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A promising orthopedic implant material with enhanced osteogenic and antibacterial activity: Al2O3-coated aluminum alloy

Yuchang Weng,* Hanxia Liu,† Shunping Ji, Qian Huang, Hao Wu, Zhibin Li,* Zhongzhen Wu,* Huaiyu Wang, Liping Tong, Ricky K.Y. Fu, Paul K. Chu,*, Feng Pan

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1. Introduction

Osteoporosis, a senile disease, is becoming more prevalent as baby boomers age [1], thus, artificial implants are often needed. Most orthopedic implants are made of titanium (Ti) alloys with high strength, good corrosion resistance, and moderate biocompatibility [2–4]. Nevertheless, the elastic modulus of Ti-based implants is larger than that of natural bones which will cause potential resorption of tissues adjacent to the implants due to the stress shielding effect. Although biodegradable magnesium (Mg) alloys have merits such as the light weight, good biodegradability, and similar elastic modulus compared to human bone which make them promising for orthopedic applications [5–7], Mg-based implants may not have the proper corrosion resistance and mechanical properties under weight-bearing conditions.

As a common-used light alloy, aluminum (Al) alloys are similar to Mg alloys in terms of weight and elastic modulus and have comparable yield strength as Ti alloys, which makes them potential candidates of orthopedic implant materials. In addition, aluminum alloys also possess favorable properties such as good casting ability and low price, which can remarkably reduce the economic cost of manufacturing and using of implant materials. Nonetheless, potentially toxic Al ions can be released and cause biocompatibility problems. Besides, owing to their relatively poor anti-corrosion characteristics and large intrinsic friction coefficient, Al alloys can easily be worn and eroded during using. Hence, Al alloys have barely been used as orthopedic implants. However, if appropriately protective surface modification can be implemented on Al alloys, it is possible to carry forward their good mechanical properties and to mitigate, even to eliminate hazardous effects of Al corrosion.

With good wear resistance, intrinsic wettability, chemical stability, and biocompatibility, alumina ceramics have been widely used as orthopedic implant materials, especially as dental and bone replacement [8–10]. However, like most of the ceramic materials, alumina ceramics have intrinsically poor elasticity and bear a high risk of fracture [11]. In recent studies, alumina ceramics have been used as coatings in conjunction with metallic substrates for biomedical use [12–15]. The ceramic provides the hardness and wear resistance while the metallic component provides toughness and high strength for load bearing applications [16]. Therefore, to combine the advantages of aluminum and alumina, we designed and fabricated an Al2O3-coated aluminum alloy

Keywords: Al2O3 coating, Al alloy, Micro-arc oxidation, Element doping, Biological properties

Better orthopedic implants improve the life quality of patients and elderly people. Compared to Ti and Mg alloys commonly found in bone implants, Al alloys have merits such as high specific strength, excellent casting capability, and low cost, but unfortunately, the poor wear resistance, corrosion resistance, and insufficient bio-compatibility have hampered wider application to bone replacements. In this study, well structure designed and excellent adhered Al2O3 coating is fabricated by micro-arc oxidation (MAO) on Al alloys to improve the wear resistance and corrosion resistance and inhibit release of potentially harmful Al ions from the Al alloy substrate. Ca, Fe, and Zn are respectively doped into the Al2O3 coating to enhance the biological properties. In particular, the Al2O3 coating doped with Zn delivers outstanding osteogenic performance and the antibacterial rates against E. coli and S. aureus are 99.5 ± 0.56% and 98.77 ± 0.52%, respectively. The improved mechanical and biological properties reveals that the Al2O3 coated Al alloys prepared by MAO coatings have large potential in orthopedic applications.
with a ceramic-on-metal structure. By using micro-arc oxidation (MAO) [17–19], a thick and double-layered structured Al2O3 coating can be grown in-situ on aluminum alloy [20]. The dense inner layer of the Al2O3 coating with high bonding strength to the substrate can protect the underlying Al alloys from the external environment and prevent the exfoliation of toxic Al ions. In addition, by mimicking the microstructure of natural bones, the porous outer layer of the Al2O3 coating can facilitate bone growth in vitro and in vivo. Moreover, in this work, specific biologically important elements, such as Zn, Ca, and Fe, are incorporated into the Al2O3 coatings in situ to obtain the favorable biological functions by MAO in special modified electrolyte. The samples with and without the Al2O3 coatings are systematically characterized and in vitro experiments are performed to investigate the osteogenic and antibacterial capabilities.

