, 7H$_2$O. 0.1g/l KH$_2$PO$_4$, H$_2$O, 0.06g/l Na$_2$HPo$_4$, 7H$_2$O, pH 7.4, at a temperature of 37°C. A three-electrode cell consists of platinum foil (1 cm$^2$) as a counter electrode, saturated calomel electrode (SCE, Lei CI 232, Shang Hai, China) as reference and one the samples of an area of 0.79 cm$^2$ were the working electrodes.

The PDP curves were obtained by scanning the electrode potential from -2.0 to -1.0 V $\text{SCE}$ at a scanning rate of 1mV s$^{-1}$. The polarization plots were analyzed by linear extrapolation to detect cathodic Tafel slope and specify the values of free corrosion potentials $E_{corr}$, corrosion current density $i_{corr}$. The taken region for data calculation was about 50mV more negative than the free corrosion potential. The EIS spectrum was obtained at the fixed DC potential when the stabilization stopped. The excitation signal of a sinusoidal alternating voltage 10mV was superimposed on the DC potential, which was scanned from 200kHz to 0.01Hz. The acquired EIS data were recorded using the Zsim Demo software.

**Results and Discussion**

**Coating morphology and wettability**

The optical microscope, scanning electron microscope and water contact angle are illustrated in Figure 2. MAO@AZ31 film presented porous structure that transformed into cracked morphology at PA@MAO@AZ31 films. The disappearance of the porous structure could be attributed to the ability of Phytic acid to attack the surface [13]. On the contrary, the DOP@MAO@AZ31 film revealed agglomerated particles of dopamine that blocked the porous MAO structure. Dopamine can effuse through the discharge channels and quenched to yield a compact layer [8], consequently a thick film of 19μm is formed. The water contact angle (WCA) of both MAO@AZ31 and PA@MAO@AZ31 films presented hydrophilic behavior that could be related to their morphology. On the other hand, DOP@MAO@AZ31 coating was hydrophobic with water contact angle 112°C.
Figure 2: Optical microscopy, scanning electron microscope and water contact angle of MAO@AZ31, PA@MAO@AZ31 and DOP@MAO@AZ31 composite coatings.

XRD & FTIR

The composition of the surface coating was analyzed using the XRD patterns in Figure 3a. The XRD pattern detected the phases of the micro-arc oxidized film MAO@AZ31, PA@MAO@AZ31 and DOP@MAO@AZ31. The main phase that can be seen is Mg$_2$SiO$_4$ mixed with NaMgPO$_4$. The weakest intensity of Mg$_2$SiO$_4$ peak appeared at DOP@MAO@AZ31 as due to the presence of the second layer covers well MAO film. FTIR spectra Figure 3b illustrated the peaks that rely on organic function groups. The PA@MAO@AZ31 film revealed OH stretched and bonded groups at 3600 cm$^{-1}$ and 2500 cm$^{-1}$, respectively [12] as an indication of the organic Phytic acid. The DOP@MAO@AZ31 film presented the amine peak at 2150 cm$^{-1}$ [8] to prove the organic dopamine deposition. Both of those analyses confirmed the successful hybridization between the inorganic silicate (Mg$_2$SiO$_4$) with the organic film such as dopamine or Phytic.

Figure 3: XRD and FTIR spectra of MAO@AZ31, PA@MAO@AZ31 and DOP@MAO@AZ31 composite coatings.

Corrosion behavior

The electrochemical potentiodynamic curves, Nyquist plots and optical microscope for the corroded film are illustrated in Figure 4. The Phytic Acid (PA) presented a slight shift in the potential accompanied by passive current (Figure 4a). This result is an indication that cracked structure (PA@MAO@AZ31) yielded no clear difference than the porous structure (MAO@AZ31). On
the other side, the agglomerated dopamine confirmed the barrier property due to more noble corrosion potential and less value of passive current (Table 1). The corrosion current of DOP@MAO@AZ31 was reduced 3-times the MAO single-film. The Nyquist plots supported the PDP curves because the obvious jump in the resistance belongs to the DOP@MAO@AZ31 film.

Figure 4: Potentiodynamic, Nyquist curves and OM for the corroded films of MAO@AZ31, PA@MAO@AZ31 and DOP@MAO@AZ31 in SBF.

Table 1: Electrochemical parameters for MAO@AZ31, PA@MAO@AZ31 and DOP@MAO@AZ31 compared to Bare AZ31.

| Samples         | BareAZ31 | MAO@AZ31 | PA@MAO@AZ31 | DOP@MAO@AZ31 |
|-----------------|----------|----------|-------------|--------------|
| Ecorr (VSCE)    | -1.05    | -1.3     | -1.22       | -1.13        |
| I corr (µA/cm²) | 250      | 1.1      | 0.9         | 0.4          |
| CR (mpy)        | 224      | 2.9      | 2.3         | 0.35         |

The assumed mechanism in case of dopamine which has a higher tendency to agglomerate [14], yielded a greater thickness. This polydopamine (PDOP) second layer can cover well the first layer of silicate, lowers the intensity of Mg₂SiO₄ (XRD). Hence, this double-layer can successfully isolate AZ31 from the corrosive ions, reduce the corrosion rate.

In case, Phytic acid dissolves the first silicate layer according to the following equation [15],

$$\text{Mg}_2\text{SiO}_4 + [\text{H}_4\text{PA}]^+ \rightarrow \text{SiO}_2 + \text{H}_2\text{O} + [\text{Mg}_2]\text{[PA]}^{6-}$$

This acid caused a cracked film (SEM) upon AZ31, facilitated the attack of corrosive ions. Therefore that cracked structure (PA@MAO@AZ31) presented non-remarked enhancement that the porous structure (MAO@AZ31).

Conclusion

A combination of organic film overlays MAO cant for sure provides surface protection. If the organic film such as phytic acid attacked the MAO film, a slight enhancement is obtained. The porous MAO structure when transformed into a cracked one, facilitated the corrosive ions penetration. If the organic film such as dopamine supported the MAO film to effectively isolate the AZ31, a significant improvement is obtained. The dopamine reduced the corrosion current 3 times the MAO@AZ31 due to dopamine agglomeration that blocked the porous structure.

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