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Degradation and biocompatibility of a series of strontium substituted hydroxyapatite coatings on magnesium alloys

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There has been a surge in the research on magnesium (Mg) alloys as a promising selection for biomaterials application. However, as a foremost drawback, the fast degradation of Mg alloys limits its clinical use. In this study, a series of Sr-HA coatings with the Sr content ranging between 10–100% were prepared on Mg alloys, in order to control the degradation and enhance the osteoblast response. Microstructure analysis indicated the formation of $\text{Ca}_{10-x}\text{Sr}_x(\text{PO}_4)_6(\text{OH})_2$ coatings with the thickness ranging between 28–35 μm . The degradation results suggested that an increase in the Sr content in the coatings led to the decreasing degradation rate of the Sr-HA coated Mg. 100% Sr-HA coatings provided the best corrosion protective effect with nearly no hydrogen evolution during 10 days' immersion in Hank's solution. The *in vitro* cell biocompatibility was evaluated with MC3T3-E1 osteoblasts using the extract assay. In each case the released Sr affected the osteoblast proliferation and the expression of osteogenesis markers including, ALP, Col-I and RUNX2, in a Sr concentration-dependent manner. These results suggest that Sr-HA coating is a promising combination for controlling the degradation and enhancing the cytocompatibility of Mg alloys. The degradation and osteoblast response could be simply controlled through the adjustment of Sr content in the coatings.

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1. Introduction

Magnesium (Mg) and its alloys show excellent compatibility with human bones in density, elastic modulus and compressive strength.^{1–3} Mg is essential for human metabolism and is naturally present in bone tissue. In addition, Mg can be gradually degraded in the human body, which makes it a promising candidate for orthopedic implants.^{1,4,5} In spite of these unique benefits, Mg based implants are of limited use in clinical applications. The main disadvantage is the fast degradation in the physiological environment and the associated side effects, including gas bubbles, local alkalization, fast decay of strength.^{6–8}

Applying biocompatible and protective coatings, such as hydroxyapatite (HA), which is the main inorganic component of bones and has been used for decades in medicine and dentistry, onto magnesium alloys is a practical option to control the biodegradation process.^{8–11} HA coatings could reduce the degradation of magnesium alloys with certain amounts of success.^{8,12–15}

Strontium (Sr) is believed to play an important role in the treatment of osteoporosis and enhancement of bone remineralization as it is associated with a reduction of bone resorption and an increase in the formation of new bone.^{7,16,17} It is well documented that the presence of Sr^{2+} ions has the dual benefit of promoting pre-osteoblastic cell replication and inhibiting the activity of osteoclasts.^{18–21} Strontium ranelate (Protelos®, Servier) is currently used in the treatment of post-menopausal osteoporosis, although its prescription has been recently restricted to the acute medical cases because of cardiovascular side effects.²² Additionally, Sr can replace the lattice sites of Ca in HA to form a continuous solid-solution (Sr_xHA , x is fraction of substitution) up to full substitution (Sr-apatite) with improved mechanical properties and biological performance.^{17,23–25} There have been intensive studies to incorporate Sr into HA cements^{16,26–28} or coatings^{24,29–32} for better bone response. Furthermore, the degradation and resulted osteoblast response of Sr_xHA exhibit a dose dependent behaviour.^{16,17} Sr incorporating destabilizes the structure of HA and increased the solubility and degradation of Sr_xHA with x .¹⁷ With variations in degradation rate of Sr_xHA , the micro-environment are thereby affected. Particularly Sr availability through degradation inevitably changes, which in turn influence the metabolic activity of osteoblasts.

The Sr doses would affect the properties of Sr-HA coatings such as microstructure, solubility, degradation and cell interaction with materials. Therefore, the main aim of the present

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study was to highlight the influence of Sr doses of Sr-HA coated Mg alloys on their morphology, *in vitro* degradation, osteoblast proliferation and differentiation.

2. Materials and methods

2.1 Materials preparation

AZ31B magnesium alloys were cut into $20 \times 10 \times 2 \text{ mm}^3$ pieces, mechanically polished with SiC paper up to 2000 grit and cleaned ultrasonically with acetone, ethanol and deionized water for 5 min each. Prior to the electrodeposition, the samples were pre-treated with 40% HF solution for activation, and then rinsed in distilled water and dried at room temperature.

2.2 Electrodeposition

The electrolyte for deposition was prepared by mixing a solution of $\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$, $\text{Sr}(\text{NO}_3)_2$, $\text{NH}_4\text{H}_2\text{PO}_4$, NaNO_3 and H_2O_2 . The concentrations of $\text{NH}_4\text{H}_2\text{PO}_4$, NaNO_3 and H_2O_2 were 0.025 mol L^{-1} , 0.1 mol L^{-1} and 10 mL L^{-1} . The $\text{Ca}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$, $\text{Sr}(\text{NO}_3)_2$ concentrations were adjusted so that the Sr/(Sr + Ca) ratios of the electrolytes were 0%, 10%, 20%, 50%, and 100%, respectively. The temperature was maintained at $60 \text{ }^\circ\text{C}$ using water bath.