2. Materials and methods

2.1. Sample preparation

MAO was used to produce coatings with different doping elements on the LY12 Al alloy. Before MAO, the substrate was polished with SiC abrasive paper, degreased, and rinsed with acetone and distilled water. MAO was conducted in the corresponding electrolytes composed of the primary salt of sodium hexametaphosphate ((NaPO3)6, 15–20 g/L, XiLong Science Co., Ltd) and other additives at a constant current density (5 A/dm2) at 500 Hz for 8 min with a duty cycle of 12% and 20 kW AC power supply (Chengdu JINCHUANGLI Technology Co., Ltd, JCL-WH20). In order to incorporate Ca, Fe and Zn, calcium acetate hydrate (C4H6CaO4·H2O, 3 g/L, XiLong Science Co., Ltd), ammonium iron (III) oxalate hydrate (C2H8FeN2O12-2H2O, 6 g/L, XiLong Science Co., Ltd), or zinc acetate hydrate (C2H8O2Zn·2H2O, 5 g/L, XiLong Science Co., Ltd) were introduced to the electrolyte, respectively. The undoped samples served as the control for comparison. During MAO, the temperature of the electrolyte was controlled to be below 40 °C with an external water cooling system. After the surface treatment, the samples were cut to the required dimensions by a simple cutting machine, rinsed with distilled water, and dried.

2.2. Sample characterisation

The structure of the samples was analyzed by X-ray diffraction (XRD, Bruker, D8 Advance) equipped with an X-ray source of Cu target (λ = 0.15418 nm). The data were collected at 2θ = 10–80° at a scanning rate of 2θ = 5°/min. The samples were analyzed by X-ray photoelectron spectroscopy (XPS, ESCALAB 250X, Thermo Fisher) referenced to the C1s peak (284.8 eV). The energy of Al Kα irradiation at the sample surface is 1486.6 eV. The surface morphology was examined by field-emission scanning electron microscopy (FE-SEM, Carl Zeiss, SUPRA® 55) and the elemental composition and lateral distribution were determined by energy-dispersive X-ray spectroscopy (EDS). To assess the corrosion behavior, polarization curves were acquired with an electrochemical workstation (1470E, Solartron Metrology) between -2.0 and 0 V versus the saturated calomel electrode (SCE) at a scanning rate of 10 mV/s at 25 °C in the NaCl (3.5 wt%) and simulated body fluid (SBF) solutions, respectively. The corrosion potential (Ecorr) and corrosion current density (icorr) were analyzed by the Tafel extrapolation method. Scratch tests were measured on a scratch tester (WS-2005, Lanzhou Institute of Chemical Physics, Chinese Academy of Sciences). In the test, the load on a diamond indenter with an included angle of 120° was linearly increased from 0 to 100 N at a rate of 20 N/min at 25 °C and 65 ± 1% RH. The wear resistance was evaluated on a friction-wear tester (MFT-5000, Rtec Instrument) at 25 °C and 65 ± 1% RH. GCr15 steel balls with diameter of 4 mm, surface roughness of 0.02 μm and hardness of 64 HRC were used as the counterpart materials. All wear tests were conducted with GCr15 steel balls that slide against the middle of MAO coatings in a circular motion with a radius of 5 mm. The rotating speed and normal load of the wear tester were 200 r/min and 3 N, respectively. The depth profiles and topographies of the wear track were detected using a VK-X200 laser confocal microscope. The surface contact angles were measured on contact angle meter (PT-705B, Dong Guan Precise Test Equipment Co., LTD) at 25 °C and 55 ± 1% RH. For statistical accountability, more than 6 measurements were performed on each sample and the average values were calculated. The thickness of the MAO coatings was determined on an eddy current thickness meter (CTY2300, SDCH. Co.; LTD) and multiple measurements were conducted to obtain averages. The simulated body fluid (SBF) was used as the culture medium in the immersion experiments at 37 °C and inductively-coupled plasma atomic emission spectrometry (ICP-AES, JY2000-2, Horiba) was employed to monitor the time-dependent dissolution of Al ions from the samples. The volume of SBF was calculated by the following equation [21]: \[ V_i = \frac{S_i}{10}, \] where \( V_i \) is the volume of SBF (mL) and \( S_i \) is the apparent surface area of each sample (mm²).

