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13-93 bioactive glass/alginate composite scaffolds 3D printed under mild conditions for bone regeneration

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Composite scaffolds of type 13-93 bioactive glass (13-93 BG) and sodium alginate (SA), denoted 13-93 BG/SA, in mass ratios of 0 : 4, 1 : 4, 2 : 4 and 4 : 4 were prepared for bone regeneration by 3D printing under mild conditions. All obtained scaffolds were completely interconnected. As the 13-93 BG/SA mass ratio increased from 0 : 4 to 4 : 4, the pore size improved from 340 ± 27 to 530 ± 27 μm , porosity increased from 65.5 ± 5.3 to $87.5 \pm 3.9\%$ and shrinkage ratios decreased by half. Moreover, 13-93 BG incorporation improved the compressive strength and modulus of the SA scaffold, achieving highest values of 16.74 ± 1.78 MPa and 79.49 ± 7.38 MPa, respectively, in 13-93 BG/SA with a 2 : 4 mass ratio. Furthermore, *in vitro* apatite mineralization, the release of bioactive ions (Mg^{2+} and SiO_4^{4-}) and a weakly alkaline pH were exhibited by scaffolds containing 13-93 BG, which further promoted the attachment, proliferation and osteogenic differentiation of rBMSCs on composite scaffolds. As a result, 3D-printed 13-93 BG/SA scaffold was an ideal candidate for bone regeneration.

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Introduction

A new generation of tissue engineering scaffolds with good interconnection, biodegradability, bioactivity and adequate mechanical support is required for bone regeneration.^{1,2} Tissue engineering scaffolds have many essential features, among which composition and macro/nanoporous connected morphology are thought to be critical in influencing cellular response. Biodegradable organic-inorganic composite scaffolds have recently attracted attention as bone tissue mimics.³⁻⁶ Their main advantage is that altering the organic/inorganic material composition or ratio can change the properties of the resultant composite scaffold to satisfy the requirements of bone regeneration.⁵ Organic materials used for bone tissue engineering applications are usually natural polymers (such as collagen, chitosan and alginate) or synthetic polymers (such as polylactic acid (PLA) and polycaprolactone (PCL)), because of their biocompatibility and biodegradation.^{7,8} Inorganic materials used are usually hydroxyapatite (HAp), β -tricalcium phosphate (β -TCP) and bioactive glasses because of their osteoconductivity.⁹⁻¹²

The macro/nanoporous connected morphology of composite scaffolds has been investigated using various methods, most notably solvent casting, particulate leaching and freeze-drying.¹³⁻¹⁶ However, these conventional methods have many limitations.¹⁶ For example, the pore size, morphology and distribution, and the external scaffold geometry, are difficult to control. Moreover, the application of non-physiological conditions prevents incorporation of biological components during the scaffold fabrication process. In the last decade, the development of 3D printing technology has allowed the fabrication of scaffolds with controlled and regular pore sizes, shapes and interconnectivity using computer aided design and computer aided manufacturing (CAM), suggesting that 3D printing could be used to construct individual scaffolds for bone regeneration.^{4,17-20} Furthermore, composite scaffolds for tissue engineering have been fabricated by 3D printing under mild conditions (room or physiological temperature and organic-free solvent), which should be beneficial for designing multifunctional scaffolds containing bioactive proteins or living cells for medical therapy.^{4,5,17}

Based on this background, biodegradable organic-inorganic composite scaffolds for bone regeneration were fabricated in this present study using 3D printing under mild conditions. Biocompatible and biodegradable alginate has been widely used in tissue engineering, growth factor and cell delivery,^{12,20-23} and was selected as the organic component in this study. It is also a suitable biomaterial for preparing functional hydrogels, due to its water-solubility and cross-linking with ionic response. Furthermore, the sol-gel transition of alginate induced by

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multivalent cations, such as Ca^{2+} , under mild conditions has been successfully developed as a novel stabilization method for 3D printing, which would be convenient for encapsulating bioactive proteins or cells in 3D printed scaffolds.²⁴ Type 13-93 bioactive glass (13-93 BG) was used as the inorganic component to fabricate the composite scaffolds for bone repair. 13-93 BG is modified from the 45S5 composition, but contains a higher SiO_2 content and additional network modifiers, such as K_2O and MgO . It has been stated that 13-93 BG is an ideal biomaterial for bone regeneration, because it has the same osteoconductivity as 45S5 BG, but a higher mechanical strength and faster biodegradation ratio.^{25,26} However, the appropriate conditions for fabricating 13-93 BG scaffolds with controlled morphology and suitable mechanical properties for bone regeneration require investigation. Rahaman and collaborators successfully prepared 13-93 BG scaffolds with high strength and porosity for bone regeneration by slip casting, freeze extrusion fabrication and unidirectional freezing of camphene-based suspensions, and further testified the excellent performance of 13-93 BG in bone repair.^{27–29} However, BG scaffold preparing methods are inconvenient, requiring other organic solvents, additional sintering procedures, or cool temperatures. Furthermore, the consistency and repeatability of the pore architectures in 13-93 BG scaffolds prepared by these methods needs improving. Therefore, taking advantage of the sol–gel transition of alginate induced by multivalent cations to stabilize 3D printing could be an alternative method for fabricating 13-93 BG/alginate composite scaffolds under mild conditions.

These organic–inorganic composite scaffold fabricated under mild conditions combined both the characteristics of 13-93 BG and alginate with the advantage of 3D printing to control pore size and individual architecture. 13-93 BG/alginate composite scaffolds were fabricated with mass ratios of 0 : 4 (0%), 1 : 4 (20%), 2 : 4 (33.3%) and 4 : 4 (50%) using 3D printing under mild conditions with the same geometry and plotting parameters. Their structures, mechanical properties, *in vitro* apatite mineralization abilities and cell response to rat bone mesenchymal stem cells (rBMSCs) were further investigated.

