



Cite this: *RSC Adv.*, 2017, 7, 24607

Strontium ranelate-loaded PLGA porous microspheres enhancing the osteogenesis of MC3T3-E1 cells

Zhenyang Mao,^{†a} Zhiwei Fang,^{†b} Yunqi Yang,^b Xuan Chen,^b Yugang Wang,^a Jian Kang,^b Xinhua Qu,^{ID} *^a Weien Yuan^{*b} and Kerong Dai^{*a}

Biodegradable poly lactic-co-glycolic acid (PLGA) has been used as a tissue engineering scaffold as well as a carrier for the delivery of proteins, drugs, and other macromolecules. Hydroxyapatite (HA) nanoparticle self-assembly on the surface of a PLGA microsphere can create a hydrophilic environment, and its hydroxide can neutralize the acidity of PLGA degradation products. These microspheres can be used to deliver strontium ranelate (SR), which is used to treat osteoporosis. In the present study, we fabricated porous PLGA microspheres (PM), SR-loaded (SR-)PM, and SR-PM with HA nanosuspension and polyvinyl alcohol (PVA) as a surfactant (SR-PM-HA) in a S/O/Ns method. The microspheres exhibited an interconnected porous structure. The percent cumulative release of SR from SR-PM was 90% in 22 days as compared to 85% from SR-PM-HA. In both cases, drug release was gradual (with no burst release). SR-PM and SR-PM-HA similarly stimulated the proliferation of MC3T3-E1 cells to a greater degree than PM and induced osteogenic differentiation, as determined by alkaline phosphatase staining and real-time PCR analysis of osteogenic marker gene expression. These results indicate that SR-PM-HA is biocompatible and suitable for drug delivery and osteoinduction.

Received 4th February 2017
 Accepted 1st May 2017

DOI: 10.1039/c7ra01445g

rsc.li/rsc-advances

1 Introduction

Biodegradable polymers have potential applications in medicine for drug delivery and as biomaterials for implants. One example is poly lactic-co-glycolic acid (PLGA), which has been used as a tissue engineering scaffold^{1,2} as well as a carrier for the delivery of proteins, drugs, and other macromolecules.³⁻⁶ PLGA is strong and biocompatible, making it potentially suitable for bone repair. However, it also has limitations such as poor hydrophilicity that could impact cell attachment, proliferation, and differentiation as well as drug release. Moreover, most PLGA degradation products are acidic and may elicit an inflammatory response and inhibit osteoblast activity in bone repair.⁷⁻⁹

Nanostructured materials and porous spheres have potential applications for drug delivery, healthcare, and biomedical and pharmaceutical industries.¹⁰⁻²⁰ Many physical and chemical methods have been developed to synthesize nanoparticles or porous spheres.^{10,16} Compounds of various sizes have been used

as nanostructured materials and porous spheres for drug delivery and bioregulation in diverse systems for a specific purpose or disease.^{10,16} Liposomes, polymer micelles/microspheres, dendrimers, nanocapsules, and metal nanoparticles have been used as vehicles for drug delivery and bioregulation.²¹⁻²³

Nano- or micro-scale hydroxyapatite (HA) particles measure in the order of hundreds of nanometers in length and have been widely used in bone reconstruction. In contrast to plate-shaped bone apatite, HA particles are rod-shaped and appear as enamel. Nano-sized apatite particles of 20–40 nm can serve as building blocks of bone by self-assembly in a collagen matrix to produce a biomaterial that is flexible, insensitive to dissolution/growth, and possesses mechanical strength.²⁴ In addition, HA nanoparticle self-assembly on the surface of PLGA can create a hydrophilic environment that allows cell attachment to microspheres, which can in theory promote drug release. Furthermore, HA is a hydroxide that can neutralize the acidity of PLGA degradation products, thereby improving osteoclast function.

Microspheres have been prepared by various methods, including water-in-oil-in-water (W/O/W),²⁵⁻²⁹ water-in-oil-in-oil (W/O/O),³⁰ oil-in-water (O/W),³¹ water-in-oil (W/O),³² solid-in-oil-in-water,^{33,34} solid-in-oil-in-oil,³⁵ solid-in-oil-in-oil-in-water,^{36,37} and water-in-oil-in-oil-in-water³⁸ emulsion. In addition, many techniques for generating porous microspheres have been developed.³⁸⁻⁴⁴ However, none of these approaches

^aShanghai Key Laboratory of Orthopedic Implants, Department of Orthopedic Surgery, Shanghai Ninth People's Hospital, Shanghai Jiao Tong University School of Medicine, 639 Zhizaoju Road, Shanghai 200011, China. E-mail: krdai@163.com; xinhua_qu@126.com

^bSchool of Pharmacy, Shanghai Jiao Tong University, 800 Dongchuan Road, Shanghai, China. E-mail: yuanweien@126.com

[†] Zhenyang Mao and Zhiwei Fang contributed equally to this work.



adequately counter the surface hydrophobicity that causes poor biocompatibility of PLGA or the acidity of its degradation products, which diminishes drug release and efficacy.

To address this issue, we developed a novel method for preparing microspheres, solid-in-oil-in-nanosuspension (S/O/Ns), using polyvinyl alcohol (PVA) and a HA nanosuspension as surfactants. The resultant nanoparticles were used as a carrier for strontium ranelate (SR), a drug used to treat osteoporosis that acts by reducing bone resorption and increasing new bone deposition by osteoblasts. We found that drug-loaded HA-PLGA microspheres provided sustained release of SR and enhanced osteogenesis in MC3T3-E1 cells.

2 Experimental

2.1 Preparation of microspheres

The synthesis of microspheres is illustrated in Fig. 1.

2.1.1 Materials. PLGA (65 : 35, molecular weight 20 kDa) was obtained from Lakeshore Biomaterials (Darmstadt, Germany). PVA (87–89% hydrolyzed, MW 31 000–50 000), HA nanoparticles, and SR were purchased from Sigma-Aldrich (St. Louis, MO, USA). Dichloromethane and other reagents were of analytical grade.

2.1.2 Preparation of porous PLGA microspheres (PM). Microspheres were prepared by double emulsion in water-in-oil-in-water (W/O/W). A 1 ml volume of water deionized with different volumes of NH_4HCO_3 (10% w/v) was added to 4 ml of methylene chloride with 250 mg PLGA. The solution was homogenized using a Powergen 700 homogenizer (Thermo Fisher Scientific, Waltham, MA, USA) for 3 min at 5000 rpm. The W/O emulsion was immediately poured into a beaker containing 200 ml of 0.1% (w/v) PVA, and re-emulsified at 200 rpm for 4 h using an overhead propeller (LR-400A; Thermo Fisher

Scientific). Microspheres were separated by centrifugation after evaporation of the solvent, washed three times with distilled water, and lyophilized in a freeze dryer, yielding the PM.