2.3. Cell culture

The human bone mesenchymal stem cells (hBMSCs) were obtained from ATCC and maintained in the culture medium composed of the α-minimum essential medium (α-MEM, Hyclone), 10% fetal bovine serum (FBS, Corning), and 1% penicillin/streptomycin (Invitrogen). After expanding to passage 5, the hBMSCs were seeded onto the samples with different dimensions and cultured. Before cell seeding, the samples were sterilized with 75% ethanol overnight and rinsed with the sterile phosphate-buffered saline (PBS) solution. The cultures were incubated in a humidified atmosphere of 5% CO2 at 37 °C and the culture medium was refreshed every 3 days.

2.3.1. Cell viability

The hBMSCs were seeded onto 10 mm × 10 mm samples (1 × 10⁴ cells per sample) on 24-well tissue culture plates. After culturing for 1, 3, and 7 days, the cells were quantitatively investigated with the cell viability assay using the CCK-8 kit (Donjindo). At each time point, the harvested specimens were rinsed thrice with PBS and incubated with the 10% CCK-8 reagent in the culture medium (0.7 ml) for 4 h. Afterwards, 100 μl of the incubated mixture was transferred to a 96-well plate and analyzed spectrophotometrically at 450 nm (Multiskan GO, Thermo Fisher).

2.3.2. Live/dead staining

Live/Dead Cell staining was performed on the 10 mm × 10 mm samples on 24-well plates were seeded with the hBMSCs at a density of 1 × 10⁴ cells per sample. After culturing for 3 days, the cells were rinsed thrice with PBS, stained for 15 min in darkness with the Live/Dead Cell Staining Kit (BioVision), and examined by fluorescence microscopy (Olympus BX53).

2.3.3. Flow cytometry

Apoptosis and necrosis of the hBMSCs on the samples were evaluated with the Annexin V-FITC/PI assay (TransStart) according to the manufacturer’s instructions. Briefly, the hBMSCs were cultured for 3 days, harvested, rinsed with PBS, and stained with Annexin V-FITC and PI. The apoptotic cells labeled with Annexin V-FITC and necrotic cells labeled with PI were analyzed on a cell Lab Quanta SC flow cytometry instrument (Beckman couler).

2.3.4. Cell morphology

After incubation for 3 days, the cells on the samples were rinsed thrice with PBS and fixed with 2.5% glutaraldehyde for 15 min. Prior to FE-SEM (Carl Zeiss, SUPRA® 55), the specimens were sequentially dehydrated in gradient ethanol solutions (30, 50, 75, 90, 95 and 100 v/v %) for 15 min and final dehydration was conducted in hexamethyldisilazane. Afterwards, the samples were dried, sputter-coated with...
platinum, and examined by FE-SEM.

2.3.5. Quantitative real-time PCR

The osteogenesis-related gene expressions of the hBMSCs on the samples were quantitatively analyzed by the real-time polymerase chain reaction (real-time PCR). Samples with a dimension of 20 mm × 20 mm were seeded with hBMSCs at a density of 5 × 10^4 cells per sample on 6-well plates. The cells were cultured in a basic growth medium initially for 3 days and then the osteogenic medium (basic medium with 5 μmol/ml glycerophosphate, 50 μg/ml ascorbic acid, and 100 pmol/ml dexamethasone) was refreshed for subsequent incubation. After osteogenic induction for 21 days, the cells were rinsed thrice with PBS, fixed with 95% alcohol for 10 min, and rinsed with distilled water three times. The specimens were then stained with 40 mM Alizarin Red (pH 4.2, Sigma), dissolved in 10% cetyl pyridinium chloride (Sigma), and analyzed for extracellular matrix (ECM) mineralization spectrophotometrically at 540 nm (Multiskan GO, Thermo Fisher).

