One-Step Electrodeposition of Self-Assembled Colloidal Particles: A Novel Strategy for Biomedical Coating

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ABSTRACT: A novel biomedical coating was prepared from self-assembled colloidal particles through direct electrodeposition. The particles, which are photo-cross-linkable and nanoscaled with a high specific surface area, were obtained via self-assembly of amphiphilic poly(γ-glutamic acid)-g-7-amino-4-methylcoumarin (γ-PGA-g-AMC). The size, morphology, and surface charge of the resulting colloidal particles and their dependence on pH, initial concentrations, and UV irradiation were successfully studied. A nanostructured coating was formed in situ on the surface of magnesium alloys by electrodeposition of colloidal particles. The composition, morphology, and phase of the coating were monitored using Fourier transform infrared spectroscopy, energy-dispersive X-ray spectroscopy, scanning electron microscopy, and X-ray diffraction. The corrosion test showed that the formation of the nanostructured coating on magnesium alloys effectively improved their initial anticorrosion properties. More importantly, the corrosion resistance was further enhanced by chemical photo-cross-linking. In addition, the low cytotoxicity of the coated samples was confirmed by MTT assay against NIH-3T3 normal cells. The contribution of our work lies in the creation of a novel strategy to fabricate a biomedical coating in view of the versatility of self-assembled colloidal particles and the controllability of the electrodeposition process. It is believed that our work provides new ideas and reliable data to design novel functional biomedical coatings.

1. INTRODUCTION

Poly(γ-glutamic acid) (γ-PGA) is a kind of bacterial-secreted polypeptide.1–3 γ-PGA and its derivatives have been widely used in many fields, such as drug delivery,3 tissue engineering,4 and wound dressing, because of their biodegradability and biocompatibility.5 γ-PGA has abundant free carboxyl groups that can be modified by functional groups or polymers to prepare desired polymers.6–8 With these excellent properties of γ-PGA, it is desirable to develop various γ-PGA-based multifunctional materials for wider applications.

Self-assembly has been demonstrated to be a powerful technology to make nanoscaled particles from synthetic or natural amphiphilic polymers.9,10 As reported in previous works, amphiphilic γ-PGA derivatives can self-assemble into micellar aggregates with various morphologies, such as spheres, rods, and vesicles.11,12 The attractive advantages of these nanoparticles include a high specific surface area, green nature, nontoxicity, biodegradability, and flexibility in internal or surface functionalities. Although γ-PGA-based particles have potential applications as drug delivery, protein, gene, and vaccine carriers because of these outstanding properties,13–16 research that expands the application of γ-PGA-based particles in biomedical coating systems is lacking.

The substrates used in this work are magnesium alloys, which have physical, mechanical, and biological properties very similar to those of natural bones and are promising implant materials in clinical applications.17,18 However, the widespread use of magnesium alloys has been limited because these alloys are highly susceptible to corrosion in physiological environments, especially during the initial period of implantation.19,20 To improve these weaknesses, the application of coatings is a general strategy to delay initial degradation of the alloy.21 Accordingly, a broad range of surface modification techniques, including spin-coating,22 electroless plating,23 plasma electrolytic oxidation,24 physical vapor deposition,25 spraying26,27 and dipping,28 have been adopted to fabricate coatings.

In addition to all of the techniques mentioned above, electrodeposition has also obtained considerable attention for making coatings because of the controlled integration of its application and the capability to produce uniform films on regularly or irregularly shaped metal substrates at ambient temperature.29–31 Furthermore, the procedure is simple and uses...
 inexpen$$ive equipment. The formation of the coating can be controlled by the deposition time, bath concentration, and applied potential. In previous reports, however, the use of electrodeposition to prepare coatings on magnesium alloys still has been relatively infrequently explored because of the high corrosion rate of magnesium alloys in aqueous solution. Moreover, some nondegradable oxides, such as Mg(OH)$_2$ and MgO, may be produced during the procedure.\(^{32,33}\) In order to avoid these drawbacks, we fabricate a Mg-based biomedical coating by conducting electrodeposition in organic solution. It is another advantage of our investigation, since the magnesium alloy can maintain its original surface composition without a chemical reaction with an electrolyte during the electrodeposition procedure.