2.1.3 Preparation of 5% SR-loaded (SR-)PM. A 4 ml volume of methylene chloride was combined with 250 mg PLGA, 13.4 mg SR (freeze-dried powder), and 1 ml deionized water containing NH_4HCO_3 (10%, w/v) and dextran (5%). The mixture was homogenized at 5000 rpm for 3 min to obtain a primary solid-in-oil (S/O) emulsion, which was immediately poured into a beaker containing a 200 ml solution of 0.1% (w/v) PVA and re-emulsified at 200 rpm for 4 h using an overhead propeller. The microspheres were separated by centrifugation after evaporating the solvent, washed three times with distilled water, and lyophilized by freeze drying to obtain SR-PM.

2.1.4 Preparation of microspheres with surface nanoparticles. A suspension was formed by adding 100 mg HA nanoparticles to 4.5 ml of an aqueous solution of 2% (w/w) PVA followed by homogenization at 2000 rpm. A 10% (w/w) PLGA solution was dissolved in dichloromethane and the solution was combined with the HA nanoparticle suspension. The resultant O/Ns emulsion was transferred to 1000 ml of 5% (w/w) sodium chloride solution at 4 °C with gentle stirring (Xinhang JJ-1, Jintan Xinhang Co., Jintan, China) at 100 rpm for 2 h in order to evaporate the organic solvent. The lyophilized microspheres were rinsed with distilled water prior to storage.

2.1.5 Preparation of SR-PM with self-assembled HA nanoparticles on the surface (SR-PM-HA). A 4 ml volume of methylene chloride was combined with 250 mg PLGA, 13.4 mg SR, and 1 ml deionized water containing NH_4HCO_3 (10%, w/v) and homogenized at 5000 rpm for 3 min to obtain the primary S/O emulsion. This was immediately poured into a beaker containing 200 ml of 0.1% (w/v) PVA and 100 mg HA nanoparticle nanosuspension (S/O/Ns); the mixture was transferred to

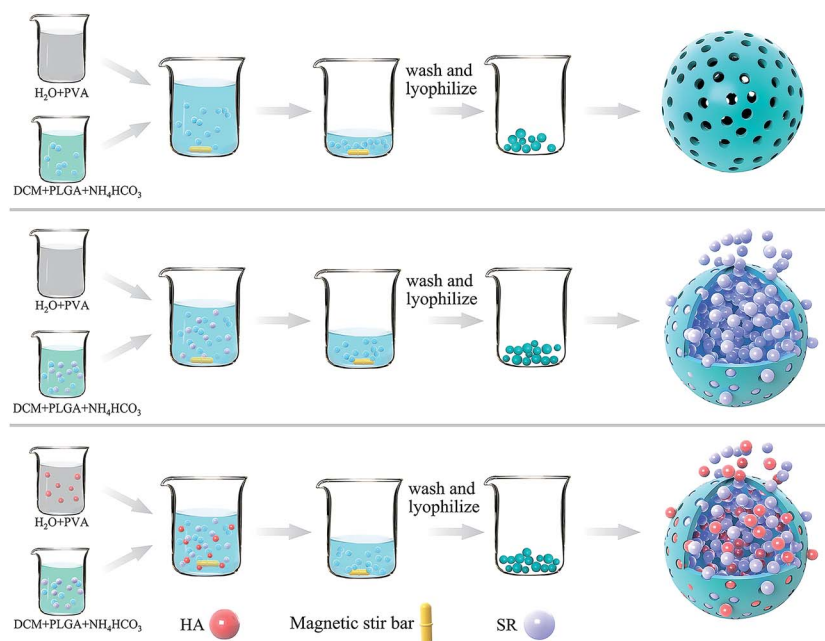


Fig. 1 Scheme of SR-PM-HA synthesis.



a beaker with 200 ml of 0.1% (w/v) PVA and re-emulsified at 200 rpm for 4 h using an overhead propeller. The microspheres were separated by centrifugation after evaporating the solvent, and washed three times with distilled water and lyophilized in a freeze dryer, yielding 5% SR-loaded PLGA microspheres with self-assembled HA (2%) nanoparticles (SR-PM-HA).

2.2 Characterization

2.2.1 Morphology. The morphology of PM, SR-PM, and SR-PM-HA was visualized by scanning electron microscopy (SEM) using a Sirion 200 instrument (FEI Tecnai, Hillsboro, OR, USA). Samples were coated with gold vapor at 5 keV under an argon atmosphere prior to scanning.

2.2.2 Encapsulation efficiency test. Drug-loaded PLGA microspheres (SR-PM or SR-PM-HA; 15 mg) were immersed in 5 ml dichloromethane and centrifuged for 5 min at 12 000 rpm in order to remove the PLGA. After re-dissolving in 3 ml phosphate-buffered saline (PBS, pH 7.4) followed by centrifugation for 5 min at 12 000 rpm, the SR content in the supernatant was analyzed by high-performance liquid chromatography (HPLC). The efficiency of SR encapsulation in the microspheres was calculated using the following equation:

$$\text{Encapsulation efficiency (\%)} = P/P_t \times 100$$

where P_t is the total theoretical weight of SR and P is the total actual weight of SR encapsulated in PLGA microspheres. The value was obtained from the results of three independent experiments.

SR release was measured by reversed-phase (RP)-HPLC at room temperature. A Diamonsil C18 column (250 × 4.6 mm, 5 μm) was used as the stationary phase, whereas the mobile phase consisted of 0.2% acetic acid buffer (pH = 5.5, adjusted with triethylamine):methanol(95 : 5). The eluent was monitored with an ultraviolet detector (234 nm) at a flow rate of 1. The linear response was above the concentration range of 10–200 μg ml⁻¹. The limits of detection and quantification were 20.0 and 50.0 ng, respectively.

2.2.3 Concentrations of key elements in PM. Concentrations of key elements (Sr for SR-PM and SR-PM-HA and P and Ca for SR-PM-HA) were determined by polarized Zeeman atomic absorption spectrophotometry (AAS) (Z-2000, Hitachi, Tokyo, Japan). SR content in SR-PM and SR-PM-HA, and SR and HA contents in SR-PM-HA were calculated.

2.2.4 Size distribution of microspheres. The average particle size and size distribution in microsphere preparations were determined with a particle size and shape analyzer (CIS-100; Ankersmid, Nijverdal, the Netherlands). Dry microsphere sample (10 mg) was placed in a quartz cuvette filled with isopropyl alcohol and measurements were made while stirring using a magnetic bar.

2.2.5 Changes in pH with PLGA degradation. PLGA microspheres (150 mg) were incubated at 37 °C in vials containing 5 ml PBS (100 mM, pH 7.4) with constant shaking. A 50 μl solution of penicillin (100 U ml⁻¹) and streptomycin (100 g ml⁻¹) was added. The pH of the buffer was measured every 5 days using an FE20 digital pH meter (Mettler Toledo,

Columbus, OH, USA). The experiment was repeated three times to obtain average pH values.

2.2.6 In vitro drug release. Drug-loaded PLGA microspheres (50 mg), including SR-PM and SR-PM-HA, were incubated at 37 °C in vials containing 5 ml PBS (100 mM, pH 7.4) with constant shaking. The buffer was collected at pre-determined time points and SR content was determined by RP-HPLC (as for the encapsulation efficiency test, Section 2.2.2). Three measurements were obtained for each sample to obtain average release profiles.

