Corrosion Behavior and Electrochemical Properties of As-cast Mg-2Zn-0.5Ca-Y Series Magnesium Alloys in Hank's Solution and NaCl Solution

Ren Xiaodong, Li Xuesong, Yang Yue, Yang You, Wu Hua

Key Laboratory of Advanced Structural Materials, Ministry of Education, Changchun University of Technology, Changchun 130012, China

Abstract: Microstructure and bio-corrosion behavior of as-cast Mg-2Zn-0.5Ca-Y series alloys were investigated for biomedical application by optical microscopy, scanning electronic microscopy (SEM), immersion tests and electrochemical measurements in Hank's solution and NaCl solution (3.5wt%). The morphologies of the Mg-2Zn-0.5Ca-Y series magnesium alloys indicate that the corrosion is not uniform and a lot of cracks are formed on the surface of the corrosion layer after immersion tests. The corrosion resistance of as-cast Mg-2Zn-0.5Ca-Y series alloys increases owing to the addition of the rare earth element Y and that of Mg-2Zn-0.5Ca-1.0Y alloy is the best. All of the potentiodynamic polarization curves obtained in the tests in Hank's solution and NaCl solution show current plateaus which are due to the presence of a protective corrosion product film. The electrochemical impedance curves of the alloys in the test solutions have only one capacitive loop. The appearance of a capacitive loop at high frequency is caused by the stable double-layer capacitance. The capacitive loop at low frequency is attributed to diffusion control at film-free areas.

Key words: Mg-2Zn-0.5Ca-Y series alloy; corrosion behavior; potentiodynamic polarization; electrochemical impedance

Magnesium alloys show potential application as biodegradable materials due to their outstanding biological performance. Recently, it has even been reported that the presence of magnesium in the bone system is beneficial to bone strength and growth^[1]. Unfortunately, rapid corrosion is an intrinsic response of magnesium alloys to chloride containing solutions^[2], including human body fluid or blood plasma. In general, it is accepted that Mg is easily oxidized to form a thick hydroxide film when it is in contact with humid air or water^[3,4]. In solutions containing Cl⁻, the corrosion resistance of magnesium alloy is much worse. Enhancement of the corrosion resistance of magnesium alloys is one of important issues if their application is to be increased.

The corrosion behavior of magnesium alloys with rare earth (RE) element additions has been investigated during past decades^[5,6]. It is well known that the addition of rare earth elements is an effective way to enhance the corrosion

resistance of magnesium alloys. The improvement has been attributed primarily to the formation of metastable RE-containing phases along the grain boundaries. Another important effect of RE on the corrosion resistance of these alloys is the so-called "scavenger effect", i.e. some impurity elements in the alloys, such as Fe, will deteriorate their corrosion resistance significantly. RE additions are reported to cancel the influence of such impurities by the formation of intermetallic compounds^[7].

Yttrium (Y) is a particularly interesting alloying element for magnesium alloys because Y has a standard electrochemical potential (−2.372 V) equal to that of magnesium (−2.372 V)^[8]. More recently, researchers focused on the effects of Y on magnesium alloys. The corrosion behavior and electrochemical properties of Mg-2Zn-0.5Ca-Y series alloy have been evaluated in Hank's solution and NaCl solution.

In the present study, a new Mg-2Zn-0.5Ca-Y series alloy,

Received date: December 9, 2015

Foundation item: National Natural Science Foundation of China (51171030); Science and Technology Department of Jilin Province Key Project (20120345)

Corresponding author: Li Xuesong, Ph. D., Professor, Key Laboratory of Advanced Structural Materials, Ministry of Education, Changchun University of Technology, Changchun 130012, P. R. China, Tel: 0086-431-85716260, E-mail: lixuesong@mail.ccut.edu.cn

Copyright © 2017, Northwest Institute for Nonferrous Metal Research. Published by Elsevier BV. All rights reserved.

used as a high performance structural material as well as a biomaterial containing different Y element concentrations, was prepared by casting and was tested to determine the amount of elemental Y that could provide higher corrosion resistance for Mg-2Zn-0.5Ca-Y series alloys.

1 Experiment

The chemical composition of as-cast Mg-2Zn-0.5Ca-Y series alloy used in this work is listed in Table 1. The alloys were prepared by conventional casting methods under the protection of molten salt. The melts were then poured into a steel die with 90 mm×45 mm×12 mm in size at 800 °C. The specimens for the tests were machined to form a cast sheet.