2.3.6. Mineralization assay

Mineralization of the hBMSCs was assessed by Alizarin Red staining. Samples with dimensions of 20 mm × 20 mm were seeded with hBMSCs at a density of 5 × 10^4 cells per sample on 6-well plates. The cells were cultured in a basic growth medium initially for 3 days and then the osteogenic medium (basic medium with 5 μmol/ml glycerophosphate, 50 μg/ml ascorbic acid, and 100 pmol/ml dexamethasone) was refreshed for subsequent incubation. After osteogenic induction for 3, 7, and 14 days, the total RNA of the cultured cells was extracted using the Trizol reagent (Life Technologies) and 2 μg of total RNA from each specimen was reversely transcribed into cDNA by using a RevertAid First Strand cDNA Synthesis Kit (Thermo Scientific) according to the manufacturer’s instructions. RT-PCR was performed on a real-time PCR system (BIO-RAD) to analyze the gene expressions of alkaline phosphatase (ALP), bone sialoprotein (BSP), and osteocalcin (OCN) with β-actin serving as the endogenous gene for normalization. The primer sequences are listed in Table 1 and quantification of the gene expressions was based on the comparative cycle-threshold (Ct) method expressed as 2^−ΔΔCt ± SD [22].

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward primers</th>
<th>Reverse primers</th>
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<tbody>
<tr>
<td>ALP</td>
<td>CCCAAAGGCTTCTTCTTG</td>
<td>CTGGTAGTTGTTGAGC</td>
</tr>
<tr>
<td>BSP</td>
<td>CTGGAAAGGTCGTCTTGGG</td>
<td>TTGAGCGCGTGTATTGAC</td>
</tr>
<tr>
<td>OCN</td>
<td>GCCCGTGTGCAAGGC</td>
<td>GGACCCACATCCATAG</td>
</tr>
<tr>
<td>β-actin</td>
<td>TGGCACCACACACAAAGAA</td>
<td>CTAAGTCATAGTCCGCCTAG</td>
</tr>
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Fig. 1. SEM images and EDS spectra of the MAO samples: (A) Undoped, (B) Ca-doped, (C) Fe-doped, and (D) Zn-doped. For each sample, the typical water droplet image is shown on the top-right.
where \( \bar{C} \) was the average number of bacteria on pristine Al alloy substrate (CFU/sample) and \( \bar{T} \) was the average number of bacteria on the various MAO samples (CFU/sample).

2.5. Statistical analysis

The in vitro assays were performed in triplicate and each value was expressed as mean ± standard deviation. Each in vitro experiment was repeated three times with the typical data shown. Statistical evaluation was performed by one-way ANOVA, where \( p < 0.05 \) was considered significant and \( p < 0.01 \) was considered highly significant.

3. Results

3.1. Surface characterizations

Fig. 1 shows the surface morphology of the samples and volcano-like porous structures are observed. The pore size varies from several nanometers to more than ten micrometers. In particular, the surface of the Fe-doped sample has the least amount of pores and smallest average pore size but the undoped and Zn-doped samples show more pores. According to EDS results, the coatings are mainly composed of Al, P, and O. Besides, Ca, Fe, and Zn are also detected from the corresponding doped coatings, respectively. The surface wettability is indicated by the typical water droplet images shown on the top-right in Fig. 1. The Ca, Zn, and Fe-doped samples show a similar water contact angle of about 35° whereas the water contact angle of the undoped sample is 55°. The \( \text{Al}_2\text{O}_3 \) coatings formed on the Al alloy are mainly composed of Al, P, and O in the phosphate electrolyte [23] in addition to the respective dopant element (Ca, Fe, or Zn).