The electrodeposition was conducted in a standard electrolysis cell with three electrodes in which the AZ31B alloy, the graphite plate and a saturated calomel electrode (SCE) acting as working electrode, counter electrode and reference electrode, respectively. Deposition was performed at -2.5 V/SCE for 70 min through an electrochemical workstation (CHI660E). After deposition, the samples were further treated in 1 mol L^{-1} NaOH solution at $80 \text{ }^\circ\text{C}$ for 2 h.

2.3 Characterization

The phase of Sr-HA coatings was analyzed using a Bruker D8 multipurpose X-ray diffractometer (XRD) with a scanning range (2θ) of $20\text{--}70^\circ$, and a scanning speed of 4° min^{-1} . An environmental scanning electron microscopy (ESEM, FEI Quanta 250FEG) equipped with an energy dispersive spectrometry (EDS) was then used to examine the surface and cross-sectional morphology of the samples as well as the elemental compositions of the coatings.

2.4 Immersion tests

Immersion tests were carried out in Hank's solution (8.00 g NaCl , 0.40 g KCl , 0.14 g CaCl_2 , 0.35 g NaHCO_3 , $0.20 \text{ g MgSO}_4 \cdot 7\text{H}_2\text{O}$, $0.12 \text{ g Na}_2\text{HPO}_4 \cdot 12\text{H}_2\text{O}$, $0.06 \text{ g KH}_2\text{PO}_4$) according to the ASTM-G31-72. The Mg samples were coated with Sr-HA coatings on all sides. The uncoated and Sr-HA coated samples were completely immersed in Hank's solution. The pH value was adjusted to 7.40 ± 0.5 and the temperature was maintained at $37 \pm 0.5 \text{ }^\circ\text{C}$. The hydrogen evolution behaviour was measured during the immersion. After 10 days immersion, the samples were removed from the solution, gently rinsed with distilled water, followed by drying at room temperature. Surface morphologies of the samples were analysed by ESEM and EDS. The ion concentrations were determined by inductively coupled

plasma optical emission spectrometry (ICP-OES; Leeman, Profile).

2.5 Electrochemical tests

The electrochemical corrosion was carried out in Hank's solution using a CHI660E electrochemistry workstation. A three-electrode cell was used for electrochemical measurements. The counter electrode was a platinum plate and the reference electrode was a saturated calomel electrode. The sample was pre-incubated in Hank's solution for 30 min to obtain a relatively steady open circuit potential (OCP). Afterwards, the potentiodynamic polarization test was conducted with a scan rate of 1.0 mV s^{-1} with the initial potential at about 300 mV below the OCP.

2.6 Osteoblast proliferation

The proliferation of osteoblast-like cells (MC3T3-E1, ATCC CRL-2593) was evaluated using Cell Counting Kit-8 (CCK-8, Dojindo) assay according to standard ISO 10993-5. Prior to the cell experiments, samples were sterilized using ultraviolet radiation for at least 2 h. Extracts were prepared with a surface area to Dulbecco's modified Eagle's medium (DMEM, low glucose, Gibco) ratio of 1 mL cm^{-2} in a humidified atmosphere with 5% CO_2 at $37 \text{ }^\circ\text{C}$ for 72 h incubation. The extract was centrifuged twice, and then the supernatant solution was withdrawn and filtered with milipore filters ($0.22 \text{ }\mu\text{m}$). The pH values and osmolality of the extracts were measured. Ion concentrations were determined by ICP-OES. The extract was diluted to 50%.

MC3T3-E1 were cultured in DMEM, 10% fetal bovine serum (FBS, Gibco), 100 U mL^{-1} penicillin and $100 \text{ }\mu\text{g mL}^{-1}$ streptomycin at $37 \text{ }^\circ\text{C}$ in a humidified atmosphere of 5% CO_2 . Cells were seeded in 96-well cell culture plates at 3×10^4 cells per mL and incubated for 24 h to allow attachment. The medium was then replaced with different extracts. After 1, 3 and 5 days, the extracts were replaced with DMEM, to which was then added 10% CCK-8 solution. After incubation for a further 3 h, the absorbance was read at 450 nm by a microplate reader (Vari-oskan LUX, Thermo), and expressed as the optical density (OD). Cell morphologies were observed by fluorescence staining. Cells were fixed with 4% polyoxymethylene and permeabilized with 0.1% Triton X-100. Cellular actin and nuclei were stained with FITC phalloidin and DAPI simultaneously before observing by a fluorescence microscope (Olympus).

2.7 ALP activity

MC3T3-E1 cells were seeded at a density of 3×10^4 cells per mL and incubated for 24 h, at which time the medium was replaced with the extracts. After 5 days, the cells were lysed in 0.2 mL of 0.1% Triton X-100, shaken and conducted two standard freeze-thaw cycles. The alkaline phosphatase (ALP) activities were determined by a colorimetric assay using an ALP assay kit (Beyotime) and the absorbance was measured at 405 nm. The intracellular total protein levels were determined using the bicinchoninic acid (BCA) protein assay kit (Beyotime) simultaneously. The ALP activity was finally normalized to the total



protein content correspondingly and was expressed as $\mu\text{mol min}^{-1} \text{mg}^{-1}$ protein.