Rectangular samples for the immersion test with a dimension of 10 mm×10 mm×2 mm were cut from the cast alloy directly. Before the immersion tests, the surface of the specimens was polished with 2000 grit emery paper and finely polished with 0.5 μm diamond powder, cleaned with alcohol solution, and then dried in a warm air blast. Then the sample was immersed in 150 mL Hank's solution under ambient conditions at room temperature (the ratio of the volume of the solution to the surface area is 150 mL:2 cm²)^[9] for up to 120 and 168 h. The chemical composition of Hank's solution is listed in Table 2.

The corrosion rate was cross-checked by measuring the mass of the specimens before and after the immersion tests. The latter was done after cleaning and removal all of corrosion products in 15%CrO₃+1%AgNO₃+400 mL H₂O solution.

The corrosion morphology observation of the alloy before and after immersion was performed using optical microscopy (OM) and scanning electron microscopy (SEM). The phase composition of the alloys was studied using X-ray diffraction (XRD).

A three-electrode test cell was used for electrochemical measurements. The electrochemical impedance spectroscopy (EIS) studies were performed at open circuit potential with amplitude of 10 mV over the frequency range of 10⁵~0.01 Hz on specimens. The specimens were exposed to the corrosive electrolytes of NaCl solution (3.5wt%) and Hank's solution to

investigate the corrosion mechanism. All the tests were carried out at room temperature.

The electrochemical tests used a 'Bio-logic' VSP potentiostat/frequency response analysis system to evaluate the corrosion behavior of the specimens at room temperature. A saturated Ag/AgCl (saturated with KCl) was used as the reference electrode. A platinum mesh and the investigated specimen were used as the counter electrode and the working electrode, respectively. The electrochemical tests were conducted in solutions at a scan rate of 5 mV/s. The polarization curves were used to estimate corrosion and passivation potentials (E_{corr} , E_{pp}), and corrosion current density (I_{corr}) at the corrosion potential (E_{corr}) by the Tafel extrapolation of the cathodic branch. Five tests were performed on each sample to obtain the average response and the standard deviation.

2 Results and Discussion

2.1 Morphology and corrosion products

Fig.1 shows the microstructures of the as-cast Mg-2Zn-0.5Ca-Y series alloys before immersion testing. It can be seen that the structure of alloy I is composed of multiphase structure and has no continuous grain. The structure of the others are composed of many fine equiaxed crystal grains that have been elongated and the second phase precipitates appear at the grain boundaries. After immersion in Hank's solution for 120 h and 168 h, the specimen surface was covered with a film of corrosion products. The samples were cleaned in a solution comprising 15% CrO₃, 1% AgNO₃, and 400 mL H₂O for 10 min and then ultrasonically cleaned in alcohol for 7 min. The samples then were dried in a warm air blast. The morphologies of the cleaned Mg-2Zn-0.5Ca-Y series alloys are shown in Fig.2. The surface of Mg-2Zn-0.5Ca alloy is covered by a corrosion layer with many cracks, and some white particles could be found on the corroded surface. Few cracks and white particles could be seen on the surface of alloy II, III, IV, and V, which indicates that the corrosion layers of these alloys are denser than that of alloy I. However, there are many precipitates could be found on the surface of alloy IV and V. Representative XRD patterns from the alloy samples before immersion in the test solutions are shown in Fig.3. Samples are mainly consisted of Mg, Mg₃YZn₆, Mg₂Zn, and MgY for alloy IV and V.

2.2 Immersion tests

During the immersion tests, hydrogen was generated continuously on the surface of the samples and the surface gradually became black corrosion product. After the samples were cleaned, their surface exhibited signs of corrosion attack. It is evident that the corroded areas are uneven and pitting has occurred on the surface. There is evidence of crack formation and some of the surface layer has become detached. The uneven surface originates from the attack by the chloride ions and the dissolution of the Mg. Fig.4a and 4b indicate that the oxygen contents of alloy I and alloy II are significantly higher

Table 1 Chemical composition of as-cast Mg-2Zn-0.5Ca-Y series alloy (wt%)