Materials and methods

Synthesis and characterization of 13-93 BG powders

According to a previous study,²⁷ 13-93 BG was prepared by melting the required quantities of SiO_2 , CaCO_3 , Na_2CO_3 , K_2CO_3 , MgCO_3 and NaH_2PO_4 (analytical grade, Sinopharm Chemical Reagent Co., Ltd., China) in a platinum/rhodium crucible in air for 2 h at 1300 °C. The melt was stirred twice at the isothermal temperature and quenched between two steel plates. Each glass powder was crushed with a hardened steel mortar and pestle, and then sieved through a stainless-steel sieve to give particle sizes of $<40\ \mu\text{m}$. The 13-93 BG composition (in mol%) was $6\text{Na}_2\text{O}\cdot 8\text{K}_2\text{O}\cdot 8\text{MgO}\cdot 22\text{CaO}\cdot 54\text{SiO}_2\cdot 2\text{P}_2\text{O}_5$. X-ray diffraction (XRD, Bruker, Germany) and Fourier translation infrared spectroscopy (FTIR, Spectrum 2000, PerkinElmer, USA) were used to analyze the 13-93 BG powders, and their morphologies were characterized by scanning electron microscopy (SEM, Jeol JSM6510, Tokyo, Japan).

Fabrication of 13-93 BG/SA scaffolds by 3D printing

The scaffolds were 3D printed as shown in Fig. 1. Sodium alginate powders (SA, Mingyue Chemical Reagent, China) and 13-93 BG powders were mixed until homogeneous using 13-93 BG/SA mass ratios of 0 : 4 (0% 13-93 BG), 1 : 4 (20%), 2 : 4 (33.3%) and 4 : 4 (50%). Printable inks were then prepared by mixing 1 g of 13-93 BG/SA with 5 g of polyvinyl alcohol (PVA) in a 6 wt% solution (Sigma-Aldrich, $M_w = 13\ 000\text{--}23\ 000$). After stirring until homogeneous, the inks were loaded into a printing tube and the scaffolds were printed using a printing needle (406 μm diameter) on a Bioscaffolder 2.1 platform (GeSiM, Germany) with computer-assisted design (Fig. 1a). The composite inks were extruded at room temperature using dosing pressures ranging from 400 to 500 kPa at a constant dispensing speed of $25\ \text{mm s}^{-1}$. The preformed scaffolds (Fig. 1b) were soaked in 1 M CaCl_2 aqueous solution for 5 h to achieve crosslinking, and then washed with deionized water three times. Finally, the scaffolds were lyophilized for use in subsequent experiments (Fig. 1c).

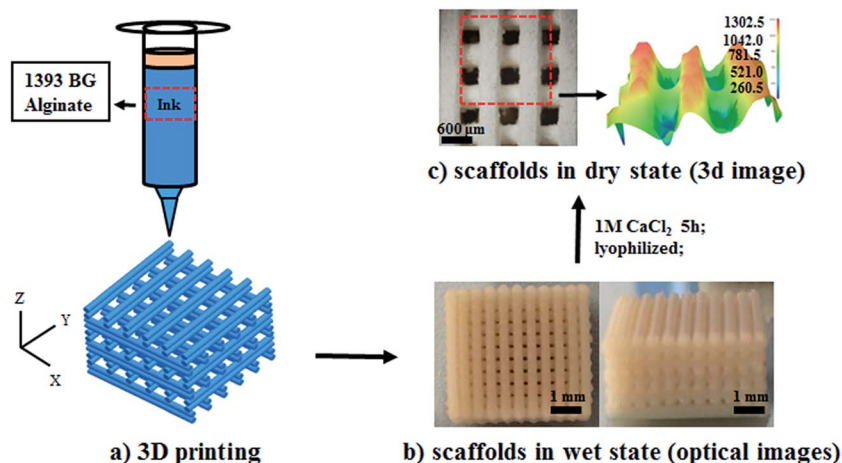


Fig. 1 Fabrication process for BG/SA composite scaffolds: (a) computer assisted design model and bio-inks; (b) optical images of scaffolds in a wet state; and (c) 3D micrographs of scaffolds in a dry state. (Print model: XXYY pattern, 406 μm , 400–500 kPa at a speed of $25\ \text{mm s}^{-1}$).



Characterization of scaffold morphology, porosity and shrinkage

The morphology and pore sizes of the 3D-printed 13-93 BG/SA scaffolds were characterized by SEM.

The scaffold porosity (ρ) was measured using a liquid replacement method.^{4,30} In brief, a scaffold was placed in a graduated cylinder containing dimethyl formamide (DMF) of a recorded volume (v_1). The cylinder was then placed under vacuum to force DMF into the scaffold pores. When no more bubbles appeared, the final volume was recorded as v_2 . Finally, the scaffold was removed and the volume of remaining DMF was recorded as v_3 . The porosity, ρ , of the scaffold was calculated according to eqn (1):

$$\rho = \frac{v_1 - v_3}{v_2 - v_3} \times 100\% \quad (1)$$

Shrinkage was calculated by measuring the scaffold length and height before and after drying. The values of ρ and shrinkage were averages from five samples.

Mechanical properties of 13-93 BG/SA scaffolds

The compressive strengths of the 13-93 BG/SA scaffolds ($6 \times 6 \times 6$ mm) were measured using an electromechanical universal testing machine (CMT6000, China) at a crosshead speed of 0.5 mm min^{-1} . A compressive deformation of 70% was achieved for all scaffolds at room temperature. The experiment was repeated using five samples.

In vitro apatite mineralization and ionic release of 13-93 BG/SA scaffolds

The apatite mineralization abilities of the 13-93 BG/SA scaffolds were investigated by soaking scaffolds in simulated body fluid (SBF), using 200 mL g^{-1} of scaffold mass, at 37°C for 0 and 10 days. Three samples were used as replicates. The morphologies and elemental compositions of the scaffold apatite mineralization layers were measured by SEM and energy-dispersive spectroscopy (EDS).

Ions released by the 13-93 BG/SA scaffolds were studied by soaking the scaffolds in 1.5 mL PBS at 37°C for 1, 3, 5 and 7 days. At predetermined time points, 1 mL aliquots of the release solution were taken and centrifuged at $10\,000 \text{ rpm}$ for 3 min, with 1 mL of fresh PBS solution added to the scaffolds for further incubation. The solution pH and the release of calcium, magnesium and silicate ions were measured using a pH meter and inductively coupled plasma atomic emission spectroscopy (ICP-OES, Optima 7100 DV, PerkinElmer Co. USA), respectively.