The present study attempted to fabricate a biomedical coating of $\gamma$-PGA particles on magnesium alloy by combining the techniques of polymeric self-assembly and electrodeposition. In our previous work,\(^{34,35}\) we used a similar approach to prepare a molecular-imprinted polymeric (MIP) sensor coating in a water system. In this study, we expand the technique to the fabrication of a biomedical coating. The entire strategy is described in Figure 1. A photosensitive amphiphilic poly($\gamma$-glutamic acid)-g-7-

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2. EXPERIMENTAL SECTION

2.1. Materials. $\gamma$-PGA ($M_n = 200000–500000$) was purchased from Wako Pure Chemical Industries Co., Ltd. (Osaka, Japan). 7-Amino-4-methylcoumarin (AMC), 1-ethyl-3-(3-(dimethylamino)propyl)-carboxidimide hydrochloride (EDC·HCl), and 1-hydroxybenzotriazole (HOBt) were obtained from Aladdin Reagent Co., Ltd. (Shanghai, China). Dimethyl sulfoxide (DMSO), absolute ethanol, triethylamine, and acetic acid were purchased from Sinopharm Chemical Reagent Co., China). Dimethyl sulfoxide (DMSO), absolute ethanol, triethylamine, and acetic acid were purchased from Sinopharm Chemical Reagent Co., Ltd. (Shanghai, China). Dulbecco’s modified Eagle’s medium (DMEM) (Gibco), fetal bovine serum (FBS) (Hyclone), and penicillin streptomycin (Gibco) were purchased from WuXi Triolv Biotechnology Inc. (Wuxi, China). All of the reagents and chemicals were used without further purification.

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fabricated by UV irradiation of the coated sample for 1.5 h on each side (UV light was generated by a spot-curing system with a wavelength of 365 nm and a power of 200 W). The coated samples were marked as coated-Mg-CP1.66, coated-Mg-CP2.66, and coated-Mg-CP3.33.

2.5. Physical and Chemical Properties of the Coatings. The surface morphologies of the obtained samples were determined by scanning electron microscopy (SEM) at 2.0 kV (model S-4800, Hitachi, Tokyo, Japan). X-ray photoelectron spectroscopy (XPS) (PerkinElmer, Waltham, MA, USA), and attenuated total reflectance Fourier transform infrared spectroscopy (ATR-FTIR) (Nicolet 6700, Thermo Electron Corp., Madison, WI, USA) were used to investigate the chemical composition of the coatings. The phase compositions of the prepared samples were studied by X-ray diffraction (XRD), and the instrument (D8, Bruker, Karlsruhe, Germany) was operated from 3° to 90° in steps of 0.02° at a scan rate of 4° deg min⁻¹.

2.6. Corrosion Resistance of the Coating. Electrochemical tests were performed to evaluate the anticorrosion properties of the colloidal particle coating and bare magnesium alloys in simulated body fluid (SBF) at 37 ± 0.5 °C. The constituents of the SBF solution were the same as in a previous report. A three-electrode cell was used to carry out the measurements. The sample, a platinum plate, and a saturated calomel electrode (SCE) were used as the working, counter, and reference electrodes, respectively. The scanning range was from −0.6 to 0.6 V of the open-circuit potential at a rate of 1 mV s⁻¹.

3. RESULTS AND DISCUSSION

3.1. Preparation of γ-PGA-g-AMC Copolymer. AMC was grafted to γ-PGA chains through the reaction between carboxylic groups on the linear γ-PGA chains and the amino groups of AMC. In our work, EDC-HCl and HOBT were used to activate carboxylic groups, which were then attached to the amino groups of AMC to produce the γ-PGA-g-AMC copolymer. The ¹H NMR spectrum of the resulting copolymer is shown in Figure S1 in the Supporting Information. The degree of grafting was calculated by comparing the peak areas of aromatic ring protons in the coumarin group and CH protons in the PGA main chain; the results showed that nearly 20% of AMC was grafted to the γ-PGA chains.

3.2. Self-Assembly of Colloidal Particles. To prepare the self-assembled colloidal particles, absolute ethanol was gradually added dropwise to γ-PGA-g-AMC copolymer solutions. The changes in the absorbance of the different solutions were studied using UV–vis measurements. The morphology of colloidal particles in the dried state was observed by TEM.