Alloy No.	Zn	Ca	Mg	Y
I	2.0	0.5	Bal.	0
II	2.0	0.5	Bal.	0.5
III	2.0	0.5	Bal.	1.0
IV	2.0	0.5	Bal.	1.5
V	2.0	0.5	Bal.	2.0

Table 2 Chemical composition of Hank's solution (g/L)

NaCl	CaCl ₂	Na ₂ HPO ₄ ·12H ₂ O	KH ₂ PO ₄	MgSO ₄	H ₂ O	C ₆ H ₁₂ O ₆
8.00	0.14	0.06	0.06	0.20		1.00

Fig.1 Microstructures of the as-cast Mg-2Zn-0.5Ca-Y series alloys before the immersion tests: (a1, a2) alloy I, (b1, b2) alloy II, (c1, c2) alloy III, (d1, d2) alloy IV, and (e1, e2) alloy V

Fig.2 Morphologies of the Mg-2Zn-0.5Ca-Y series alloy that had been cleaned after immersion in Hank's solution: (a) alloy I, (b) alloy II, (c) alloy III, (d) alloy IV, and (e) alloy V

at corrosion region. Combined with Fig.2, the corrosion could be determined to be localized corrosion. Compared with alloy I and alloy II, there are less oxygen contents for alloy III and alloy IV, and porous honeycomb structures form on the corrosion surface. The corrosion surface is more uniform and the corrosion is general corrosion as shown in Fig.2 and Fig.4c, 4d. From Fig.2 and 3, we know that the addition of Y in the alloy leads to the formation of the second phase MgY,

which can form corrosion micro-batteries and decreases the corrosion resistance of alloys as it destroys the protective film. Based on above results, it could be concluded that the proper addition of Y can enhance corrosion resistance.

The corrosion rate of Mg-2Zn-0.5Ca-Y series alloy in Hank's solution for different exposure time is shown in Fig.5. The trends of the corrosion rate were calculated by measuring the mass of the specimens before and after the immersion

Fig.6 Potentiodynamic polarization curves (or 'Tafel' curves) for the Mg-2Zn-0.5Ca-Y series alloys after immersion in 3.5wt% NaCl solution (a) and in Hank's solution (b) at room temperature

Fig.7 Nyquist plots for Mg-2Zn-0.5Ca-Y series alloys after immersion in 3.5wt% NaCl solution (a) and in Hank's solutions (b) for differing periods up to 12 h

Hank's solution for 12 h, respectively. In all cases, single capacitive loops are observed, and the sizes increase with the increasing of Y content. The change in the impedance response may reflect thickening of the passive film on the surface of the electrodes^[11]. However, independent of the increase in the content of element Y, the size of the 'semicircles' is greater for Alloy III compared with that of Alloys I, II, IV, or V, indicating an improvement in the corrosion resistance of rare earth containing electrodes.

It is observed that the corrosion mechanism in Hank's solution is similar to that in NaCl solution. The radius of the Nyquist plot is largest when the content of the element Y is 1.0%, and that of alloys IV and V decrease due to the formation of the second phase MgY which destroy the protective film. The radii of all the remained curves increase with the increasing of Y content compared with alloy I, which indicate that element Y improves the corrosion resistance of Mg-2Zn-0.5Ca-Y series alloys. Thus, the addition of Y to the Mg-2Zn-0.5Ca-Y series alloys results in the observed grain refinement.

3 Discussion

Previous report^[11] indicated that enrichment of rare earth played a role in hindering Mg grain growth and induced grain refinement and the fewer Y can promote the grain size refinement and form fewer corrosion micro-batteries^[12,13], which is consistent with our results as shown in Fig.1.

From immersion test results, it is evident that the corrosion rate increases with the increasing of immersion time (Fig.5). The surface of Mg-2Zn-0.5Ca alloy is covered by a corrosion layer with many cracks and white particles. Fewer cracks and white particles could be observed on the surface of alloy II, III, IV, and V. It means that the corrosion layer of these alloys are denser than that of alloy I (Fig.2). Compared with the others, the surface of alloy I has more corrosion points and pits. After addition of Y, the corrosion points and pits decrease. Therefore rare-earth Y can improve the corrosion resistance of Mg-2Zn-0.5Ca-Y series alloys. However, with an increase in Y content to 1.5% and 2.0% the corrosion rates become much higher. The addition of more Y in the alloy leads to the formation of a second phase MgY. The second phase MgY can form many corrosion micro-batteries, which accelerate the corrosion rate of the alloy, so that many precipitates on the surface of alloy IV and V could be found as shown in Fig.2. From above discussion, it could be concluded that when the Y content is smaller than what can induce the formation of a second phase MgY, the addition of Y could improve corrosion resistance. In the present work, the corrosion rate is lowest when the amount of Y addition is 1.0% and the corrosion resistance of Mg-2Zn-0.5Ca-1.0Y alloy is the best.