Seeding and cultivation of rBMSCs on scaffolds

Primary rat bone mesenchymal stem cells (rBMSCs) were obtained from femurs and tibiae of Sprague-Dawley (SD) rats ($140\text{--}160 \text{ g}$) according to a known method,³¹ and cells at passage 8 were used for experiments. The cell experiment protocols were approved by Shenzhen Institute of Advanced Technology, Chinese Academy of Science. All experiments were performed in

compliance with the Regulations for the Administration of Affairs Concerning Experimental Animals in China.

rBMSCs were cultivated in Dulbecco's modified Eagle's medium (DMEM, low glucose) containing 10% fetal calf serum, 10 U mL^{-1} penicillin and $100 \mu\text{g mL}^{-1}$ streptomycin at 37°C under 5% CO_2 in air. Before cell seeding, the cell culture scaffolds ($5 \times 5 \times 1.56 \text{ mm}^3$) were immersed in 75% ethanol for 6 h, washed three times with PBS and then immersed in culture medium without fetal calf serum for 12 h. After removing the culture medium, the scaffolds were transferred to 24-well plates and 2×10^4 cells were seeded onto each sample. The medium was renewed every 2 days.

Viability and morphology of rBMSCs on scaffolds

After cell culture for 1, 3 and 5 days, the proliferation of rBMSCs seeded on the scaffolds was evaluated using cell counting kit-8 (CCK-8, Dojindo).³² Absorbance was measured at 450 nm using a microplate reader. All experiments were performed in triplicate.

Morphologies of rBMSCs on the scaffolds were determined by fluorescence microscopy (Olympus, Tokyo) after 36 h of culture. In this experiment, the rBMSCs were stained with $2 \mu\text{g mL}^{-1}$ fluorescent dye (5/6-(*N*-succinimidylloxycarbonyl)-3',6'-*O*,*O*-diacetyl fluorescein) before use.³³

Alkaline phosphatase (ALP) activity assay

rBMSCs were seeded on the scaffolds under culture conditions in growth medium (DMEM, 10% FBS). ALP activity was measured after 7 and 14 days of culture, as previously described.² Briefly, the cells were washed three times with PBS and disrupted at 4°C for 30 min using cell lysis buffer. The cell lysis buffer was then collected by centrifugation and stored at -20°C . The ALP activity of the cell lysis buffer was determined using an alkaline phosphatase kit (Beyotime Institute of Biotechnology, Shanghai, China). The total protein content of the cell lysis buffer was determined using a BCA protein assay kit (Pierce Chemical Co., Rockford, USA), read at 562 nm and calculated according to a series of bovine serum albumin (BSA) standards. ALP activity was expressed as absorbance at 405 nm (OD value) per milligram of total cellular proteins. Cell culture plates were used as the control. All experiments were performed in triplicate.

Quantitative real-time polymerase chain reaction (qRT-PCR)

The osteogenic differentiation of rBMSCs seeded on scaffolds (0 : 4 and 2 : 4 groups) was further assessed by qRT-PCR after coculture for 7 and 14 days. The relative mRNA expression levels of commonly used bone markers, including osteopontin (OPN), osteocalcin (OCN), type I collagen (Col-I), and alkaline phosphatase (ALP) (primer pairs used are shown in Table 2), were measured. Total RNA isolation for qRT-PCR samples was conducted using an RNAsimple Total RNA Kit (Tiangen, Beijing, China). Complementary DNA (cDNA) was then reverse-transcribed from $1 \mu\text{g}$ of total RNA using a RevertAid First Strand cDNA Synthesis Kit (ThermoFisher Scientific, Massachusetts, USA) according to manufacturer instructions. Real-time PCR was performed on SYBR Green PCR Master Mix



(Toyobo Life Science, Osaka, Japan) and the reaction was carried out using an ABI 7500 system (Applied Biosystems, Massachusetts, USA). Finally, housekeeping gene glyceraldehyde-3-phosphate dehydrogenase (GAPDH) was used to normalize the relative mRNA expression level of each gene and quantification was based on the cycle threshold (CT) values.

Statistical analysis

For all experiments, the data were expressed as mean \pm standard deviation (SD). One-way ANOVA was performed to analyze variables, with p -values < 0.05 considered statistically significant differences.

Results

Characterization of 13-93 BG powders

The 13-93 BG powders ($6\text{Na}_2\text{O} \cdot 8\text{K}_2\text{O} \cdot 8\text{MgO} \cdot 22\text{CaO} \cdot 54\text{SiO}_2 \cdot 2\text{P}_2\text{O}_5$) were successfully prepared using appropriate quantities of the raw materials. The structure and morphology of the 13-93 BG powder are shown in Fig. 2. The FT-IR spectrum (Fig. 2a) showed distinctive absorption bands at $490\text{--}500\text{ cm}^{-1}$ (symmetric stretching, $\nu_{\text{sym}}(\text{Si-O-Si})$), $1000\text{--}1300\text{ cm}^{-1}$ (asymmetric stretching, $\nu_{\text{asym}}(\text{Si-O-Si})$), 750 cm^{-1} (bending mode $\delta(\text{Si-O-Si})$) and 930 cm^{-1} (SiO_{NBO} shoulder). The XRD pattern of the as-prepared glass (Fig. 2b) showed a band centered at $\sim 30^\circ$ in the 2θ range that was typical of amorphous glass. Fig. 2c shows the morphology of 13-93 BG powder, which had particle sizes of less than $40\text{ }\mu\text{m}$, which facilitated extrusion in 3D printing.