2.8 Real-time PCR assay (QPCR)

MC3T3-E1s were cultured in 6-well plates at 5×10^5 cells per well and incubated for 24 h. The medium was then replaced with 2 mL extracts and incubated for 5 days. Total RNA was extracted by using Trizol reagent (Gibco) and used as template (2 μg) for reverse-transcribed into cDNA according to the manufacturer's instructions (Takara). To detect gene expression, quantitative real-time polymerase chain reaction (QPCR) was performed using the SYBR Premix Ex Taq kit (Takara, Shiga, Japan) according to the manufacturer's instructions and an iQ5 quantitative real-time PCR System (Bio-Rad Laboratories, Berkeley, CA). Forward and reverse primer sequences of alkaline phosphatase (ALP), collagen type I (COL-1) and runt-related transcription factor 2 (RUNX2) are shown in Table 1. All reactions were pre-denatured at 95 °C for 3 min, followed by 40 cycles at 95 °C for 10 s and 55 °C for 30 s. After the PCR reaction, a melting curve was generated to confirm the PCR product's specificity and identity. Triplicate reactions were performed, and the relative fold-change of gene expression was determined by normalizing to GAPDH and calculating the $2^{-\Delta\Delta C_T}$.

2.9 Statistical analysis

All data were expressed as means \pm standard deviation. Statistical analysis was conducted by one-way analysis of variance (ANOVA). A *p* value of less than 0.05 was considered to be significant.

3. Results and discussion

3.1 Coating characterization

Fig. 1 depicts the XRD spectra of the Sr-HA coatings on Mg alloys. XRD results demonstrate the formation of $\text{Ca}_{10-x}\text{Sr}_x(\text{PO}_4)_6(\text{OH})_2$ phase in all coatings, with the peaks corresponding closely to those of hydroxyapatite (JCPDS no: 72-1243) and pure strontium hydroxyapatite (JCPDS no: 70-1511). The diffraction peaks shift to lower 2θ values with increasing amount of Sr addition in the Sr-HA coatings, which is attributed to the larger ionic radius of Sr.^{17,22} The crystallinity of Sr-HA phase deteriorates with increasing Sr up to 50%, as judged by the broaden diffraction peaks. When the Ca is completely substituted by Sr, $\text{Sr}_5(\text{PO}_4)_3(\text{OH})$ is formed and the crystallinity increases again evidenced by the sharp XRD peaks.

The Sr-HA coatings show similar FTIR spectra (Fig. 2). The characteristic peaks due to PO_4^{3-} are clearly seen at 1025 cm^{-1} ,

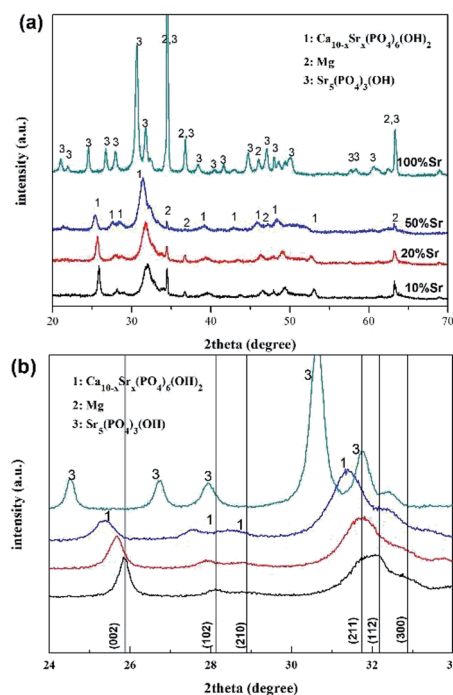


Fig. 1 XRD patterns of the Sr-HA coatings obtained from the electrolytes with Sr/(Sr + Ca) ratios ranging 10–100%. (a) Full diffraction spectra (20–70°) and (b) detailed spectra (24–35°) of XRD patterns.

600 cm^{-1} and 569 cm^{-1} .²² The differentiation between various coatings locates in the carbonate adsorption band at around $1351\text{--}1540 \text{ cm}^{-1}$,³³ the intensity of which increases with increasing Sr content up to 50%. For 100% Sr substitution, the intensity of the carbonate band is significantly reduced. In addition, the bands located at 3442 cm^{-1} and 1643 cm^{-1} are attributed to the stretching and bending mode of adsorbed water (H_2O) molecules.²² The absorbed water molecules are bonded with the surface phosphate groups and form an amorphous hydrated layer.³⁴ The increasing peak intensity is also observed with increasing Sr up to 50% and is reduced for 100% Sr-HA coating.