Figure 2 shows the absorbance changes when ethanol was added to the γ-PGA-g-AMC solutions. As the results show, the absorbance of all of the samples initially decreased with increasing ethanol amount because the copolymer solutions were diluted by the addition of ethanol. When the ethanol content attained a certain value, the absorbance began to increase, indicating that the γ-PGA-g-AMC copolymer chains started to aggregate and form self-assembled particles, except for the sample with an initial copolymer concentration of 1 mg mL⁻¹, as the driving force for the self-assembly in the 1 mg mL⁻¹ sample was too low to form colloidal particles. The ethanol content at this point is defined as the critical ethanol content (CEC). After the addition of more ethanol, the absorbance continued to increase because of the aggregation of self-assembled particles. In order to make a uniform distribution of colloidal particles, twice the volume equivalent of ethanol with respect to the initial copolymer solution was used to make the self-assembled colloidal particles. In the self-assembly process, the driving force is mainly related to the amount of precipitating agent (ethanol) added and the initial copolymer concentration. Since the concentration of 1 mg mL⁻¹ is dilute, the interaction and chain entanglement between the copolymer segments are very weak compared with other higher concentrations. Therefore, when ethanol was added, the copolymer chain could not aggregate effectively and form particles.

The morphology and size of colloidal particles in the dried state were observed by TEM. As shown by the picture in the Figure 2 inset, the self-assembly of γ-PGA-g-AMC occurred and the aggregated colloidal particle size was almost 100 nm in the
dried state. Compared with the size determined by DLS ($R_h = 115.6 \text{ nm at } 1.66 \text{ mg mL}^{-1}$; see Figure 3), the TEM results show a smaller particle size. This result is understandable because the swelling of the particles in solution comes from the larger amount of absorbed solvent. As solvent is evaporated, the particles would shrink. These results imply that adding ethanol as a precipitating agent could induce self-assembly of the γ-PGA-g-AMC copolymer. The mechanism can be explained as follows: the copolymer molecular chains contain abundant −COOH groups, which attract each other by hydrogen bonding, thus decreasing their affinity for ethanol. When precipitating agent is added, the chain segments containing −COOH groups start to collapse into insoluble complexes because of interpolymer hydrogen bonding. However, those containing −COO− groups experience mutual electrostatic repulsion and are mainly distributed in the shell of resulting particles during the process of ethanol addition, thus forming stable colloidal particles.

3.3. Effect of Initial Copolymer Concentration and pH. Self-assembly is a dynamic equilibrium process. The polymer concentration can affect the interaction of macromolecular chains and the aggregate degree of self-assembled particles in the resulting colloidal particles. As shown in Figure 3, the average hydrodynamic radii ($R_h$) of 0.33, 1.00, 1.66, 2.66, and 3.33 mg mL$^{-1}$ colloidal particles fabricated from 2, 3, 5, 8, and 10 mg mL$^{-1}$ initial γ-PGA-g-AMC copolymer solutions, respectively.

![Figure 3. Average sizes and size distributions of 0.33, 1.00, 1.66, 2.66, and 3.33 mg mL$^{-1}$ colloidal particles fabricated from 2, 3, 5, 8, and 10 mg mL$^{-1}$ initial γ-PGA-g-AMC copolymer solutions, respectively.](image)

As shown in Figure 3, with increasing irradiation time (365 nm, total intensity 200 W, 4 mL of solution), $R_h$ decreased from 115.6 to 71.3 nm. The photoinduced decrease in average $R_h$ due to photo-cross-linking is around 52%. It is implied that under UV irradiation, the interchain photo-cross-linking leads to shrinkage of whole colloidal particles. A similar diameter-decreasing effect of initial copolymer concentration and pH on (a) the surface charge and (b) the average hydrodynamic radius of the colloidal particles. The concentration of the particle solution was 1.66 mg mL$^{-1}$.

![Figure 4. Effect of pH on (a) the surface charge and (b) the average hydrodynamic radius of the colloidal particles. The concentration of the particle solution was 1.66 mg mL$^{-1}$.](image)

As shown in Figure 5, the variations of surface charge and average $R_h$ of colloidal particles as a function of pH. Obviously, the surface charge became more negative with increasing pH from 6 to 10. As the pH increased from 6 to 9.5, the average $R_h$ increased from 95.5 to 227.3 nm, and then the particle size dramatically increased for pH >9.5. These results indicate that the particle size and surface charge in ethanol are both influenced by the pH of the solution. In our work, the self-assembled colloidal particles can keep their particle state at pH 6–10. For pH <6, the samples formed precipitates as a result of protonation of −COO− groups. For pH >10, the solutions became transparent, demonstrating that the particles changed into soluble molecular chains or a random coil state. These results demonstrate that electrodeposition of colloidal particles can be conducted at pH 6–10 in order to fabricate the nanostructured coating.