Fabrication and characterization of 3D printed 13-93 BG/SA scaffolds

Highly concentrated SA, with or without incorporating 13-93 BG, was used to prepare printing pastes to fabricate scaffolds, as shown in Fig. 1. The prepared 13-93 BG/SA composites had mass ratios of 0 : 4 (0% 13-93 BG), 1 : 4 (20%), 2 : 4 (33.3%) and 4 : 4 (50%), and the obtained scaffolds were denoted as pure SA, BG/SA 1 : 4, BG/SA 2 : 4 and BG/SA 4 : 4, respectively. All scaffolds were constructed in air and cured by incubation in CaCl_2 solution for 5 h at room temperature. The structures and

morphologies of the 3D printed 13-93 BG/SA scaffolds are presented in Fig. 3 and their parameters summarized in Table 1. Top-view and side-view images (Fig. 3a) showed that all printed 13-93 BG/SA scaffolds were completely interconnected and that macropore sizes were better maintained by higher BG contents. High magnification SEM images of the BG/SA 2 : 4 scaffold (Fig. 3b) not only confirmed a macropore size of around $500\text{ }\mu\text{m}$, but also revealed that there were plenty of micropores in the scaffold strands, due to the dissolution of incorporated PVA during scaffold incubation in CaCl_2 solution. Moreover, the porosity of 13-93 BG/SA scaffolds (Fig. 3c) was improved from 65.5 ± 5.3 to $87.5 \pm 3.9\%$ as 13-93 BG content in the scaffolds increased from 0 : 4 to 4 : 4. Fig. 3d depicts the shrinkage ratios in three directions (X , Y and Z) of all 13-93 BG/SA scaffolds. After lyophilization, the pure SA scaffolds demonstrated the most shrinkage, around 20% in all directions, while the shrinkage ratios decreased markedly to around 10% (BG/SA 4 : 4) as the 13-93 BG content increased.

Mechanical properties of 3D printed 13-93 BG/SA scaffolds

The compressive strength and modulus of the 3D printed 13-93 BG/SA scaffolds in a dry state were determined, as shown in Fig. 4, and used to calculate 70% deformation. The compressive strength and modulus initially increased, and then decreased, with increasing 13-93 BG content. Pure SA scaffolds had the weakest mechanical properties, with a compressive strength of $9.08 \pm 0.58\text{ MPa}$ and modulus of $42.62 \pm 4.05\text{ MPa}$, while BG/SA 2 : 4 had the most impressive mechanical properties, with a compressive strength of $16.74 \pm 1.78\text{ MPa}$ and modulus of $79.49 \pm 7.38\text{ MPa}$. However, in BG/SA 4 : 4 scaffolds the compressive strength and modulus sharply declined to $10.94 \pm 1.46\text{ MPa}$ and $53.07 \pm 3.33\text{ MPa}$, respectively.

In vitro mineralization and ionic release of the 3D printed 13-93 BG/SA scaffolds

The *in vitro* apatite mineralization of an implanted material is usually related to its osteoinductive ability for bone regeneration.^{9,29} In this study, the pure SA scaffolds had smooth surfaces without apatite deposition before or after soaking for 10 days in SBF, while incorporation of 13-93 BG endowed the 13-93 BG/SA

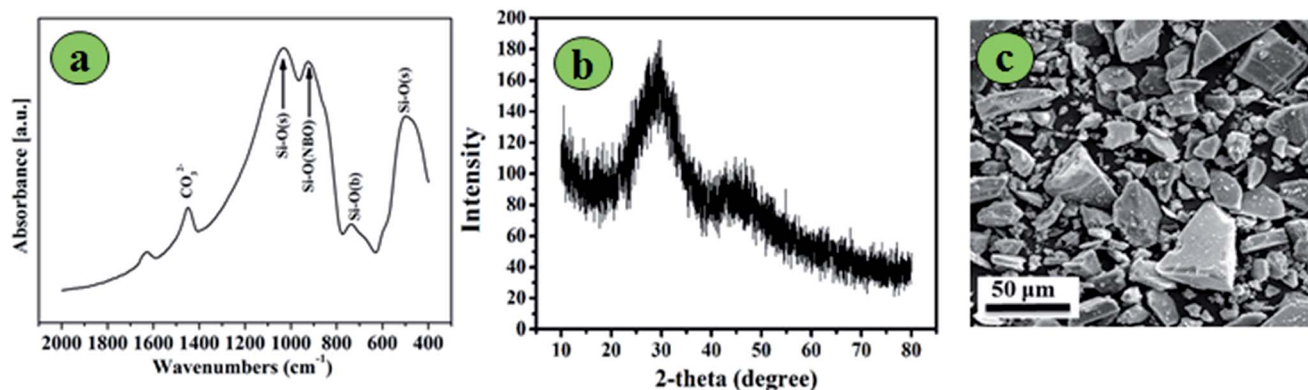


Fig. 2 Characterization of 13-93 BG powders: (a) FT-IR spectrum; (b) XRD pattern; and (c) morphology and size ($<40\text{ }\mu\text{m}$).