The surface morphologies of the coatings with different Sr contents are shown in Fig. 3. When the Sr content varies in the range of 10–50%, the coatings are constituted by small size spheroids and exhibit some apparent porosity (Fig. 3a1–c1). The coating with 50% Sr exhibits more compact, uniform surface morphology. For 100% Sr-HA coating, it is composed of well packed flower-like crystals (see Fig. 3d1). The atomic strontium proportion is quantified by EDS for each coating to determine the substitution percentage by calculating the Sr/(Ca + Sr) ratio

Table 1 Primer sequence for quantitative real-time PCR

Gene	Forward primer sequence 5'–3'	Reverse primer sequence 5'–3'
ALP	AACGTGGCCAAGAACATCATCA	TGTCATCTCCAGCCGTGTC
COL-1	GCCTCCGAGAACATCACCTA	GCAGGGACTTCTTGAGGTTG
RUNX2	CCATAACGGTCTTCACAAATCCT	TCTGTCTGTGCCCTTCTGGTTC
GAPDH	GGCACAGTCAAGGCTGAGAATC	ATGGGTGGTGAAGACGCCAGTA



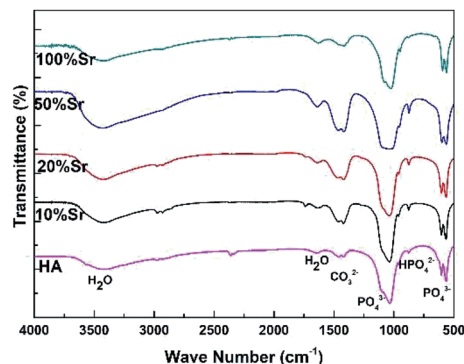


Fig. 2 FTIR spectra of the Sr-HA coatings obtained from the electrolytes with Sr/(Sr + Ca) ratios ranging 10–100%.

(Fig. 3e) and (Ca + Sr)/P ratio (Fig. 3f). The obtained coating contains a lower strontium content than the designated Sr/(Sr + Ca) ratio in the electrolyte. They have the (Ca + Sr)/P ratios of ranging from 1.73 to 1.82 for 10–50% Sr substituted Sr-HA coating. The higher (Ca + Sr)/P ratio can possibly be explained by the carbonate substitution in the apatite lattice (Fig. 2), which has been observed previously for carbonated HA.^{17,35} Fig. 4a and b shows the typical cross-section morphology and elements profiles of the coating. The coating thickness ranges from 28–35 μm . No apparent interface gap between the coating and magnesium substrate is observed, indicating the coating adhere well on the substrate.

3.2 *In vitro* degradation

The primary concern of these Sr-HA coatings is their protection efficacy of magnesium degradation. Thus the *in vitro*

degradation of the experimental samples was characterized as a function of immersion time in Hank's, as shown in Fig. 5. The hydrogen evolution volume and the amount of Mg ions released decrease with increasing Sr content (Fig. 5a and c). The increment of pH values is significantly slowed down with Sr-HA coating. The pH increases to 10.0 after 5 d immersion for 10–50% Sr-HA coated samples. While for 100% Sr-HA coated samples, the pH only increases 9.0 after 10 d immersion in Hank's solution (Fig. 5b). The potentiodynamic polarization curves and the corresponding electrochemical parameters are shown in Fig. 6. It is evident that anodic kinetics of all the Sr-HA coated samples had been reduced, with the anodic polarization current approximately one order of magnitude lower than the uncoated Mg. The restrictive effect on anodic kinetics leads to more noble corrosion potential for the coated samples compared with the uncoated control, indicating the coatings provide protection from corrosion. The calculated corrosion current density values are a function of the Sr/(Sr + Ca) ratio in the coatings. The 100% Sr-HA coating reduced the corrosion current density to $17.55 \mu\text{A cm}^{-2}$, a $\sim 53\%$ reduction as compared to the 10% Sr-HA coated sample and a $\sim 93\%$ reduction as compared to the uncoated sample. As such, the Sr content has a critical influence on the corrosion protectiveness, in which the 100% Sr-HA coating demonstrates the best inhibiting ability from corrosion (Fig. 5d).

After 10 d immersion in Hank's solution, the SEM observations indicate fewer small-sized spheroids for 10% Sr-HA coatings after immersion than the as-prepared coatings. Massive laminar precipitation is deposited and show denser coating morphology, while some micro cracks are still visible (Fig. 3a2). The EDS reveals that the Sr proportions in the coatings slightly increase to 7.4 at% (Fig. 3e). 50% Sr-HA coatings exhibit much