3.4. Photoresponsive Properties. It is well-known that direct irradiation ($\lambda > 310 \text{ nm}$) of coumarin and its derivatives leads to photochemical dimerization. Because the colloidal particles contain many coumarin units that are photoresponsive to alternative irradiation with UV light, the photo-cross-linking properties of coumarin are expected to be reflected in the self-assembled particles. To monitor this property, the photosensitivity of the colloidal particles was demonstrated using DLS.

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![Figure 5. Hydrodynamic radius and distribution of the colloidal particles irradiated with a spot curing system (200 W, 365 nm) at different times. The concentration of the colloidal particle solution was 1.66 mg mL$^{-1}$.](image)
phenomenon has also been obtained in other coumarin-containing self-assembled systems.\textsuperscript{45,46} It is believed that this photo-cross-linking property of the particles can also affect the properties of coating systems. As reported in previous work,\textsuperscript{47} UV cross-linking of colloidal particles at the oil–water interface can make a dense shell of colloidosomes and lock the interfacial particles in a Pickering or Mckering emulsion. Thus, we hypothesize that when the photo-cross-linkable colloidal particles are electrodeposited on the Mg–Ca alloy surface, a dense coating surface can be fabricated by further UV irradiation of the electrodeposited particles.

### 3.5. Electrodeposition of Colloidal Particles
\(\gamma\)-PGA is a naturally biodegradable and nontoxic biomaterial. It consists of alternating units of D-glutamic acid and L-glutamic acid.\textsuperscript{48} The negative charge of the \(\gamma\)-PGA colloidal particles in solution is attributed to deprotonated –COOH groups.\textsuperscript{59} The nanostructured coating on the magnesium alloy can be obtained by anodic electrodeposition in ethanol. The proposed mechanism of coating formation is based on the classical Derjaguin–Landau–Verwey–Overbeek (DLVO) theory of colloidal stability.\textsuperscript{50,51} This theory states that the total pair interaction between colloidal particles consists of two parts, the Coulombic double-layer repulsion and the van der Waals attraction. As reported by Lyklema,\textsuperscript{52} these forces promote the stability of colloidal particles in nonaqueous media. The DLVO theory describes the potential energy curve for pair interaction, as shown in Figure S2 in the Supporting Information. We can assume that coagulation of the colloidal particles would occur when the energy barrier is overcome. When an electric field is applied, the negatively charged \(\gamma\)-PGA-\(g\)-AMC colloidal particles would migrate, and this migration can induce a particle concentration gradient around the magnesium alloy surface. As determined by previous work,\textsuperscript{53} the potential energy peak decreases as the electrolyte concentration increases, and thus, there exists a critical electrolyte concentration (flocculation value) for coagulation in an applied electric field. While the energy barrier between colloidal particles disappears around the Mg–Ca anode in the applied field, coagulation becomes possible and the nanostructured coating forms. As shown in Figure S3a–c in the Supporting Information, some particles are immobilized onto the substrate at short deposition times. This behavior indicates that charged colloidal particles can move toward the anode in the electric field within a given time. After 15 min, the nanostructured surface was formed as a result of accumulation of colloidal particles (Figure S3d). This demonstrates that coating formation can be controlled by electrodeposition.

FTIR and XPS were used to confirm the presence of nanostructured coatings on the Mg–Ca alloy. In the FTIR spectrum of the nanostructured coating (spectrum c in Figure 6A). The characteristic peak at 3258 cm\(^{-1}\) is related to O–H stretching vibrations of –COOH groups. The peaks at 1725 and 1645 cm\(^{-1}\) are attributed to C=O stretching vibrations of the –COOH and –CO—NH– groups, respectively. The observed band at 1545 cm\(^{-1}\) is assigned to C≡C. The peak at 1155 cm\(^{-1}\) is attributed to C–N stretching. These bands are well in accordance with vibrational peaks of native colloidal particles (spectrum b in Figure 6A). As expected, no such peaks are seen on bare metal substrates devoid of the nanostructured coating (spectrum a in Figure 6A). Moreover, the observed band at 1006 cm\(^{-1}\) in the FTIR spectrum of the nanostructured coating is related to the absorption of triethylamine. The XPS results imply that main elements on the surface of the coated sample are C, N, and O (Figure 6B). The results indicate the formation of nanostructured coatings on the surface of magnesium alloys.