2.3 Evaluation of MC3T3-E1 cell proliferation

MC3T3-E1 cells were purchased from the Cell Bank of the Chinese Academy of Sciences (Shanghai, China). MC3T3-E1 cells were cultured in α-Minimal Essential Medium (α-MEM; Gibco, Grand Island, NY, USA) supplemented with 100 mg ml⁻¹ streptomycin, 100 U ml⁻¹ penicillin, and 10% fetal bovine serum (all from Hyclone, Logan, UT, USA) at 37 °C in a humidified atmosphere of 5% CO₂, with the medium changed every 3 days.

The effects of PM, SR-PM, and SR-PM-HA on cell proliferation were evaluated with Cell Counting Kit (CCK)-8 (Dojindo Laboratories, Kumamoto, Japan) according to the manufacturer's instructions. Cells were seeded at a density of 3 × 10³ per well in 96-well plates with each sample prepared in triplicate, and treated with the microspheres at a concentration of 5 mg ml⁻¹ 24 h later. CCK-8 reagent (10 μl) was added to each well and the plates were incubated for 2 h at 37 °C. The optical density was measured with an ELX800 microplate reader (Bio-Tek, Winooski, VT, USA) at a wavelength of 450 nm (650 nm for the reference).

2.4 Osteogenic differentiation of MC3T3-E1 cells

MC3T3-E1 cells (5 × 10⁵ per ml) were cultured with PM, SR-PM, or SR-PM-HA (5 mg ml⁻¹) in a 12-well plate. The medium was changed every 3 days. After culturing in α-MEM for 3 days, the medium was replaced with osteogenic medium composed of α-MEM supplemented with 50 mM L-ascorbic acid, 10⁻⁸ M dexamethasone, and 10 mM β-glycerophosphate (all from Sigma-Aldrich). At predetermined time points, cells were collected by digestion with trypsin and transferred to a 24-well plate.

Alkaline phosphatase (ALP) staining was carried out using a kit (Rainbow, Shanghai, China) according to the manufacturer's instructions. The expression of osteogenic differentiation marker genes such as ALP, runt-related transcription factor 2 (Runx-2), osteocalcin (OCN), and collagen type I (Col I) was assessed by real-time (RT)-PCR. RNA was extracted using TRIzol reagent (Invitrogen, Carlsbad, CA, USA) and cDNA was synthesized with the OmniScript Reverse Transcription kit (Qiagen, Valencia, CA, USA). SYBR Green (Bio-Rad, Hercules, CA, USA) was used along with the Gene Amp 2400 system (Perkin-Elmer, Waltham, MA, USA) for PCR amplification.⁴⁵

2.5 Statistical analysis

Data were analyzed using SPSS v.19.0 software (SPSS Inc., Chicago, IL, USA). Continuous variables are expressed as mean ±



SD. Between-group differences in demographic and clinical variables were evaluated by one-way analysis of variance, with the least significant difference and Dunnett's post-hoc tests used for variables with homogeneous and heterogeneous variances, respectively. A P value < 0.05 was considered statistically significant.

3 Results and discussion

3.1 Morphology and composition of microspheres

PLGA is biocompatible and biodegradable, applying in medicine for drug delivery and as biomaterials. However, poor hydrophilicity and acidic degradation products of PLGA limit its application.^{7–9} Poor hydrophilicity results in slow degradation rate in water. Although the hydrophilicity of PLGA could be modified by adjusting the ratio of lactic acid and glycolic acid,^{4,46,47} it is still undesirable. The novel S/O/Ns method we used is to solve the issues. The advantages of S/O/Ns were that we could not only fabricate PM loading specific drug, but bring HA nanoparticles self-assembled on the PM. After the fabrication, the morphology of PM, SR-PM, and SR-PM-HA was characterized by SEM. The porous structure could be clearly observed in three kinds of microspheres (Fig. 2a–c). The surfaces of PM and SR-PM were smooth whereas that of SR-PM-HA was rough. This was because that nanoparticles were uniformly self-assembled on the microsphere surface (Fig. 2c). The microspheres had an interconnected porous structure that was favorable for cell adhesion and proliferation (Fig. 2d). Particle size distribution results showed that the sizes of three kinds of microspheres were similar. They ranged in sizes from 400–1000 nm, 4–20 μm , and 20–200 μm (Fig. 3). The mean size was in the range of 20–200 μm . Generally, the size and the shape were appropriate for the further fabrication of porous microspheres.

In order to confirm that SR was loaded and HA nanoparticles were self-assembled on the microspheres, we

analyzed microsphere composition AAS (Fig. 4). PM did not contain Sr, P, or Ca; SR-PM contained Sr ($0.92\% \pm 0.14\%$) but not P or Ca; and SR-PM-HA contained Sr ($0.96\% \pm 0.15\%$), P ($0.78\% \pm 0.15\%$), and Ca ($0.21\% \pm 0.01\%$). The SR content of SR-PM and SR-PM-HA was $2.69\% \pm 0.46\%$ and $2.81\% \pm 0.44\%$, respectively; and the HA content of SR-PM-HA was $19.6\% \pm 0.52\%$ (Fig. 5). These results indicate that SR was effectively encapsulated by PM and HA on the PM surface.

3.2 Degradation and drug loading-release property of SR-PM and SR-PM-HA

In an aqueous environment, PLGA is degraded by hydrolysis of its ester linkages into lactic and glycolic acids, which can activate an inflammatory response. Microspheres with HA nanoparticles can neutralize the acidity of PLGA degradation products. After 2 weeks in solution, pH decreased markedly, with no difference observed between PM and PM-HA (Fig. 6). However, in the subsequent 3 weeks, PM-HA stabilized the pH above 7.1. It is possible that alkaline HA nanoparticles released from PM quenched hydrogen ions that dissociated from lactic and glycolic acids.

ASS results showed that the SR content of SR-PM and SR-PM-HA was $2.69\% \pm 0.46\%$ and $2.81\% \pm 0.44\%$ (Fig. 5). Real total weight of SR encapsulated in PM was regarded as A mg PM; the total weight of SR added to the PM formulation was 13.4 mg and the total amount of obtained PM was B mg. SR loading in microspheres was calculated with the following equations.

$$\text{Encapsulation efficiency} = (A) \times (B \div 15) \times 100\% \div 13.4$$

$$\text{Encapsulation efficiency of SR-PM} = 54.02\% \pm 9.26\%$$

$$\text{Encapsulation efficiency of SR-PM-HA} = 56.45\% \pm 8.35\%$$

The percent cumulative release of SR from SR-PM was 90% in 22 days as compared to 85% from SR-PM-HA (Fig. 7). In both