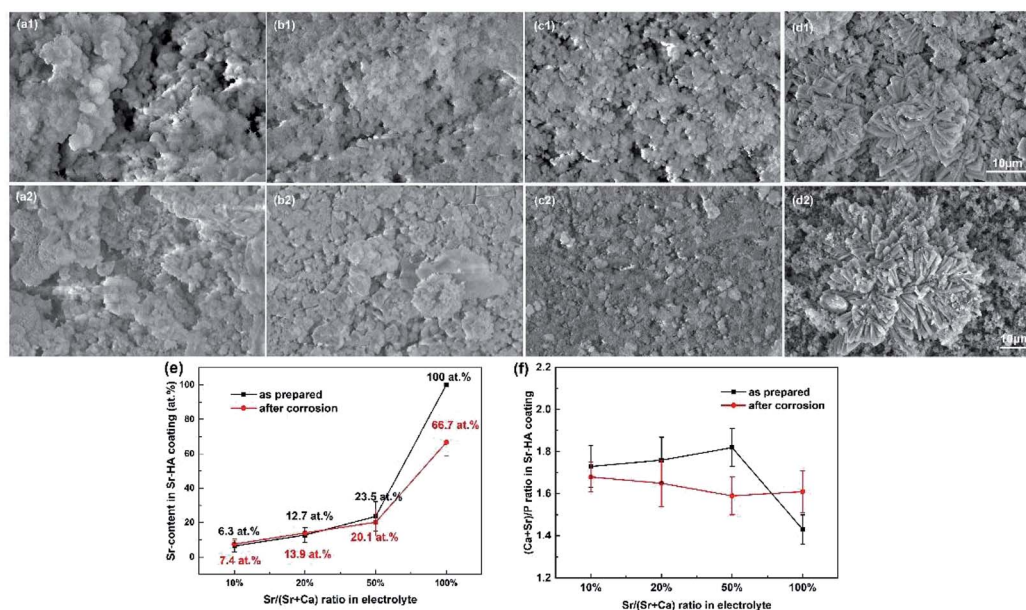


Fig. 3 Surface observations of the Sr-HA coatings with various Sr/(Sr + Ca) ratios of (a1) 10%, (b1) 20%, (c1) 50%, (d1) 100%. (a2–d2) The surface morphologies of the coatings after immersion in Hank's solution for 10 d. (e) The obtained Sr/(Sr + Ca) ratio and (f) the (Ca + Sr)/P ratio of the coatings before and after immersion tests.



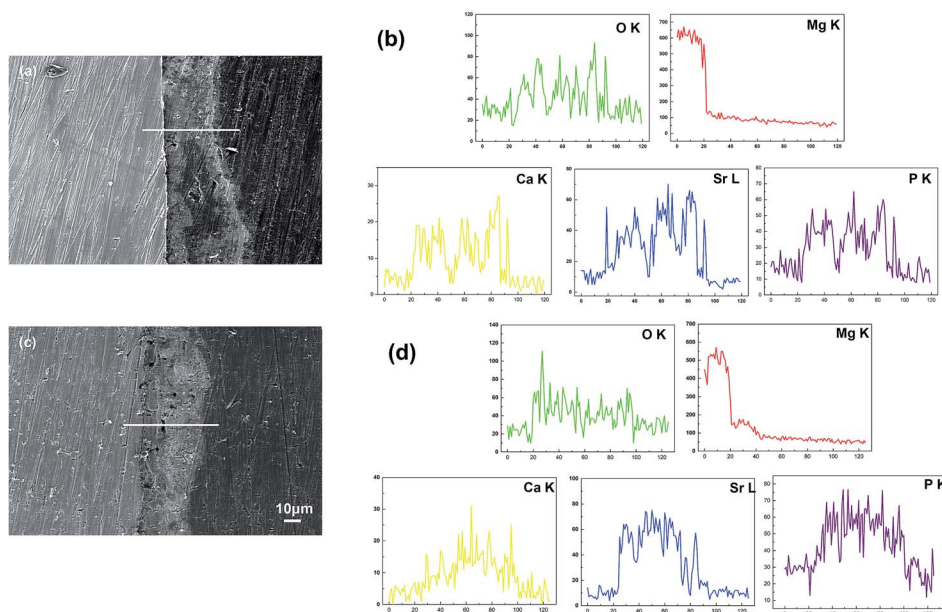


Fig. 4 The cross-sectional observations of the Sr-HA coatings with 50% Sr (a) before and (c) after immersion in Hank's solution for 10 d. The EDS line-scan of the 50% Sr-HA coating (b) before and (d) after corrosion.

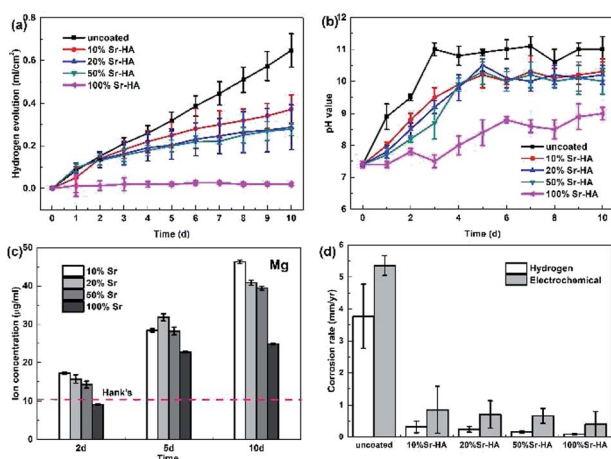


Fig. 5 (a) Hydrogen evolution, (b) pH change and (c) Mg release of the Sr-HA coated Mg alloys immersed in Hank's solution. (d) The corrosion rates calculated from the electrochemical and immersion tests.