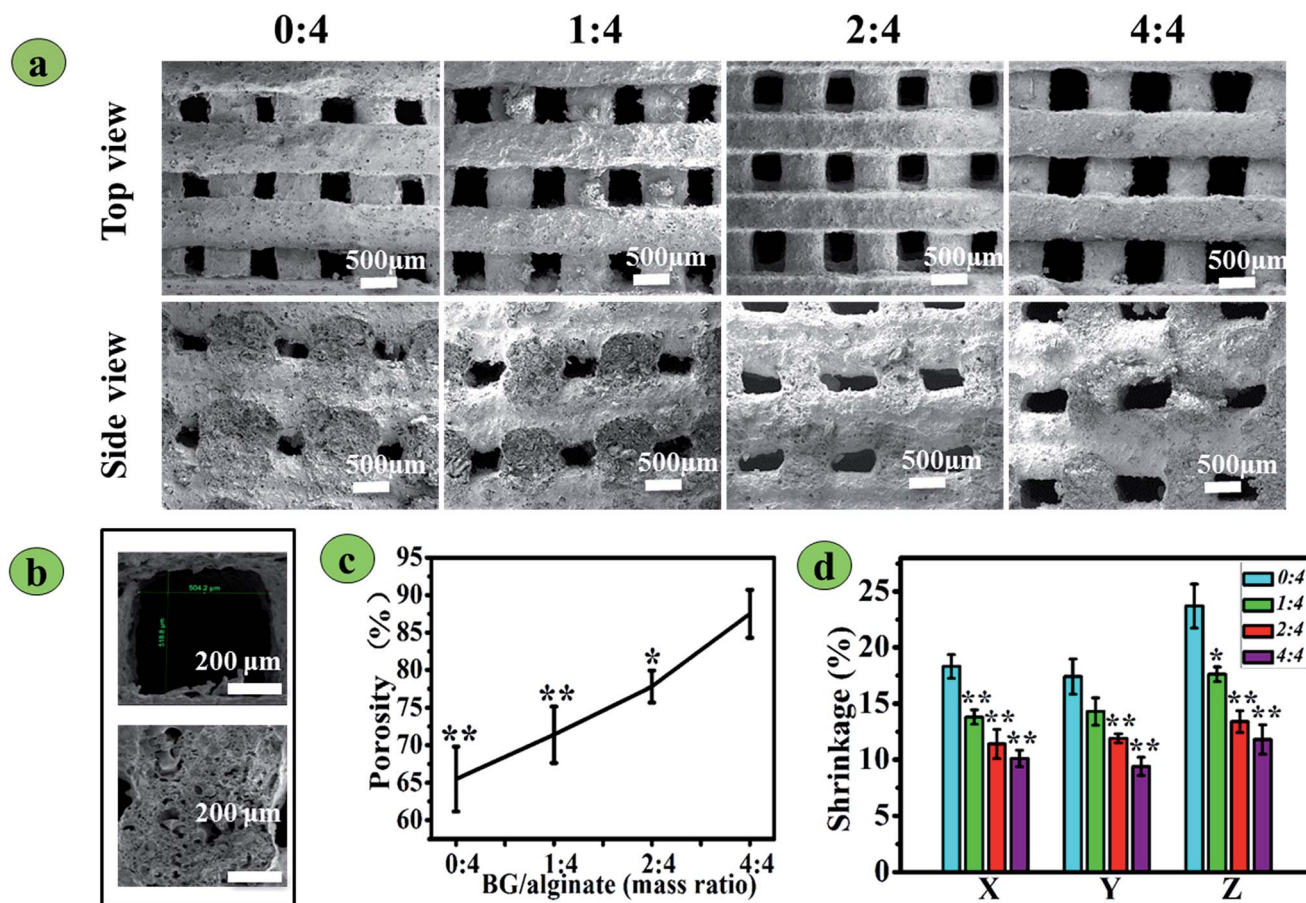


Fig. 3 Morphology, porosity and shrinkage of 3D-printed BG/SA composite scaffolds: (a) top and side-view SEM images of scaffolds. BG/SA mass ratios were 0 : 4, 1 : 4, 2 : 4, and 4 : 4; (b) high-magnification SEM images showing a pore and a vertical section of strands of the BG/SA 2 : 4 scaffold; (c) porosity; (d) scaffold shrinkage (%) in X, Y and Z directions. Scale bar is 500 μm . (*Denotes $P < 0.05$, compared with 13-93 BG/SA 0 : 4; **denotes $P < 0.01$, compared with 13-93 BG/SA 0 : 4).

Table 1 Parameters of 3D printed BG/SA composite scaffolds

| Scaffold code | 13-93 BG/SA (mass ratio) | Shrinkage (%) | | | Pore size top view (μm) | Pore size side view (μm) | Porosity (%) | Compressive strength (MPa) | Modulus (MPa) |
|---------------|--------------------------|----------------|----------------|----------------|--------------------------------------|---------------------------------------|----------------|----------------------------|------------------|
| | | X | Y | Z | | | | | |
| SA | 0 : 4 | 18.3 \pm 1.3 | 17.4 \pm 1.9 | 23.7 \pm 2.4 | 340 \pm 27 | 250 \pm 10 | 65.5 \pm 5.3 | 9.08 \pm 0.58 | 42.62 \pm 4.05 |
| BG/SA 1 : 4 | 1 : 4 | 13.8 \pm 0.8 | 14.3 \pm 1.5 | 17.6 \pm 0.8 | 460 \pm 17 | 310 \pm 12 | 71.4 \pm 4.6 | 10.87 \pm 1.10 | 52.49 \pm 7.03 |
| BG/SA 2 : 4 | 2 : 4 | 11.4 \pm 1.6 | 11.9 \pm 0.5 | 13.4 \pm 1.2 | 500 \pm 24 | 360 \pm 18 | 77.8 \pm 2.6 | 16.74 \pm 1.78 | 79.49 \pm 7.38 |
| BG/SA 4 : 4 | 4 : 4 | 10.1 \pm 0.9 | 9.4 \pm 1.0 | 11.8 \pm 1.6 | 530 \pm 27 | 400 \pm 26 | 87.5 \pm 3.9 | 10.94 \pm 1.46 | 53.07 \pm 3.33 |

Table 2 Primer pairs used in real-time PCR analysis

| Gene | Forward primer (5'-3') | Reverse primer (5'-3') | Accession no. |
|-------|------------------------|------------------------|---------------|
| GAPDH | GGCACAGTCAAGGCTGAGAATG | ATGGTGGTGAAGACGCCAGTA | NM_017008.4 |
| OPN | CCAAGCGTGGAACACACAGCC | GGCTTTGGAAGTCCGCTGACTG | M14656.1 |
| OCN | GGTGCAGACCTAGCAGACACCA | AGGTAGCGCCGGAGTCTATCA | J04500 |
| COL-I | GCCTCCCAGAACATCACCTA | GCAGGGACTTCTTGAGGTTG | NM_053304 |
| ALP | AACGTGGCCAAGAATCATCA | TGTCCATCTCCAGCCGTGTC | NM_013059 |

scaffolds with excellent apatite mineralization ability in SBF (Fig. 5a). After soaking for 10 days in PBS, alveolate-like apatite crystals formed on the surface of BG/SA 1 : 4, BG/SA 2 : 4 and

BG/SA 4 : 4. Newly formed apatite on the surface of the 13-93 BG/SA scaffolds was further confirmed by EDS analysis, as shown in Fig. 5b, with characteristic peaks observed for Ca, Mg,



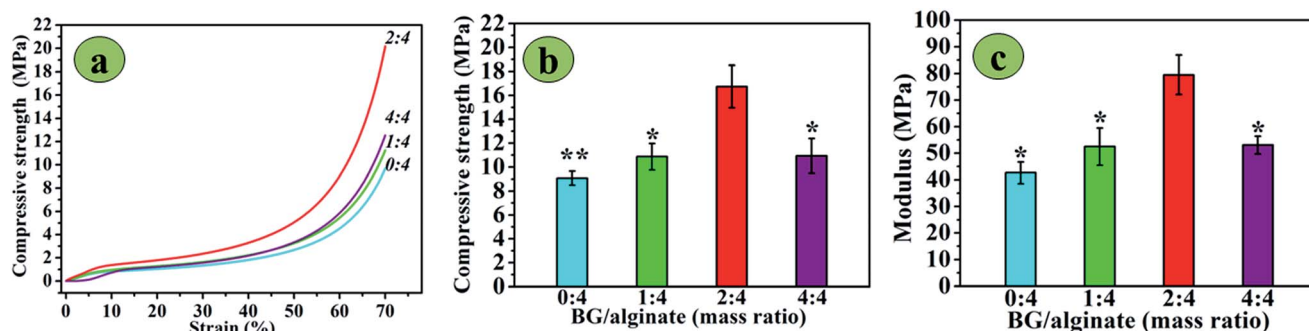


Fig. 4 Mechanical properties of 3D-printed BG/SA composite scaffolds: (a) compressive stress–strain curves; (b) compressive strength; and (c) modulus. (*Denotes $P < 0.05$ compared with BG/SA 2 : 4; **denotes $P < 0.01$ compared with BG/SA 2 : 4, $n = 4$).