The surface microstructure of the coated samples was investigated by SEM (Figure 7). The results show that the morphologies of the nanostructured coatings are uniform, and some colloidal particles can be clearly seen before UV irradiation (Figure 7A1–C1). This indicates that homogeneous nanostructured coatings are fabricated by electrodeposition of self-assembled colloidal particles. After UV irradiation for 1.5 h, the surfaces of the cross-linked coatings (Figure 7A2–C2) are denser and smoother as a result of the photo-cross-linking properties of self-assembled particles. Photo-cross-linking of the particles is demonstrated to improve the homogeneity and also the sealability of the nanostructured coating. Generally, a denser coating on the magnesium alloy provides better physiological corrosion resistance of the substrate. Therefore, photo-cross-linking of the nanostructured coating supposedly contributes to the anticorrosion performance of the magnesium alloy, and we will discuss this topic in detail in section 3.7.

### 3.6. XRD Analyses of the Coated Samples
The XRD patterns of the bare and coated magnesium alloys are shown in Figure 8. The intensity of the Mg phase of the bare substrate (curve a) is very strong. As the concentration of the colloidal particle electrolyte for electrodeposition increases, the intensity of the Mg phase of the coated sample decreases accordingly (curves b–d). The increase in the electrolyte concentration is suggested to result in the increase in deposited mass. In addition,
the phases of oxides, such as Mg(OH)$_2$ and MgO, are not observed in the XRD patterns. This result implies that non-biodegradable oxides are barely formed in our organic systems. Furthermore, similar experiments were also performed in water systems, and the Mg(OH)$_2$ phase was observed in the XRD patterns (Figure S4 in the Supporting Information). These results obviously identify that the original properties of the magnesium alloys are maintained throughout the electrodeposition process in our organic systems. This observation is good news for magnesium alloys as implant materials because non-biodegradable oxides are not formed and the original biocompatibility and biodegradability of the magnesium alloys have been protected.

3.7. Corrosion Properties of Samples. The protective properties of the coatings were evaluated through potentiodynamic polarization tests in SBF: the more positive the corrosion potential ($E_{\text{corr}}$) (or the lower the polarization current), the better the corrosion resistance. Generally, the cathodic polarization curves are assumed to represent cathodic hydrogen evolution through water reduction, whereas the anodic polarization curves represent the dissolution or destruction behaviors of the protective coating.$^{34}$

As shown in Figure S5 in the Supporting Information, $E_{\text{corr}}$ for the nanostructured-coating-modified sample is more positive than that for the bare magnesium alloy. The corrosion current density ($I_{\text{corr}}$) of the coated magnesium alloy correspondingly decreases. In addition, the corrosion resistance is further enhanced by 1.5 h of UV irradiation because photo-cross-linking induces the dense and smooth surface of the coated sample (as shown in the SEM measurements). These results indicate that photo-cross-linking of nanostructured coatings provides good corrosion resistance to the magnesium alloy.

The polarization curves for the bare magnesium alloy, coated-Mg-CP$_{1.66}$, coated-Mg-CP$_{2.66}$, and coated-Mg-CP$_{3.33}$ after 1.5 h of UV photo-cross-linking on each side are shown in Figure 9.

<table>
<thead>
<tr>
<th>sample</th>
<th>$E_{\text{corr}}$ (V)</th>
<th>$I_{\text{corr}}$ (A cm$^{-2}$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>bare magnesium alloy</td>
<td>$-1.80$</td>
<td>$8.71 \times 10^{-4}$</td>
</tr>
<tr>
<td>coated-Mg-CP$_{1.66}$</td>
<td>$-1.52$</td>
<td>$3.31 \times 10^{-4}$</td>
</tr>
<tr>
<td>coated-Mg-CP$_{2.66}$</td>
<td>$-1.30$</td>
<td>$9.98 \times 10^{-4}$</td>
</tr>
<tr>
<td>coated-Mg-CP$_{3.33}$</td>
<td>$-1.03$</td>
<td>$4.63 \times 10^{-4}$</td>
</tr>
</tbody>
</table>