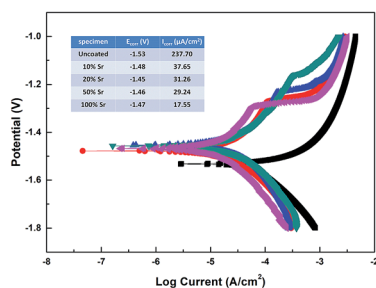


Fig. 6 Potentiodynamic polarization curves of the Sr-HA coated Mg alloys in Hank's solution.

more compact surface morphology with the decreasing values of Sr content in the coating. The flower-like crystals are still seen on the surface of 100% Sr-HA coating with some loose cotton like precipitation in the surrounding area (Fig. 3d2). The Sr content is significantly reduced to 66.7 at%. It is also found that the (Ca + Sr)/P ratio is reduced for 10–50% Sr-HA coating after immersion test, while it increases from 1.43 to 1.59 for 100% Sr-HA sample (Fig. 3f). From the cross-sectional images of the samples, some micro-pores appear in the coatings and a reduction of Sr content is observed in the outer layer in the coatings (Fig. 4c and d), indicating the dissolution of Sr-HA coating. The XRD analysis reveals no difference in the coatings before and after immersion tests (Fig. 7). We observe a reduction of the diffraction peaks from Mg substrate, indicating the coatings become more compact. This speculation is in line with the SEM observation in Fig. 3a2–d2. An interesting point to be noted is that there is no $\text{Mg}(\text{OH})_2$ peaks in the XRD diffraction spectra. It suggests that the Sr-HA coating could maintain its corrosion resistance and keep the Mg substrate nearly unattacked during the 10 days immersion test.

The Ca, Sr and P release for different immersion intervals is revealed in Fig. 8. With increasing designed Sr proportions up to 50%, the increasing concentrations of Sr and P is observed with higher Sr substitution coatings. On day 5, 50% Sr-HA coating shows the disproportionate amount of Sr releasing relative to the Sr content in coatings. The ion releasing results also demonstrate the dissolution of coating. The coatings become more soluble with the elevation of the Sr content and the results are consistent with previous observations for Sr-HA powders and SrCaP coatings.^{16,36} Ca and P concentrations decrease with longer immersion time.

For 100% Sr-HA coatings, Pan *et al.*¹⁷ reported the highest solubility of fully substituted Sr-HA powders, while our study



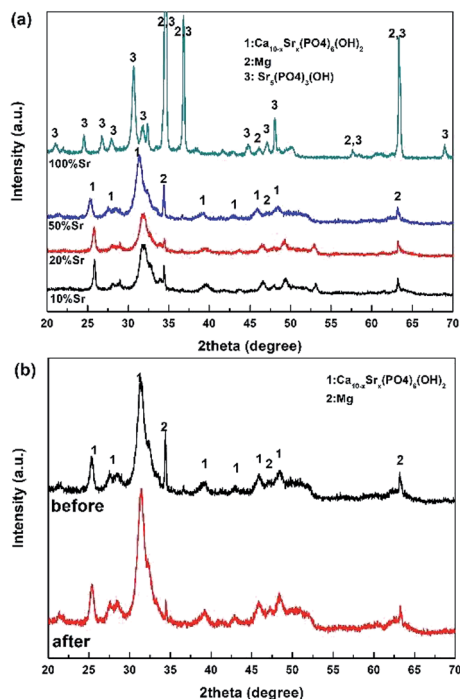


Fig. 7 (a) XRD patterns of the Sr-HA coated Mg alloys after 10 d immersion in Hank's solution. (b) The comparison of the XRD patterns for 50% Sr-HA coated Mg alloys before and after immersion test.

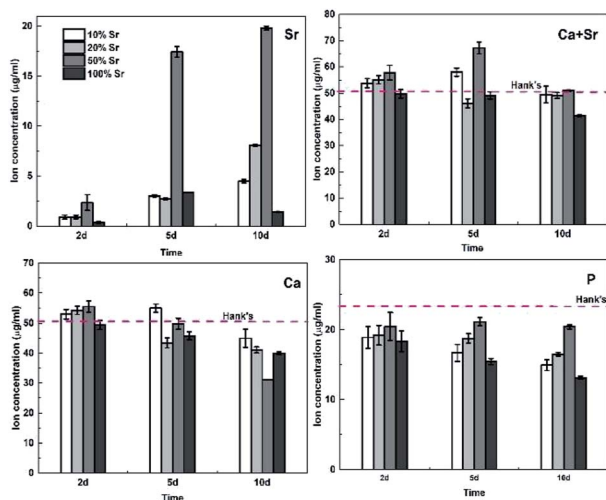


Fig. 8 The Ca, Sr and P concentrations of the Hank's solution incubating with Sr-HA coated Mg alloys for different time intervals.

indicated an opposite behavior. 100% Sr-HA coatings exhibit the least Sr releasing during the immersion tests even with higher Sr content, which demonstrate its slowest dissolution rate. On the one hand, this could be interpreted by the less contamination of carbonate in 100% Sr-HA coatings (Fig. 2), which results in lower solubility.^{35,37} On the other hand, the high crystallinity also contributes to the highest corrosion resistance of 100% Sr-HA coatings.