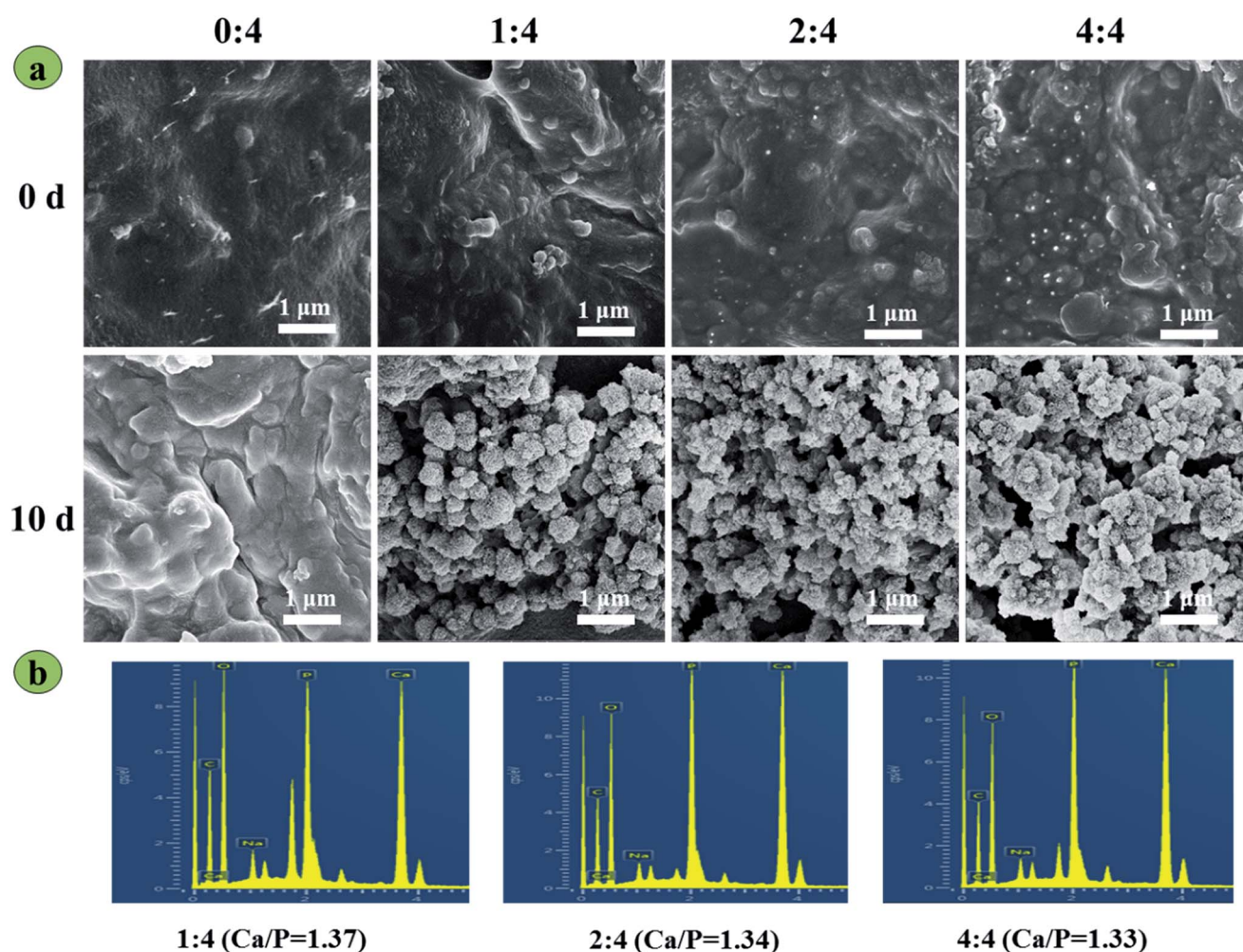


Fig. 5 *In vitro* apatite mineralization of 3D-printed BG/SA composite scaffolds: (a) surface morphologies of scaffolds before and after soaking in SBF for 10 days; (b) EDS analyses of apatite layers. Scale bar is 1 μm .

Si and P. Moreover, the Ca/P ratios in BG/SA 1 : 4, BG/SA 2 : 4 and BG/SA 4 : 4 were 1.37, 1.34 and 1.33, respectively. All Ca/P ratios were less than that of hydroxyapatite (1.67), which might be attributed to the presence of Mg^{2+} replacing Ca^{2+} in the mineralized apatite layer.

The pH values of PBS solutions containing four types of scaffolds were measured over 7 days, as shown in Fig. 6a. A pH

variation of solutions containing BG/SA scaffold rapidly increased over 7 days, while the pH of that containing pure SA scaffold increased slowly. After immersing the scaffolds for 3 days, the pH values of each BG/SA scaffold were higher than that of the pure SA scaffold, with the pH value higher with increasing 13-93 BG content. In BG/SA 4 : 4, the pH value was greater than 8.5 after immersion for 7 days in PBS. Meanwhile, the