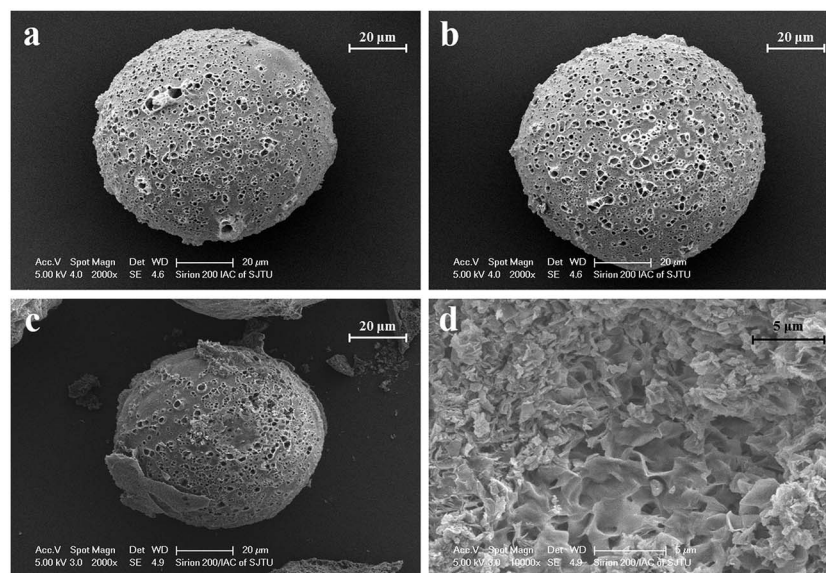


Fig. 2 Scanning electron micrographs of (a) PM, (b) SR-PM, and (c) SR-PM-HA. (d) PM cross section.



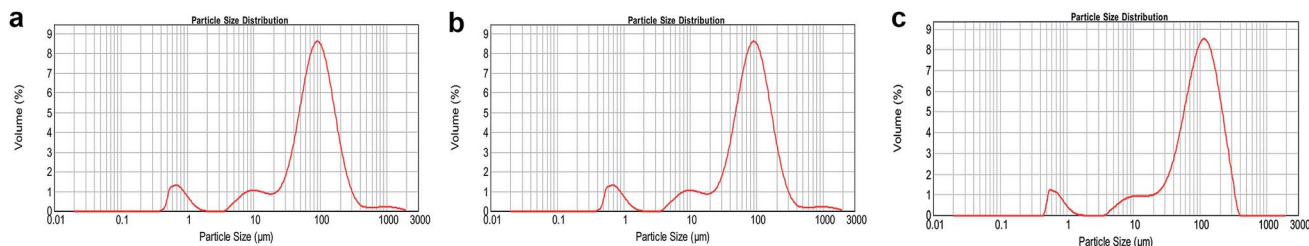


Fig. 3 Size distributions of particles. (a) PM, (b) SR-PM, and (c) SR-PM-HA size ranged from 400–1000 nm, 4–20 μm , and 20–200 μm . The mean PM droplet size was 20–200 μm .

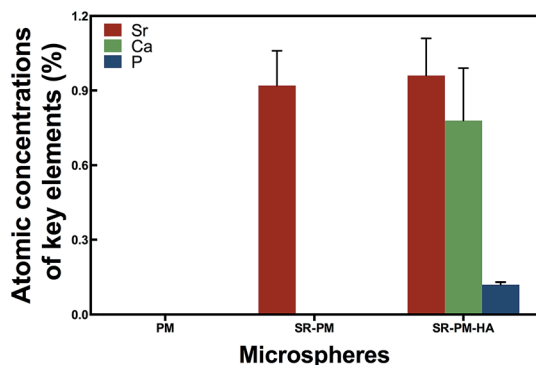


Fig. 4 Concentrations of key elements in PM. Data represent mean \pm standard deviation. SR-PM contained Sr ($0.92\% \pm 0.14\%$) and SR-PM-HA contained Sr ($0.96\% \pm 0.15\%$), P ($0.785 \pm 0.15\%$), and Ca ($0.21\% \pm 0.01\%$). PM contained none of these elements.

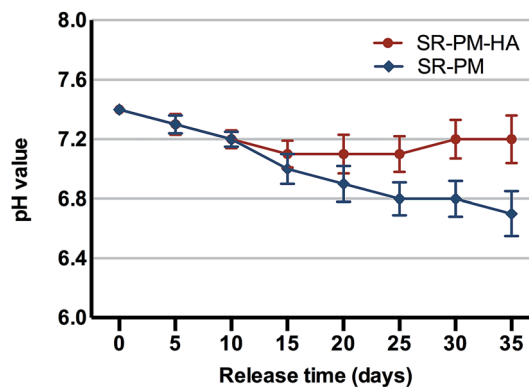


Fig. 6 Changes in pH with PLGA degradation. Data represent mean \pm standard deviation. During the first 2 weeks in solution, pH decreased markedly and there were no differences between samples. In the subsequent 3 weeks, PM with HA nanoparticles stabilized the pH above 7.1.

cases the release was gradual, with no burst release observed. PLGA is a biocompatible and biodegradable material that can be used for drug delivery.^{2,4,6,48,49} Drugs are released by diffusion, erosion of the polymer, or both. In our study, drug-loaded microspheres showed sustained rather than burst release, which is a desirable attribute for a drug carrier. The time-course of release from SR-PM can be described by first-order kinetics

(*i.e.*, gradual decrease in release rate as the amount of remaining drug decreases) (Fig. 7). However, SR-PM-HA exhibited zero-order kinetics (*i.e.*, release rate is constant even when the amount of remaining drug decreases). This is reasonable because nanoparticles self-assembled on the surface of microspheres are surfactants; moreover, microspheres with self-assembled HA nanoparticles have a large specific surface area and HA nanoparticles occupy a large portion of the microsphere

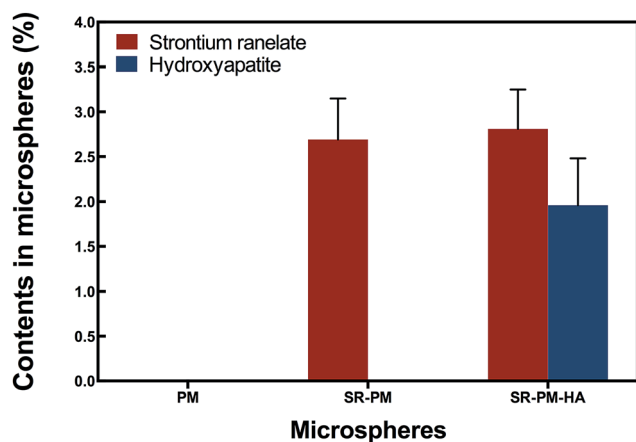


Fig. 5 SR and HA concentrations in microspheres. Data represent mean \pm standard deviation. The SR contents of SR-PM and SR-PM-HA were $2.69\% \pm 0.46\%$ and $2.81\% \pm 0.44\%$, respectively. The HA content of SR-PM-HA was $19.6\% \pm 0.52\%$.