4 Conclusions

1) The best corrosion resistance of Mg-2Zn-0.5Ca-Y series

alloys is achieved when the content of Y is 1.0%.

2) When the content of Y is smaller than 1.0%, the corrosion resistance of as-cast Mg-2Zn-0.5Ca-Y series alloy is improved with the increasing Y content because it results in the grain refinement.

3) The addition of more Y in the alloy leads to the formation of a second phase MgY. The second phase MgY can form many of corrosion micro-batteries, which accelerate the corrosion rate of the alloy.

References

- 1 Zreiqat H, Howlett C R, Zannettion A et al. *Mater Res*[J], 2002, 62: 175
- 2 Song G L. *Adv Eng Mater*[J], 2005, 7: 563
- 3 Song G L, Altrens A, StJohn D. *Corros Sci*[J], 1997, 39: 855
- 4 Song G L, Altrens A. *Advanced Engineering Materials*[J], 1999, 1: 11
- 5 Witte F, Kaese V, Haferkamp H et al. *Biomater*[J], 2005, 26: 3557
- 6 Bae D H, Lee M H, Kim K T et al. *J Alloy Compd*[J], 2002, 342: 445
- 7 Sudholz A D, Gusieva K, Chen X B et al. *Corrosion Science*[J], 2011, 53: 2277
- 8 He W W, Zhang E I, Yang K. *Materials Science and Engineering*[J], 2010, 30: 167
- 9 Liu W J, Cao F H, Chang L R et al. *Corrosion Science*[J], 2009, 51: 1334
- 10 Tamar Y, Mandler D. *Electrochemical Acta*[J], 2008, 53: 5118
- 11 Qi Q J, Liu Y b, Yang X H. *Journal of Rare Earth Society*[J], 2002, 20: 428
- 12 Inoue H, Sugahara K, Yamamoto A et al. *Corrosion Science*[J], 2002, 44(3): 603
- 13 Makar G L, Kruger J. *Int Mater Rev*[J], 1993, 38(3): 13824

Mg-2Zn-0.5Ca-Y 系列镁合金在模拟体液和氯化钠溶液中的腐蚀行为和电化学性质

任晓东, 李雪松, 杨悦, 杨友, 吴化

(长春工业大学 先进结构材料教育部重点实验室, 吉林 长春 130012)

摘要: 利用金相显微镜和扫描电镜观察 Mg-2Zn-0.5Ca-Y 系列镁合金的金相结构及表面微观形貌, 使用电化学测试系统测试铸态 Mg-2Zn-0.5Ca-Y 系列镁合金在模拟体液和 3.5%NaCl 溶液中的电化学腐蚀性能, 通过在模拟体液和 3.5%NaCl 溶液中的浸泡试验测试其腐蚀速率。Mg-2Zn-0.5Ca-Y 系列镁合金的微观腐蚀形貌表明, 腐蚀并不均匀, 在浸泡试验后其腐蚀表面有大量的裂纹。稀土 Y 的加入能够提高其耐蚀能力, 并且 Mg-2Zn-0.5Ca-1.0Y 合金的耐蚀性优于其他同系合金。由于腐蚀保护层的存在, 所有在模拟体液和氯化钠溶液中的极化曲线均显示出高容电流。在实验条件下的电化学阻抗曲线均仅有一个循环电容。出现高频率的循环电容是由于稳定的双电层电容的存在。出现低频率的循环电容是由于自由区域的扩散控制膜。

关键词: Mg-2Zn-0.5Ca-Y 系列合金; 腐蚀行为; 极化; 电化学阻抗

作者简介: 任晓东, 男, 1987 年生, 硕士, 长春工业大学材料科学与工程学院, 吉林 长春 130012, 电话: 0431-85716260, E-mail: 279143449@qq.com