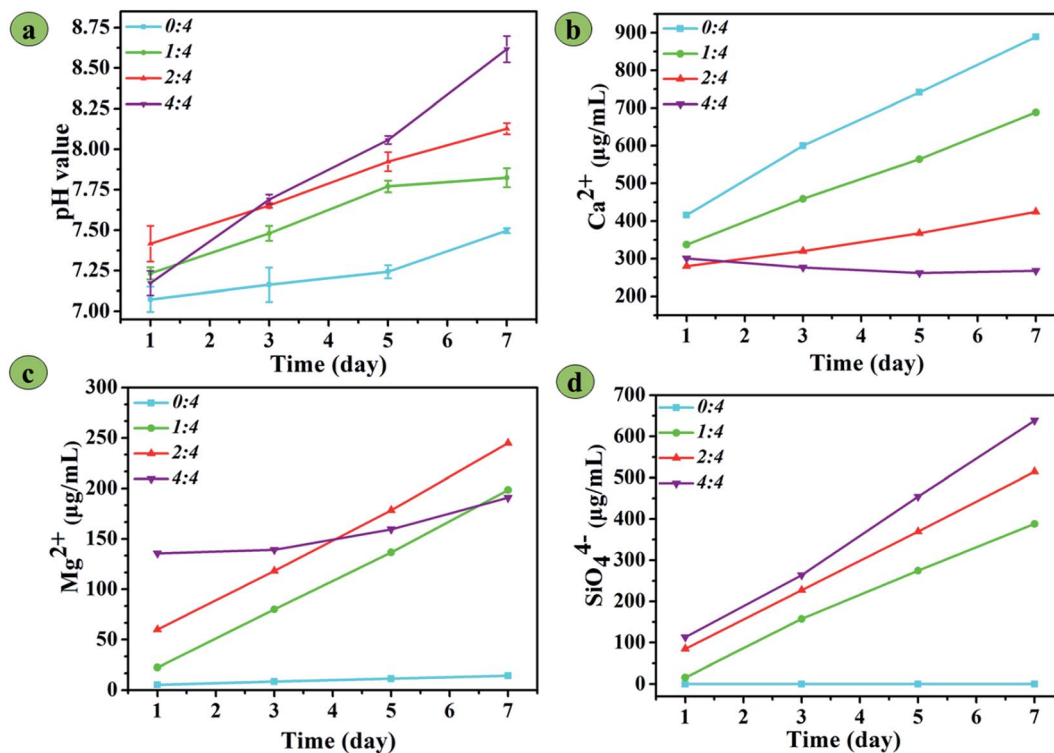


Fig. 6 pH value and ion release of 3D-printed BG/SA composite scaffolds in PBS solution at 1, 3, 5 and 7 days: (a) pH variation; (b) Ca²⁺ release; (c) Mg²⁺ release; and (d) SiO₄⁴⁻ release.

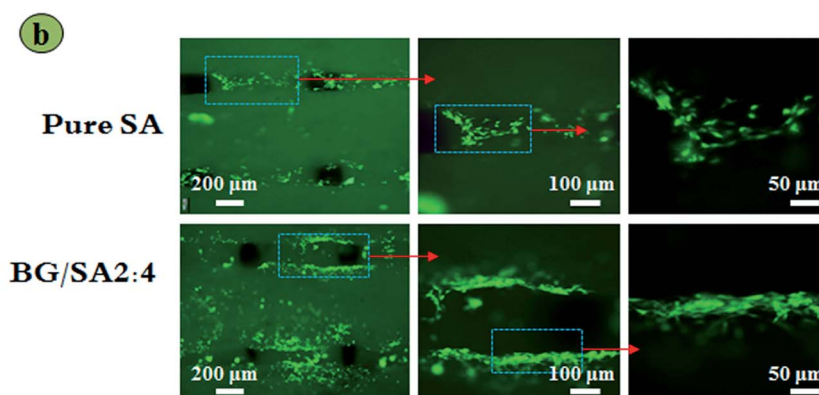
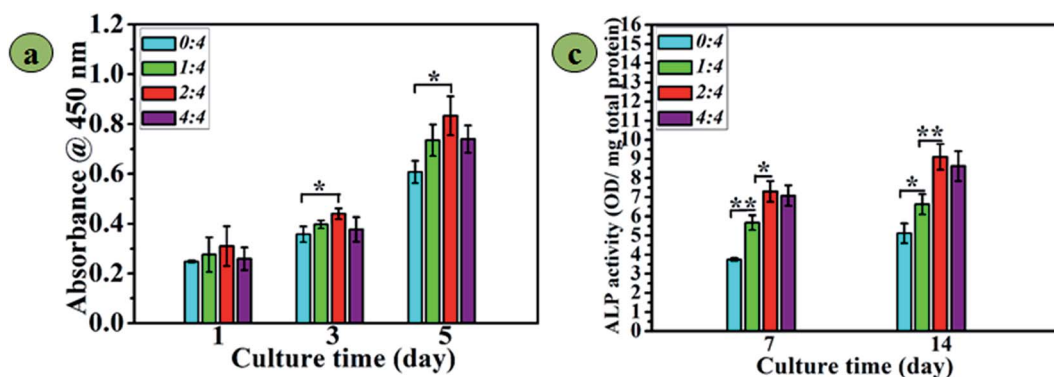


Fig. 7 rBMSC response on 3D-printed BG/SA composite scaffolds: (a) proliferation determined using CCK-8 at 1, 3 and 5 days; (b) attachment on pure SA and BG/SA 2 : 4 scaffolds determined by fluorescence images of cells after 36 h of culture; and (c) osteogenic differentiation of rBMSC seeded on scaffolds at 7 and 14 days. ($n = 3$, mean \pm SD, *denotes $p < 0.05$, **denotes $p < 0.01$).



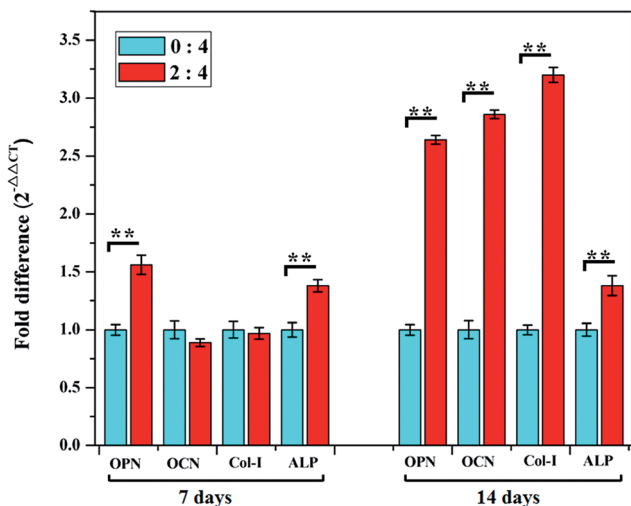


Fig. 8 Osteogenic gene expression (OPN, OCN, Col-I and ALP) of rBMSCs cultured on pure SA and BG/SA 2 : 4 scaffolds at 7 and 14 days of culture. (*Denotes $p < 0.05$, **denotes $p < 0.01$).