3.3 Degradation mechanism of Sr-HA coated Mg alloy

The degradation of the Sr-HA coatings accelerates with increasing Sr content in the coatings up to 50%. This is largely due to the larger ionic radius of Sr, which typically results in an increase in the lattice parameters (Table 2) with higher levels of Sr substitution for Ca. Surprisingly, the more soluble 50% Sr-HA coatings show much better corrosion resistance than the coatings with lower degrees of Sr substitution. It can be explained by the dissolution and reprecipitation type mechanism of Sr-HA coatings.^{38,39} The schematic diagram is shown in Fig. 9. It begins with the surface dissolution of the Sr-HA coating when in contact with the corrosive media. Ca^{2+} , Sr^{2+} and PO_4^{3-} are released into the interface between coating and solution. Simultaneously, the corrosion of Mg cannot be completely restricted, which is largely due to the penetration of the corrosive media through the micro- and nano-pores in the coatings (see Fig. 4a) to the coating-Mg interface. This results in the instant dissolution of underlying Mg substrate with the generation of hydrogen gas (Fig. 5a) and OH^- . This locally alkaline environment favors the apatite continuously precipitate back to the sample surface.²³ As a consequence, the concentrations of Ca and P in Hank's solution decrease with longer immersion period (Fig. 8c and d) with the formation of a more compact coating morphology (Fig. 3). During this process, some micro-pores could be left in the original coatings or newly formed precipitation layer due to the hydrogen evolution from Mg substrate (Fig. 4c). This suggests that rapid coating dissolution may induce increased local release of Ca^{2+} , Sr^{2+} and PO_4^{3-} , which readily leads to increased apatite formation^{7,40} to repair the dissolved coating, evidenced by the reduced concentration of Ca and P with longer immersion period (Fig. 8). The corrosion of Sr-HA coated Mg alloys is therefore suppressed. In the case of 100% Sr-HA coatings, the local release of multiple ions is less though, good apatite formation ability is observed. It is confirmed by the more rapid reduction of Ca and P concentrations in Hank's solution during the immersion period (Fig. 8), as compared to the 10–50% Sr-HA coatings. In addition, the newly formed cotton like precipitation (Fig. 3d2) with close Ca/P ratio (1.59) to HA (Fig. 3f) also demonstrates the good mineralization ability. Fredholm *et al.*⁴⁰ also indicated enhanced apatite formation for 100% Sr substituted bioactive glass.⁴⁰ The slowest degradation of 100% Sr-HA coated sample maybe attributed to both the slowly dissolution of the coating and the newly formed apatite. The degradation behavior of Sr-HA coating is a consequence of a number of different factors, while the level of Sr substitution is seen as highly significant considering its effect on the organization of HA lattice.

Table 2 The calculated unit cell parameters for Sr-HA coatings

Sample	<i>a</i> (nm)	<i>c</i> (nm)	<i>V</i> (nm ³)
10% Sr	0.9553	0.6882	0.5439
20% Sr	0.9594	0.6933	0.5526
50% Sr	0.9645	0.7056	0.5685
100% Sr	0.9747	0.7261	0.5974



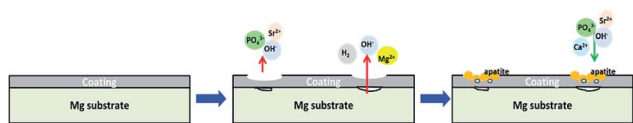


Fig. 9 Schematic diagram reveals the dissolution-precipitation of the Sr-HA coating.

According to the obtained results in this study, it is obvious that the degradation of magnesium substrate can be regulated by controlling the degree of Sr incorporation in the Sr-HA coatings.

3.4 Cell biocompatibility of Sr-HA coated Mg alloy

The effects of Sr-HA coatings on the cell proliferation were investigated by exposure of the cells to 50% diluted extracts of the experimental samples. Fig. 10 reveals the osmolality, pH value and ion concentrations of the extracts. The extracts of Sr-HA coated samples exhibit significantly lower osmolality, pH value and Mg concentration than that of the uncoated samples. It shows no significantly difference in the three values of different Sr-HA coated samples. While the Sr concentration in the extracts increases with increasing Sr content in the coatings up to 50%. The Ca and P levels for all the samples are significantly lower than the DMEM control. Fig. 11 shows the metabolic activity tested by CCK-8 assay and the ALP activity in different extracts. The cells show increased proliferation in the Sr-HA extracts. It reveals that cells cultured in four Sr-HA coating extracts exhibit relatively similar OD value until day 3. The difference between the four groups become distinct at day 5 ($p < 0.05$) and OD value of the 50% Sr-HA coating extracts is significantly higher than the other three groups. The images of osteoblasts cultured in the extracts are shown in Fig. 11c to be consistent with the CCK-8 results. The cell ALP activity normalized to total protein show no significantly difference between four Sr-HA groups at day 3 and 5. Both the OD value and ALP activity of Sr-HA groups are higher than those of the uncoated one.

Osteogenic gene expression was also studied on Sr-HA coated samples to further analyse the capability of the coatings for enhanced differentiation (Fig. 12). The gene expression is normalized to that of cells cultured in the extracts of uncoated sample. Three osteogenesis markers, ALP (an early marker for osteogenic differentiation), RUNX2 (a key transcript factor for bone formation) and Col-1 (main content of bone ECM), were

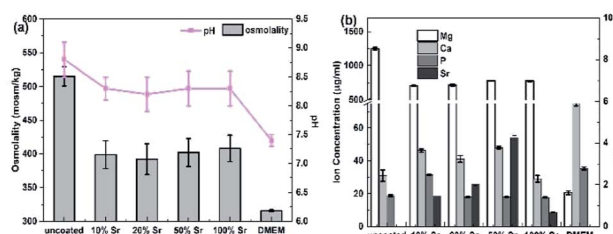


Fig. 10 (a) The osmolality, pH value and (b) ion concentrations of the extracts for the Sr-HA coated Mg alloys.