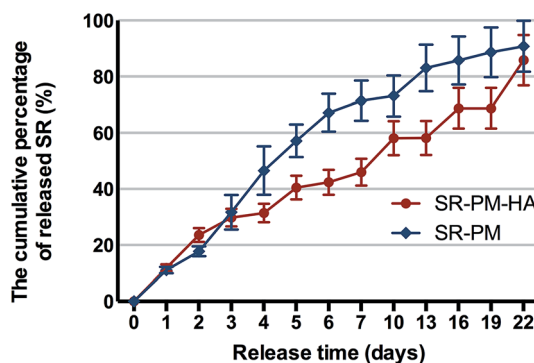


Fig. 7 Cumulative drug release percentage. Data represent mean \pm standard deviation. Cumulative percentage of released SR was higher for SR-PM than for SR-PM-HA (90% vs. 85% of total loaded drug in 22 days).



surface, which could reduce the amount of drug that can bind. Finally, we concluded that HA nanoparticles could regulate the pH of the microenvironment with PLGA degradation, which causes local acidity; moreover, SR contains carboxyl, which dissolves in alkaline environments. This can explain the constant release rate of SR-PM-HA. We propose that the S/O/Ns method reduced the amount of drug on the microsphere surface and thereby prevented burst release. Self-assembled HA enabled sustained drug release from PLGA microspheres.

3.3 Biocompatibility and osteoinductivity of SR-PM-HA

MC3T3-E1 cells were cultured with PM, SR-PM, and SR-PM-HA for various times and proliferation was evaluated with the CCK-8 assay (Fig. 8). Compared to PM, SR and SR-PM-HA showed no cytotoxicity. Moreover, after 3 days, SR-PM and SR-PM-HA stimulated cell proliferation to similar degrees. These results indicate that the microspheres are biocompatible and are thus suitable as biomaterials. SR has been shown to induce preosteoblast growth⁵⁰ and has a positive effect on MC3T3-E1 cell proliferation and differentiation, consistent with our observations.⁵¹ SR was also found to influence cellular processes *via* interactions with calcium-sensing receptors, which leads to upregulation of ETS-related gene 1 and *c-fos*, two regulators of osteoblast proliferation.⁵²

The osteoinductive activity of PM, SR-PM, and SR-PM-HA was investigated by examining MC3T3-E1 cell differentiation after 7 and 14 days based on ALP staining and the expression of osteogenic differentiation markers (Fig. 9). At both time points, ALP staining intensity was highest for SR-PM-HA, followed by SR-PM and PM. We also examined the expression of the osteoblast markers ALP, Col I, OCN, and Runx-2 (ref. 53–55) by RT-PCR. After 3 days, there was no change in ALP expression in cells cultured with PM, SR-PM, or SR-PM-HA ($P > 0.05$). However, at 7 and 14 days, ALP level was higher in cells cultured with SR-PM-HA than in those cultured with SR-PM or PM ($P < 0.05$). Similarly, levels of Col I and OCN were highest in the SR-

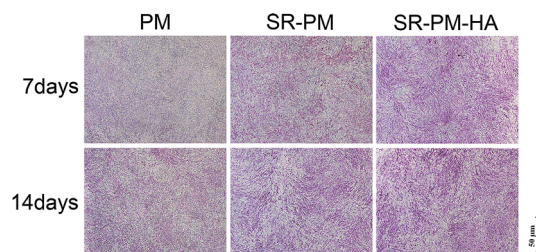


Fig. 9 ALP activity in MC3T3-E1 cells cultured for 7 and 14 days in osteogenic medium with PM, SR-PM, or SR-PM-HA. At both time points, ALP staining intensity was highest in the presence of SR-PM-HA, followed by SR-PM and PM.

PM-HA group ($P < 0.05$), followed by SR-PM and PM at 14 days. Runx-2 expression was upregulated in cells cultured with SR-PM and SR-PM-HA at 3 and 7 days as compared to those cultured with PM ($P < 0.05$). The level was decreased at 14 days, although there were no differences observed between SR-PM and SR-PM-HA at any time point ($P > 0.05$) (Fig. 10).

Osteoporosis is a systemic disease that is characterized by osteopenia and bone tissue damage, which increases bone fragility and risk of fracture.⁵⁶ Several pharmacologic options are available for osteoporosis treatment, including selective estrogen receptor modulators, bisphosphonates, calcitonin, strontium, and parathyroid hormone. Whereas most of these only inhibit bone resorption by osteoclasts, the latter two are dual-action bone agents that also induce osteogenesis by osteoblasts.^{57–59} In the present study, SR stimulated osteogenesis, as evidenced by ALP staining and expression of osteogenic differentiation markers. Osteoblast differentiation is characterized by cell proliferation and matrix mineralization and maturation. ALP is mainly expressed in matrix vesicles or at the cell surface and degrades organic phosphoesters in cartilage and bone.⁶⁰ As differentiation proceeds, ALP expression gradually increases, peaking as cells enter the matrix maturation stage before decreasing during matrix mineralization. Col I is the most abundant component of mature osteoblasts. OCN is a late bone marker that plays an important role in bone mineralization, which is the final step of osteogenesis. Thus, these three marker genes are sequentially expressed during osteogenesis. On the other hand, Runx-2 is expressed at an early stage.⁵³ ALP staining intensity and the expression of these markers were higher in cells cultured with SR-PM or SR-PM-HA than with PM, indicating that SR enhanced osteogenesis; this is likely achieved *via* activation of Wnt and other signaling pathways.⁶¹ Our observation that SR stimulates the expression of osteoblastic differentiation markers is similar to previous findings in primary murine osteoblasts⁶² and is in accordance with reports that SR induces osteogenesis by increasing osteoblast differentiation and bone matrix synthesis and mineralization.^{63–68}

PM containing HA have excellent biocompatibility and can be used to engineer bone tissue, since they can enhance osteoblast migration, attachment, and differentiation.⁶⁹ We suggest that preparing PM by the S/O/Ns method can improve surface hydrophilicity and consequently, biocompatibility; moreover, HA may neutralize the acidity of PLGA degradation products

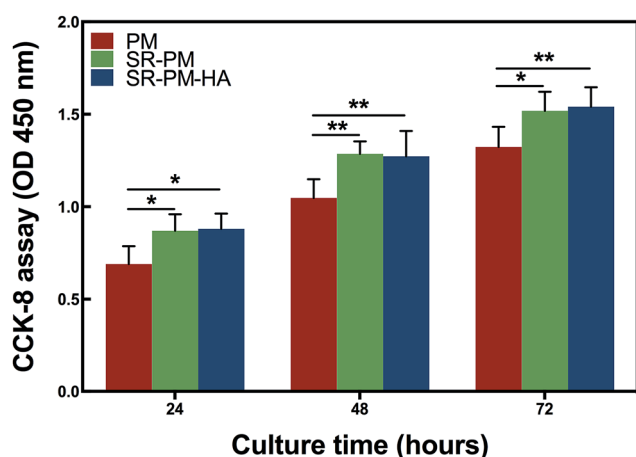


Fig. 8 Proliferation of MC3T3-E1 cells cultured with PM, SR-PM, or SR-PM-HA for indicated times. Data represent mean \pm standard deviation. After 3 days, SR-PM and SR-PM-HA similarly stimulated cell proliferation relative to PM (* for $P < 0.05$, ** for $P < 0.01$).