Generally, the micro-arc oxidation coatings have the double-layer structure [20,24]. Fig. 2 shows both outer and inner layers in the Zn-doped sample. As shown in Fig. 2A, the loose and volcano-like porous structure exists in the outer MAO layer, which is formed by the liquid plasma containing molten Al and alumina during the breakdown of the weakest location repeatedly under the electric field [25,26]. Compared to the porous outer layer, the inner layer (Fig. 2B) in the coating is much more compact with few pores. The inner layer in the Zn-doped sample has a thickness of about 3 μm and is abundant with Al, O, and P (Figs. 2C, D, and S2 in Supporting information), whereas more Zn is detected from the outer layer. The results of other \( \text{Al}_2\text{O}_3 \) samples are presented in Figs. S3–S5 and the similar results are observed. All the \( \text{Al}_2\text{O}_3 \) coatings have similar thicknesses with an outer layer thickness of 10–15 μm and an inner layer thickness of 3–5 μm (Fig. S6). Contrary to the loose structure of the outer layer, the compact structure of inner MAO layer, which enhances the anti-corrosion performance and inhibits release of Al ions from the Al alloy substrate, is formed in the last stage of the MAO process [27].

Supplementary data associated with this article can be found, in the online version, at https://doi.org/10.1016/j.apsusc.2018.06.233.

Fig. 3A shows the crystal phases determined by XRD and the pristine Al alloy substrate serves as the control. The \( \text{Al}_2\text{O}_3 \) samples show typical
peaks of $\gamma$-$\text{Al}_2\text{O}_3$ and $\alpha$-$\text{Al}_2\text{O}_3$ arising from the Al alloy substrate. Since the instantaneous temperature in a small arc discharge region is over 2000 °C, the thermodynamically stable $\alpha$-$\text{Al}_2\text{O}_3$ phase can formed in the inner layer [26]. In contrast, the $\gamma$-$\text{Al}_2\text{O}_3$ phase mostly presents in the outer layer due to fast cooling by the electrolyte[28]. Besides $\text{Al}_2\text{O}_3$, no other diffraction peaks of the dopants can be observed, which maybe results from the small concentration and amorphous state. However, the XPS results shown in Fig. 3B and D reveals the presence of Ca, Fe, and Zn with the concentrations of 4.5%, 3.0%, and 1.0%, respectively (Fig. 3C), corresponding to metal oxides or metal phosphates, which indicates the presence of iron, calcium, zinc oxide or phosphate in the films.

3.2. Adhesion, corrosion and wear resistance performance

Acoustic emission produced in the scratch test is utilized to evaluate the interface bonding strength of the $\text{Al}_2\text{O}_3$ coatings. Fig. 4A shows that the adhesion strength of all the $\text{Al}_2\text{O}_3$ coatings is above 30 N and follows the order of Fe-doping (62.0 N) > Zn-doping (46.3 N) > Ca-doping (46.0 N) > No doping (38.2 N), indicating that the adhesion strength of the $\text{Al}_2\text{O}_3$ coating can be improved by doping. As shown in the Fig. S7, the discharge of the Fe-doped sample is more intense compared to those of Ca-doped and Zn-doped samples, which induces the much stronger arc stirring at the interface between the coating and Al substrate. Therefore, the Fe-doped sample exhibits the largest adhesion. The improved adhesion between the MAO coating and substrate mitigates the risk of aseptic loosening caused by delamination of the coating[29].

Fig. 4B shows the dynamicpotential polarization curves of pristine Al alloy substrate and $\text{Al}_2\text{O}_3$ samples measured in 3.5 wt% NaCl solution. The $i_{\text{corr}}$, $E_{\text{corr}}$, and $\beta$ values are calculated by Tafel extrapolation from the linear cathodic polarization region and the results are shown in Table 2. The pristine Al alloy exhibits the worst anti-corrosion behavior with the smallest corrosion potential and largest corrosion current density. After MAO treatment, $E_{\text{corr}}$ increases and $i_{\text{corr}}$ decreases obviously, demonstrating a significant enhancement in the corrosion resistance. Similar results are also obtained from the polarization test in the SBF solution (Fig. S8 and Table S1). It has been reported that the dense inner layer can prevent penetration of the solution to the underlying substrate leading to well corrosion resistance[30]. Besides, the corrosion resistance depends on the structure of the MAO coatings[31]. On account of the least pores and the smallest average pore size, the Fe-doped sample shows the best anti-corrosion performance.