The $E_{\text{corr}}$ and $I_{\text{corr}}$ values are summarized in Table 1. The results imply that all of the coatings exhibit a better protective function as reflected from their more positive $E_{\text{corr}}$ and lower $I_{\text{corr}}$ values. Compared with the value for the bare magnesium alloy ($-1.80$ V), $E_{\text{corr}}$ increases by 15.6% for coated-Mg-CP$_{1.66}$, 27.8% for coated-Mg-CP$_{2.66}$, and 42.8% for coated-Mg-CP$_{3.33}$. The shift of $E_{\text{corr}}$ in the positive direction can be attributed to the protective properties provided by photo-cross-linking of the nanostructured coatings formed on magnesium alloys. $I_{\text{corr}}$ decreases by 62.0%, 88.5%, and 94.7% for the coated-Mg-CP$_{1.66}$, coated-Mg-CP$_{2.66}$, and coated-Mg-CP$_{3.33}$ samples, respectively.
CP₂.₆₆₆ and coated-Mg-CP₃.₃₃₃ samples, respectively. These results illustrate that the protective properties of the nanostructured coatings can be adjusted by controlling the concentration of the colloidal particle electrolyte in the electrodeposition process. These results support the conclusion that nanostructured coatings fabricated by electrodeposition of the colloidal particle electrolyte can act as protective layers and improve the initial anticorrosion properties of magnesium alloy substrates in an SBF environment.

3.8. Cytotoxicity Assay. The morphology of NIH-3T3 cells was examined by inverted optical microscopy with a digital camera. Figure 10 shows that viable cells were observed on culturing plates after 2 days, and higher magnification images are shown in Figure S6 in the Supporting Information. Compared with the control group, the majority of NIH-3T3 cells cultured in the medium with immersion extracts exhibit a healthy morphology of cells with an elongated and flattened spindle shape. More importantly, the cells cultured with immersion extracts of the coated samples show good cell spread and cell densities in contrast to those of the uncoated one, suggesting that better cytocompatibility is obtained with coated magnesium alloys.

The cell viability for different samples was evaluated by the MTT assay using NIH-3T3 cells through an indirect method in which immersion extracts were used to conduct the assay. The MTT assay is based on the ability of a mitochondrial dehydrogenation enzyme in viable cells to cleave the tetrazolium rings of pale-yellow MTT and form formazan crystals with a dark-blue color. Figure 11 shows cell viability after 2 days of incubation with immersion extracts of uncoated and coated Mg−Ca samples; the group cultured in DMEM without immersion extracts was used as a control. The results demonstrate that cells cultured in the extracts show significant differences (p < 0.05) in the average cell viability relative to the control. Also, all of the extracts of coated samples (c−e) were well-tolerated by the NIH-3T3 cells with average cell viabilities ranging from nearly 112.2% to 151.5%, in contrast to that of control group. Instead, the cells cultured with immersion extracts of uncoated Mg−Ca alloy showed relatively low viability. Therefore, all of the coated samples exhibited low cytotoxicity toward NIH-3T3 cells, revealing improved cytocompatibility after surface treatment. As reported by previous work, the viability of cells grown in extracts is strongly correlated with the corrosion products of the magnesium alloy sample. The main corrosion products of immersion extracts used in the cell experiment are Mg²⁺ and OH⁻; the ion contents would influence the cellular ion balance, cellular metabolism, and other features, thus influencing the viability of cells. The low viability of cells cultured with immersion extracts of uncoated Mg−Ca alloy may be caused by the damage due to higher Mg²⁺ and OH⁻ concentrations produced through the rapid corrosion process of magnesium alloy, while the higher viability of cells cultured with those of coated magnesium alloys is probably due to the fact that appropriate ion contents can promote the cellular metabolism and growth.

4. CONCLUSION

In this work, we have presented a new and efficient way to fabricate biomedical coatings via electrodeposition of self-assembled colloidal particles of an amphiphilic photo-cross-linkable copolymer on magnesium alloys. The resulting self-assembled colloidal particles are responsive to initial copolymer concentration, pH, and UV irradiation, as indicated by DLS
analysis, TEM characterization, and zeta potential measurements. In addition, the obtained coatings show more favorable corrosion resistance, and the coated samples exhibit low cytotoxicity toward NIH-3T3 cells compared with bare magnesium alloy. The whole strategy, which combines the advantages of self-assembled particles and electrodeposition, is novel, simple to control, and inexpensive. More importantly, this work demonstrates a general and promising approach for the development of functional biomedical coatings for not only magnesium alloys but also a wide range of metals using in medical applications.

**ASSOCIATED CONTENT**

3 Supporting Information

1H NMR spectrum of γ-PGA-γ-AMC, illustration of the electrodeposition mechanism, SEM images of nanostructured coatings obtained at different deposition times, XRD pattern of a coated sample fabricated in water solution, polarization curves of the coated-Mg-CP1.66 sample before and after UV irradiation, and higher-magnification images of NIH-3T3 cells. This material is available free of charge via the Internet at http://pubs.acs.org.

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Notes

The authors declare no competing financial interest.

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