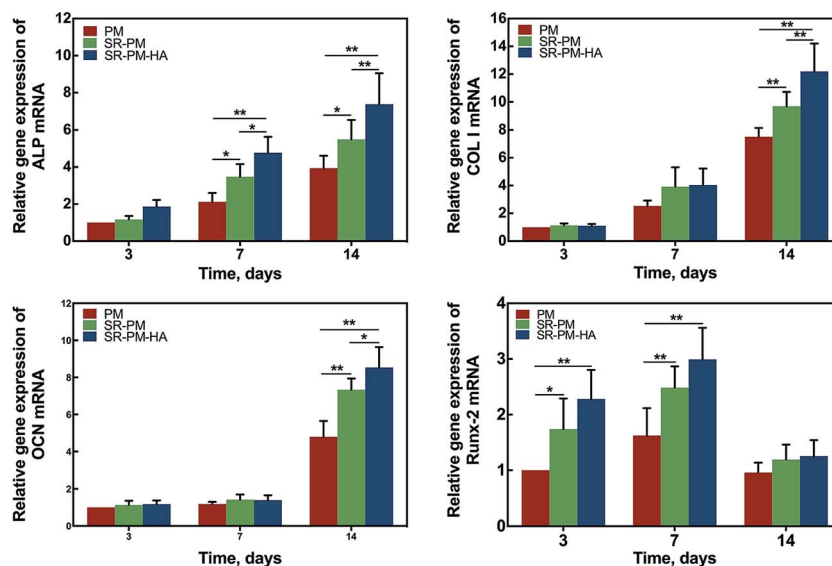


Fig. 10 Relative expression of osteogenic differentiation marker genes. Data represent mean \pm standard deviation. After 3 days, ALP expression did not differ among MC3T3-E1 cells cultured with PM, SR-PM, or SR-PM-HA. At 7 and 14 days, ALP expression was higher in cells cultured with SR-PM-HA as compared to SR-PM or PM. Col I and OCN levels were highest in the presence of SR-PM-HA followed by SR-PM and PM at 14 days. RunX-2 expression was higher in cells cultured with SR-PM and SR-PM-HA as compared to PM at 3 and 7 days; after 14 days, RunX-2 levels decreased. There was no difference in RunX-2 level between SR-PM and SR-PM-HA at any time point (* for $P < 0.05$, ** for $P < 0.01$).

and thus prevent burst release as well as an inflammatory response. We also speculate that microspheres promote osteoblast attachment, migration, and differentiation by HA self-assembly on the PM surface and drug release.

HA has similar properties to bone minerals^{70,71} and has been used in biomaterials and as adsorbents or catalysts.^{72–76} It is considered suitable for bone regeneration, particularly in orthopedic implants to accelerate the growth of surrounding bone without causing systemic toxicity or an immune reaction.^{77,78} Various methods have been developed to generate HA molecules with specific crystallinity, particle size, and morphology.^{79,80} Owing to its biocompatibility, biodegradability, inorganic properties, and ease of fabrication, HA is not only by itself an excellent biomaterial for implants^{81–85} but can also be combined with other materials for a variety of applications, such as implant coatings^{86–90} or fabrication of nanomaterials.^{76,91,92} HA in conjunction with other biodegradable microspheres has been used to construct an injectable scaffold⁹³ that promoted osteoblast attachment and proliferation and ALP activity. Previous studies have reported good biocompatibility of self-assembled HA molecules,^{86,94} consistent with our results.

4 Conclusion

SR-PM-HA has been fabricated by the S/O/Ns method according to the following stages: (i) preparation of S/O emulsion with methylene chloride combined with PLGA, SR and NH_4HCO_3 , (ii) preparation of O/Ns emulsion formed by suspension with HA nanoparticles to PVA, (iii) and combination of S/O and O/Ns. SR-PM-HA has no cytotoxicity and performs satisfying drug release property, so it is suitable to be applied as drug delivery systems. SR-PM-HA could effectively promote the proliferation and

stimulate the osteogenic differentiation of MC3T3-E1 cells. Our findings indicate that SR-PM-HA is a promising biomaterial for the treatment of bone defects by bone regeneration. To better investigate the osteoinductivity ability of SR-PM-HA, we would use human bone marrow stem cells for *in vitro* experiments and confirm our observations *in vivo* in the future.

Acknowledgements

This work was supported by the National Natural Science Foundation for Youths (Grant No. 81401852), This work was supported by the National Natural Science Foundation (Grant No. 51631009 and 81373366), the National Science Foundation of Shanghai (No. 14ZR1424000), “Chen Guang” Project of the Shanghai Municipal Education Commission, Shanghai Education Development Foundation (No. 14CG14) and Funds for Interdisciplinary Projects of Medicine and Engineering of Shanghai JiaoTong University (No. YG2014QN06, YG2015MS06, YG2015QN13).

References

- 1 C. Bouissou, J. J. Rouse, R. Price and C. F. van der Walle, *Pharm. Res.*, 2006, **23**, 1295–1305.
- 2 R. A. Jain, *Biomaterials*, 2000, **21**, 2475–2490.
- 3 J. Zhou, S. Han, Y. Dou, J. Lu, C. Wang, H. He, X. Li and J. Zhang, *Int. J. Pharm.*, 2013, **448**, 175–188.
- 4 S. C. Loo, Z. Y. Tan, Y. J. Chow and S. L. Lin, *J. Pharm. Sci.*, 2010, **99**, 3060–3071.
- 5 J. M. Chan, L. Zhang, K. P. Yuet, G. Liao, J. W. Rhee, R. Langer and O. C. Farokhzad, *Biomaterials*, 2009, **30**, 1627–1634.