Figs. 4C, D, and S9 show the friction and wear behavior of different samples. Compared to the Al alloy substrate with the friction coefficient of about 1.0 and the wear depth of about 15 μm only after 77 turns, the wear resistances of all the $\text{Al}_2\text{O}_3$-coated samples increase. Especially, the Zn-doped sample possesses the smallest friction coefficient of about 0.5 and which further decreases to about 0.3 with increasing wear time. No apparent wear scratch can be observed on all the $\text{Al}_2\text{O}_3$-coated samples after 12,000 turns, while Ti6Al4V shows a deep wear scar more than 30 μm after 12,000 turns due to its lower hardness than GCr15 steel ball, although its friction coefficient is as good as the $\text{Al}_2\text{O}_3$-coated samples (Fig. 4D). The $\text{Al}_2\text{O}_3$ coatings can improve the wear resistance of the Al alloy thereby prolonging the service lifetime and lowering inflammation risks caused by released particles[32].

3.3. In-vitro biocompatibility

Al leaching tests are conducted to evaluate the protection effects of the $\text{Al}_2\text{O}_3$ coatings. As shown in Fig. 5A, leaching of $\text{Al}^{3+}$ ions from the pristine Al alloy substrate increases from 0 mg/L to over 80 mg/L after immersion for 28 days. However, after the same immersion time, the amount of the Al ions releasing from the $\text{Al}_2\text{O}_3$-coated samples is below 0.2 mg/L, even lower than that dissolves from Ti6Al4V, which can be ignored considering the test error. Besides, according to the World Health Organization, daily ingestion of aluminum of 0–0.6 mg/kg is biosafe to human being. In this work, the dosage of Al ions leaching out...
has been greatly reduced to a concentration below the toxicity criterion by MAO treatment. Fig. S10 shows that volcano-like porous structures are observed on the coating surfaces and least and smallest pores occur on the Fe doped sample. The coatings are mainly composed of Al, P, and O and Ca, Fe, and Zn are detected from the corresponding samples. All the results show that no significant change can be found on the surface morphology and chemical compositions after immersion in SBF for 28 days.

For further evaluate the cell cytotoxicity of different samples, hBMSCs are seeded and the CCK-8 assay is employed to measure the time-dependent cell viability after incubation for 1, 3, and 7 days. As shown in Fig. 5B, the Fe-doped sample shows the most positive proliferation of hBMSCs and the other Al2O3-coated samples show similar cell viability with the Al alloy substrate and Ti6Al4V in the first day. When the incubation time increases to 3 and 7 days, the difference between the Fe-doped sample and other samples becomes statistically significant. The Fluorescence microscopy and FE-SEM results are presented in Figs. 5C and S11, and the viable and dead hBMSCs are stained green and red, respectively. All the samples show good cell viability and the cells spread well. Flow cytometry is further performed and reveals that the samples have good cytocompatibility showing low cell apoptosis and necrosis (insets in Fig. 5C). Among them, the Fe-doped sample shows the best performance of cell viability, which can be probably due to the enhanced metabolic activity of cells at an appropriate iron concentration [33].

3.4. In-vitro osteogenesis ability

Osteogenesis of cultured hBMSCs is another crucial factor to the success of bone implants. After culturing for 3, 7, and 14 days, osteogenic induction of the hBMSCs is evaluated by real-time PCR utilizing the primers for bone markers as ALP [34], BSP [35], and OCN [36]. As shown in Fig. 6A-C, the Zn-doped sample is superior exhibiting up-regulation of all the three osteogenic genes. After 14 days, the ALP, BSP, and OCN gene levels observed from the Zn-doped samples are about 2.9, 3.4 and 2.0 times of those of the Al alloy substrate and 2.4, 2.8 and 1.6 times of those of Ti6Al4V, respectively. After 21 days, the hBMSCs are examined by Alizarin Red staining and spectrophotometry. Fig. 6D indicates that ECM mineralization of hBMSCs on the Zn-doped sample is the highest. Both gene determination and mineralization assay verify that Zn-doped sample is more desirable from the perspective of osteogenesis. Besides, the other Al2O3-coated samples also show varying degrees of improving osteogenesis properties compared with the uncoated Al alloy and Ti6Al4V materials, demonstrating the important
and distinguished effect of Al₂O₃ coatings on osteogenesis.