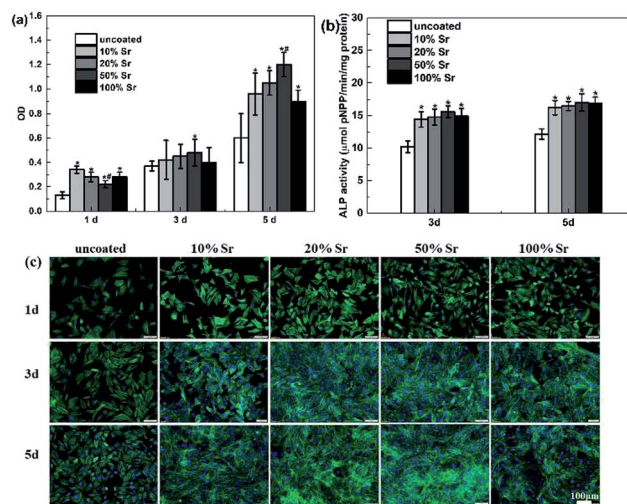


Fig. 11 Response of MC3T3-E1 osteoblasts to the extracts for Sr-HA coated Mg alloys, (a) CCK8 assay and (b) ALP activity. * $p < 0.05$ vs. uncoated, # $p < 0.05$ vs. 10% Sr-HA. (c) Fluorescent images of MC3T3-E1 osteoblasts cultured in the sample extracts.

tested. Higher ALP and Col-1 expression was observed for 20% Sr-HA coatings. A comparable RUNX2 expression was measured for 10–50% Sr-HA coatings. In addition, a distinct lower expression of Col-1, Runx2 and ALP was observed for 100% Sr-HA coatings ($p < 0.05$).

The positive results of the Sr-HA coated samples can be explained by the following reasons. First, it is attributed to the slower corrosion and the resulted lower concentration of Mg in the extracts (Fig. 10). Wang *et al.*⁴¹ reported the combined effects of the loss of LTRPC7 function and the decrease of Ca involved binding sites/reactions in high Mg level, which may influence the cell viability.⁴² 10–15 mM of Mg ion concentration is reported to be the critical dose without inhibiting cell viability. After 50% dilution, the concentration of Mg is approximately 15 mM for Sr-HA coating extracts which is in the range of the biosafety level recommended in the previous study,⁴¹ while 26 mM of Mg is observed for the uncoated samples. Second, the influence of Sr released from Sr-HA coatings cannot be excluded. The present study indicates that the osteoblast proliferation increases when treated with extracts containing higher Sr concentration and after longer incubation period. These results suggest a specific correlation between the Sr concentration in the extracts (Fig. 10b) and osteoblasts replication as well as osteogenic differentiation (Fig. 11a and

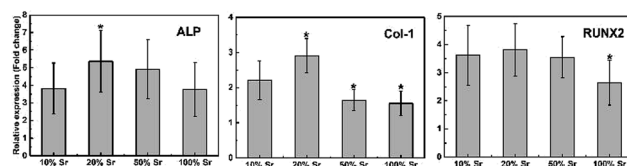


Fig. 12 Relative expression of Col-1, ALP and RUNX2 by MC3T3-E1 osteoblasts cultured in the extracts for Sr-HA coated Mg alloys for 5 d. * $p < 0.05$ vs. 10% Sr-HA.



12). It is in line with the previously report that Sr is beneficial to osteoblast proliferation and osteogenesis in a dose dependent manner.^{22,40,43} Chen *et al.*²³ also found that SrP coated Mg exhibited better hMSC proliferation when extracts contained a higher concentration of Sr ions were used. From the preliminary data, it can be concluded that Sr content is an important parameter to adjust the solubility of Sr-HA coating and consequently regulate the degradation and biocompatibility of the Sr-HA coated Mg alloys. Future work will aim to assess the biocompatible mechanism of the Sr-HA coated Mg alloy using *in vivo* animal models.

4. Conclusions

In the current study, we have developed a series of Sr-HA coatings with various Sr contents on the surface of Mg alloy substrates. The results demonstrated that the degradation performance of the Sr-HA coated Mg alloys was significantly affected by the Sr content in the coatings. The degradation was reduced with increasing degree of Sr substitution in the Sr-HA coatings. 100% Sr-HA coated Mg alloys exhibited the slowest degradation performance as compared to the other samples. The MC3T3-E1 osteoblasts cultured in the extracts of Sr-HA coated Mg alloys were evaluated for their proliferation and differentiation responses. It revealed that improved cell proliferation and differentiation responses were observed due to the delayed Mg dissolution and Sr release from Sr-HA coated Mg alloys. The present study suggests that Sr_x-HA coating is a promising and effective option to control the degradation of Mg substrates, and the variation of Sr availability through dissolution influence the cell responses.

Conflicts of interest

There are no conflicts to declare.

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