- 6 H. Wang, P. Zhao, W. Su, S. Wang, Z. Liao, R. Niu and J. Chang, *Biomaterials*, 2010, **31**, 8741–8748.
- 7 Z. Zhai, X. Qu, H. Li, K. Yang, P. Wan, L. Tan, Z. Ouyang, X. Liu, B. Tian, F. Xiao, W. Wang, C. Jiang, T. Tang, Q. Fan, A. Qin and K. Dai, *Biomaterials*, 2014, **35**, 6299–6310.
- 8 W. Liu, T. Wang, C. Yang, B. W. Darvell, J. Wu, K. Lin, J. Chang, H. Pan and W. W. Lu, *Osteoporosis Int.*, 2016, **27**, 93–104.
- 9 T. R. Arnett, *J. Nutr.*, 2008, **138**, 415S–418S.
- 10 M. Adabi, M. Naghibzadeh, M. Adabi, M. A. Zarrinfard, S. S. Esnaashari, A. M. Seifalian, R. Faridi-Majidi, H. Tanimowo Aiyelabegan and H. Ghanbari, *Artif. Cells, Nanomed., Biotechnol.*, 2017, **45**, 833–842.
- 11 K. Ariga, *Anal. Sci.*, 2016, **32**, 1141–1149.
- 12 Y. Cui, B. Li, H. He, W. Zhou, B. Chen and G. Qian, *Acc. Chem. Res.*, 2016, **49**, 483–493.
- 13 N. C. Huang, Q. Ji, T. Yamazaki, W. Nakanishi, N. Hanagata, K. Ariga and S. H. Hsu, *Phys. Chem. Chem. Phys.*, 2015, **17**, 25455–25462.
- 14 Z. Huang and S. Che, *Bull. Chem. Soc. Jpn.*, 2015, **88**, 617–632.
- 15 Z. Huang, Y. Yao, L. Han and S. Che, *Chemistry*, 2014, **20**, 17068–17076.
- 16 Y. Jiao, X. Pang, M. Liu, B. Zhang, L. Li and G. Zhai, *Colloids Surf., B*, 2016, **140**, 361–372.
- 17 B. Li, H. M. Wen, Y. Cui, W. Zhou, G. Qian and B. Chen, *Adv. Mater.*, 2016, **28**, 8819–8860.
- 18 V. Malgras, Q. Ji, Y. Kamachi, T. Mori, F.-K. Shieh, K. C.-W. Wu, K. Ariga and Y. Yamauchi, *Bull. Chem. Soc. Jpn.*, 2015, **88**, 1171–1200.
- 19 Q. Tang, J. Liu, L. K. Shrestha, K. Ariga and Q. Ji, *ACS Appl. Mater. Interfaces*, 2016, **8**, 18922–18929.
- 20 I. Uddin, S. Venkatachalam, A. Mukhopadhyay and M. A. Usmani, *Curr. Pharm. Des.*, 2016, **22**, 1472–1484.
- 21 J. Li, Y. Qiao and Z. Wu, *J. Controlled Release*, 2017, **256**, 9–18.
- 22 T. Matoba, J. I. Koga, K. Nakano, K. Egashira and H. Tsutsui, *J. Cardiol.*, 2017, DOI: 10.1016/j.jcc.2017.03.005.
- 23 N. Rahoui, B. Jiang, N. Taloub and Y. D. Huang, *J. Controlled Release*, 2017, **255**, 176–201.
- 24 X. Liu, M. Zhao, J. Lu, J. Ma, J. Wei and S. Wei, *Int. J. Nanomed.*, 2012, **7**, 1239–1250.
- 25 I. J. Castellanos, G. Cruz, R. Crespo and K. Griebenow, *J. Controlled Release*, 2002, **81**, 307–319.
- 26 A. G. Coombes, M. K. Yeh, E. C. Lavelle and S. S. Davis, *J. Controlled Release*, 1998, **52**, 311–320.
- 27 S. Freitas, H. P. Merkle and B. Gander, *J. Controlled Release*, 2005, **102**, 313–332.
- 28 J. H. Kim and Y. H. Bae, *Eur. J. Pharm. Sci.*, 2004, **23**, 245–251.
- 29 S. Takada, Y. Yamagata, M. Misaki, K. Taira and T. Kurokawa, *J. Controlled Release*, 2003, **88**, 229–242.
- 30 W. Lu and T. G. Park, *PDA J. Pharm. Sci. Technol.*, 1995, **49**, 13–19.
- 31 T. G. Park, H. Yong Lee and Y. Sung Nam, *J. Controlled Release*, 1998, **55**, 181–191.
- 32 L. Jorgensen, E. H. Moeller, M. van de Weert, H. M. Nielsen and S. Frokjaer, *Eur. J. Pharm. Sci.*, 2006, **29**, 174–182.
- 33 T. Morita, Y. Sakamura, Y. Horikiri, T. Suzuki and H. Yoshino, *J. Controlled Release*, 2000, **69**, 435–444.
- 34 T. Ren, W. Yuan, H. Zhao and T. Jin, *Micro Nano Lett.*, 2011, **6**, 70–74.
- 35 K. G. Carrasquillo, A. M. Stanley, J. C. Aponte-Carro, P. De Jesus, H. R. Costantino, C. J. Bosques and K. Griebenow, *J. Controlled Release*, 2001, **76**, 199–208.
- 36 W. Yuan, F. Wu, Y. Geng, S. Xu and T. Jin, *Int. J. Pharm.*, 2007, **339**, 76–83.
- 37 W. Yuan and Z. Liu, *Int. J. Nanomed.*, 2012, **7**, 257–270.
- 38 X. Hong, L. Wei, L. Ma, Y. Chen, Z. Liu and W. Yuan, *Int. J. Nanomed.*, 2013, **8**, 2433–2441.
- 39 Y. Cai, Y. Chen, X. Hong, Z. Liu and W. Yuan, *Int. J. Nanomed.*, 2013, **8**, 1111–1120.
- 40 C. Z. Zhang, J. Niu, Y. S. Chong, Y. F. Huang, Y. Chu, S. Y. Xie, Z. H. Jiang and L. H. Peng, *Eur. J. Pharm. Biopharm.*, 2016, **109**, 1–13.
- 41 K. Jiang, H. Zhao, J. Dai, D. Kuang, T. Fei and T. Zhang, *ACS Appl. Mater. Interfaces*, 2016, **8**, 25529–25534.
- 42 J. W. Park, Y. P. Yun, K. Park, J. Y. Lee, H. J. Kim, S. E. Kim and H. R. Song, *Colloids Surf., B*, 2016, **147**, 265–273.
- 43 Y. Wei, Y. Wang, H. Zhang, W. Zhou and G. Ma, *J. Colloid Interface Sci.*, 2016, **478**, 46–53.
- 44 Q. Xiao, K. Zhou, C. Chen, M. Jiang, Y. Zhang, H. Luo and D. Zhang, *Mater. Sci. Eng., C*, 2016, **69**, 1068–1074.
- 45 P. Kasten, R. Luginbuhl, M. van Griensven, T. Barkhausen, C. Krettek, M. Bohner and U. Bosch, *Biomaterials*, 2003, **24**, 2593–2603.
- 46 A. C. Grayson, G. Voskerician, A. Lynn, J. M. Anderson, M. J. Cima and R. Langer, *J. Biomater. Sci., Polym. Ed.*, 2004, **15**, 1281–1304.
- 47 L. Lu, S. J. Peter, M. D. Lyman, H. L. Lai, S. M. Leite, J. A. Tamada, S. Uyama, J. P. Vacanti, R. Langer and A. G. Mikos, *Biomaterials*, 2000, **21**, 1837–1845.
- 48 Q. Wang, J. Wang, Q. Lu, M. S. Detamore and C. Berkland, *Biomaterials*, 2010, **31**, 4980–4986.
- 49 T. G. Kim, H. Lee, Y. Jang and T. G. Park, *Biomacromolecules*, 2009, **10**, 1532–1539.
- 50 J. Caverzasio, *Bone*, 2008, **42**, 1131–1136.
- 51 S. Takaoka, T. Yamaguchi, S. Yano, M. Yamauchi and T. Sugimoto, *Horm. Metab. Res.*, 2010, **42**, 627–631.
- 52 N. Chattopadhyay, S. J. Quinn, O. Kifor, C. Ye and E. M. Brown, *Biochem. Pharmacol.*, 2007, **74**, 438–447.
- 53 P. Ducy, R. Zhang, V. Geoffroy, A. L. Ridall and G. Karsenty, *Cell*, 1997, **89**, 747–754.
- 54 R. T. Franceschi, *Crit. Rev. Oral Biol. Med.*, 1999, **10**, 40–57.
- 55 Y. Wang, T. Uemura, J. Dong, H. Kojima, J. Tanaka and T. Tateishi, *Tissue Eng.*, 2003, **9**, 1205–1214.
- 56 E. S. Siris, R. Adler, J. Bilezikian, M. Bolognese, B. Dawson-Hughes, M. J. Favus, S. T. Harris, S. M. Jan de Beur, S. Khosla, N. E. Lane, R. Lindsay, A. D. Nana, E. S. Orwoll, K. Saag, S. Silverman and N. B. Watts, *Osteoporosis Int.*, 2014, **25**, 1439–1443.
- 57 A. S. Hurtel-Lemaire, R. Mentaverri, A. Caudrillier, F. Cournarie, A. Wattel, S. Kamel, E. F. Terwilliger, E. M. Brown and M. Brazier, *J. Biol. Chem.*, 2009, **284**, 575–584.