3.5. In-vitro antibacterial ability

In this work, both S. aureus and E. coli are used to assess the antibacterial ability with bacteria counting and the results are shown in Fig. 7. Compared to the Al alloy substrate and Ti6Al4V, the Al₂O₃-coated samples show less re-cultivated bacterial colonies after 24 h for both E. coli and S. aureus. In particular, the Zn-doped and Fe-doped samples show significant antibacterial efficiency comparing to other samples. For the Zn-doped sample, barely visible bacterial colony can be found, indicating excellent antibacterial property. The antibacterial rates of Zn-doped sample are 99.5 ± 0.56% and 98.77 ± 0.52% against Gram-negative E. coli and Gram-positive S. aureus, respectively, indicating the potential anti-inflammation properties when the Zn-doped sample is implanted into human body.

4. Discussion

A new generation of bio-functional orthopedic implants possessing both osteoinductivity and antibacterial ability and bone modulus by simple and low-costly methods is demanded in clinical applications [37,38]. As illustrated in Fig. 8, after MAO treatment, a double-layer Al₂O₃ coating is formed on the Al alloy (Fig. 2). On the one hand, the compact inner layer enhances the mechanical and anti-corrosion performance (Fig. 4) and inhibits the release of potentially harmful Al ions from the Al alloy substrate (Fig. 5). On the other hand, the outer layer has a porous rough surface in the micrometer range similar to the micro-structure of natural bones, which generally can facilitate the osteogenic differentiation of osteoblast [39–41]. Therefore, comparing to the untreated Al substrate and the Ti6Al4V sample, not only the doped Al₂O₃ samples, but also the undoped Al₂O₃ sample show better osteogenesis activity (Fig. 6), suggesting the improvement of the osteogenesis is induced by the MAO treatment.

Moreover, Zn, Fe and other elements can be easily doped into the porous outer layer, which also play important roles in the osteoblast proliferation and differentiation in addition to antibacterial processes. As shown in Figs. 6 and 7, the Zn-doped sample shows the best osteogenic and antibacterial performance. The excellent osteogenic activity and antibacterial ability of Zn-doped sample can be mainly due to the released Zn²⁺ ions. It is believed that Zn ions at an appropriate concentration can promote bone formation and have antibacterial effect on bacteria without introducing undesired side effect [42,43]. Zn²⁺ can be transported to the bacteria cytosol via the ion channels at the expense of energy consumption [44]. The living condition of the bacteria deteriorate as the bacteria transport more Zn²⁺ ions and a high concentration of Zn²⁺ in the cytosol can be detrimental to bacterial cells [34]. Moreover, appropriate amounts of Zn ions released from the Zn-doped sample can be absorbed by cells, thus facilitate the expression of osteogenic-related genes and stimulate bone growth [45,46]. In our work, the incorporated Zn element in the MAO-coating can diffuse into the medium matrix and cause biological effects on the adhered cells and
bacteria. Summarily, the good cell compatibility, osteogenesis performance and high antibacterial of the Al2O3-coated samples are attributed to the synergistic effect of the double-layer structure of the MAO coatings and the doping elements. As a result, the Zn-doped Al2O3 coating delivers the outstanding osteogenesis and antibacterial performance, which make it a promising candidate as bone implants.

5. Conclusion

In this work, Al2O3 coatings with good adhesion strength and double-layer structure is fabricated to improve the anti-abrasion and anti-corrosion performance and to avoid the leaching out of toxic Al ions. Biological assessment demonstrates that the Fe-doped Al2O3 coating significantly improves the cytocompatibility of Al alloy samples. Moreover, the doped metallic elements show biomedical functions.
on osteogenesis and anti-bacteria. In particular, the Zn-doped sample shows outstanding osteogenic activity and antibacterial effects. The Al2O3-coated aluminum alloys have great potential in the development of novel kinds of bone substitute biomaterials in orthopedics.

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References

[33] S. Zhu, N. Huang, L. Xu, Y. Zhang, H. Liu, H. Sun, Y. Leng, Biocompatibility of pure