- 58 T. C. Brennan, M. S. Rybchyn, W. Green, S. Atwa, A. D. Conigrave and R. S. Mason, *Br. J. Pharmacol.*, 2009, **157**, 1291–1300.
- 59 G. J. Atkins, K. J. Welldon, P. Halbout and D. M. Findlay, *Osteoporosis Int.*, 2009, **20**, 653–664.
- 60 D. A. Wang, C. G. Williams, F. Yang, N. Cher, H. Lee and J. H. Elisseeff, *Tissue Eng.*, 2005, **11**, 201–213.
- 61 G. A. Fielding, W. Smoot and S. Bose, *J. Biomed. Mater. Res., Part A*, 2014, **102**, 2417–2426.
- 62 E. Bonnelye, A. Chabadel, F. Saltel and P. Jurdic, *Bone*, 2008, **42**, 129–138.
- 63 S. Choudhary, P. Halbout, C. Alander, L. Raisz and C. Pilbeam, *J. Bone Miner. Res.*, 2007, **22**, 1002–1010.
- 64 E. Canalis, M. Hott, P. Deloffre, Y. Tsouderos and P. J. Marie, *Bone*, 1996, **18**, 517–523.
- 65 W. Querido, M. Farina and K. Anselme, *Biomatter*, 2015, **5**, e1027847.
- 66 F. Yang, D. Yang, J. Tu, Q. Zheng, L. Cai and L. Wang, *Stem Cell.*, 2011, **29**, 981–991.
- 67 P. J. Marie, *Bone*, 2006, **38**, S10–S14.
- 68 P. Ammann, *Bone*, 2006, **38**, 15–18.
- 69 G. Xiao, H. Yin, W. Xu and Y. Lu, *J. Biomater. Sci., Polym. Ed.*, 2016, **27**, 1462–1475.
- 70 M. Al-Jawad, A. Steuwer, S. H. Kilcoyne, R. C. Shore, R. Cywinski and D. J. Wood, *Biomaterials*, 2007, **28**, 2908–2914.
- 71 P. Malmberg and H. Nygren, *Proteomics*, 2008, **8**, 3755–3762.
- 72 Y. Matsuura, A. Onda and K. Yanagisawa, *Catal. Commun.*, 2014, **48**, 5–10.
- 73 F. J. O'Brien, *Mater. Today*, 2011, **14**, 88–95.
- 74 S. H. Tan, X. G. Chen, Y. Ye, J. Sun, L. Q. Dai and Q. Ding, *J. Hazard. Mater.*, 2010, **179**, 559–563.
- 75 F. Ye, H. Guo, H. Zhang and X. He, *Acta Biomater.*, 2010, **6**, 2212–2218.
- 76 W. P. Wijesinghe, M. M. Mantilaka, K. G. Chathuranga Senarathna, H. M. Herath, T. N. Premachandra, C. S. Ranasinghe, R. P. Rajapakse, R. M. Rajapakse, M. Edirisinghe, S. Mahalingam, I. M. Bandara and S. Singh, *Mater. Sci. Eng., C*, 2016, **63**, 172–184.
- 77 T. Kokubo and H. Takadama, *Biomaterials*, 2006, **27**, 2907–2915.
- 78 P. O'Hare, B. J. Meenan, G. A. Burke, G. Byrne, D. Dowling and J. A. Hunt, *Biomaterials*, 2010, **31**, 515–522.
- 79 J. D. Chen, Y. J. Wang, K. Wei, S. H. Zhang and X. T. Shi, *Biomaterials*, 2007, **28**, 2275–2280.
- 80 F. Chen, Q. L. Tang, Y. J. Zhu, K. W. Wang, M. L. Zhang, W. Y. Zhai and J. Chang, *Acta Biomater.*, 2010, **6**, 3013–3020.
- 81 Z. Zhao, M. Espanol, J. Guillem-Marti, D. Kempf, A. Diez-Escudero and M. P. Ginebra, *Nanoscale*, 2016, **8**, 1595–1607.
- 82 J. Yu, H. Yang, K. Li, J. Lei, L. Zhou and C. Huang, *J. Dent.*, 2016, **50**, 21–29.
- 83 S. Pujari-Palmer, S. Chen, S. Rubino, H. Weng, W. Xia, H. Engqvist, L. Tang and M. K. Ott, *Biomaterials*, 2016, **90**, 1–11.
- 84 N. S. Remya, S. Syama, V. Gayathri, H. K. Varma and P. V. Mohanan, *Colloids Surf., B*, 2014, **117**, 389–397.
- 85 D. Gopi, M. T. Ansari, E. Shinyjoy and L. Kavitha, *Spectrochim. Acta, Part A*, 2012, **87**, 245–250.
- 86 J. Shen, Y. Qi, B. Jin, X. Wang, Y. Hu and Q. Jiang, *J. Biomed. Mater. Res., Part B*, 2017, **105**, 124–135.
- 87 A. Pommer, G. Muhr and A. David, *J. Bone Jt. Surg., Am. Vol.*, 2002, **84**, 1162–1166.
- 88 C. Knabe, G. Berger, R. Gildenhaar, F. Klar and H. Zreiqat, *Biomaterials*, 2004, **25**, 4911–4919.
- 89 J. C. Babister, L. A. Hails, R. O. Oreffo, S. A. Davis and S. Mann, *Biomaterials*, 2009, **30**, 3174–3182.
- 90 D. Li, C. Ye, Y. Zhu, Z. Gou and C. Gao, *J. Biomed. Mater. Res., Part B*, 2012, **100**, 1103–1113.
- 91 Y. Wan, C. Wu, G. Xiong, G. Zuo, J. Jin, K. Ren, Y. Zhu, Z. Wang and H. Luo, *J. Mech. Behav. Biomed. Mater.*, 2015, **47**, 29–37.
- 92 B. Ma, S. Zhang, R. Liu, J. Qiu, L. Zhao, S. Wang, J. Li, Y. Sang, H. Jiang and H. Liu, *Nanoscale*, 2017, **9**, 2162–2171.
- 93 X. Hu, H. Shen, F. Yang, X. Liang, S. Wang and D. Wu, *Appl. Surf. Sci.*, 2014, **292**, 764–772.
- 94 J. Chen, Z. Wang, Z. Wen, S. Yang, J. Wang and Q. Zhang, *Colloids Surf., B*, 2015, **127**, 47–53.