concentrations of Ca^{2+} , Mg^{2+} and SiO_4^{4-} ions in PBS after soaking the scaffolds were detected by ICP over 7 days. Fig. 6b shows that the concentration of Ca^{2+} ions from pure SA, BG/SA 1 : 4 and BG/SA 2 : 4 in PBS increased, but the concentration of Ca^{2+} ions from BG/SA 4 : 4 in PBS slowly declined. The Ca^{2+} concentration could be increased by ion exchange from the alginate strands and the release of 13-93 BG, but decreased by apatite mineralization. Therefore, Ca^{2+} release from the alginate strands motivated the sharp increase in Ca^{2+} ions from the pure SA scaffold in PBS. In contrast, BG/SA 4 : 4, the scaffold with the highest BG content, gave the highest pH value in PBS, which stimulated the spontaneous formation of apatite mineralization, causing a sharp decrease in Ca^{2+} concentration. The release of Mg^{2+} and SiO_4^{4-} ions (Fig. 6c and d) was attributed to 13-93 BG incorporation, resulting in improved Mg^{2+} and SiO_4^{4-} concentrations with increasing 13-93 BG content, except for Mg^{2+} release from BG/SA 4 : 4. This exception was due to Mg^{2+} ions, in a similar fashion to Ca^{2+} ions, precipitating under the highly alkaline conditions provided by BG/SA 4 : 4.

Cultivation of rBMSCs on 3D printed 13-93 BG/SA scaffolds

To evaluate the biological effects of the 3D printed 13-93 BG/SA scaffolds for bone regeneration, the cell morphology, proliferation, ALP activity and osteogenic activity of rBMSCs seeding on the scaffolds were systematically investigated. The proliferation of rBMSCs on different scaffolds is depicted in Fig. 7a. The cell number increased on all scaffolds during the 5 day culture, and rBMSCs on BG/SA scaffolds proliferated better than those on pure SA scaffolds. However, when the BG content was increased to 50% (4 : 4), the proliferation of rBMSCs declined. This was further confirmed by fluorescence images (Fig. 7b), which showed that rBMSCs preferred to spread and adhere on BG/SA 2 : 4 scaffolds than pure SA scaffolds.

Determination of the ALP activity of rBMSCs (Fig. 7c) indicated that 13-93 BG incorporation stimulated osteogenic

differentiation of rBMSC. After 7 and 14 days of culture, the ALP activity of rBMSCs cultured on BG/SA composite scaffolds was significantly higher than that on pure alginate scaffolds ($p < 0.05$). However, the ALP activities of rBMSCs on BG/SA 4 : 4, in accordance with the proliferation results, were slightly weaker than those on BG/SA 2 : 4 scaffolds, but better than those on pure SA scaffolds.

Fig. 8 summarizes the osteogenesis-related gene expressions of OPN, OCN, Col-I and ALP at 7 and 14 days of incubation with rBMSCs on BG/SA 2 : 4 composite and pure SA scaffolds. At 7 days, OPN and ALP of rBMSCs showed higher expression on BG/SA 2 : 4 scaffolds than on pure SA scaffolds, while OCN and Col-I expression on pure SA scaffolds and BG/SA 2 : 4 scaffolds were similar. Furthermore, at 14 days, the expression of OPN, OCN, Col-I and ALP on BG/SA 2 : 4 scaffolds were significantly higher than those on pure SA scaffolds. These results were likely due to incorporation of 13-93 BG onto BG/SA scaffolds, which could promote osteogenic differentiation of rBMSCs.

Discussion

In the present study, biodegradable 13-93 BG/SA composite scaffolds were successfully fabricated by 3D printing under mild conditions with an XYY pattern. Using SA components as the organic matrix, gelatinized by crosslinking with Ca^{2+} to form the steady scaffolds, is a popular stabilization method for mild 3D printing, while 13-93 BG particles with diameters less than 40 μm were used as the inorganic component (Fig. 2), with the intention of improving the mechanical properties and bioactivity of the SA scaffold. Compared with previous methods,²⁶⁻²⁹ including slip casting, polymer foam replication and freeze extrusion, this was a more facile method for preparing 13-93 BG scaffolds for bone regeneration. Meanwhile, the incorporation of 13-93 BG into SA scaffolds significantly improved the mechanical strength, apatite mineralization ability, and attachment and osteogenic differentiation of rBMSC.

Bioactive scaffolds with interconnected pores, suitable porosity and tailored mechanical strength are needed for bone tissue engineering.^{30,34} In particular, the 13-93 BG/SA composite scaffolds obtained herein were observed to have hierarchical macropores (around 500 μm , Fig. 3a) and plenty of micropores on the strands (Fig. 3b), suggesting completely interconnected pores, which have been reported as suitable for bone-related cell ingrowth and nutrient delivery.³⁴ Meanwhile, increasing 13-93 BG content improved porosity and decreased shrinkage ratios in three directions in the 3D printed 13-93 BG/SA composite scaffolds (Fig. 3c and d), further indicating that 13-93 BG dispersed in the SA matrix helped maintain composite scaffold stability. Furthermore, the 13-93 BG content was found to significantly regulate the mechanical properties of the 13-93 BG/SA composite scaffolds (Fig. 4), giving a maximum compressive strength and modulus of 16.74 ± 1.78 MPa and 79.49 ± 7.38 MPa, respectively, in 13-93 BG/SA 2 : 4. On one hand, the 13-93 BG particles acted as reinforcing agents in composites when the 13-93 BG content increased from 0 to 33.3%. On the other hand, when the 13-93 BG content became more dominant in the composite scaffolds, as in BG/SA 4 : 4,



they exhibited the brittleness of 13-93 BG. Accordingly, 13-93 BG/SA composite scaffolds with porosities of 65–87%, compressive strengths of 10–16 MPa and moduli of 40–80 Mpa met the mechanical demands of human cancellous bone.²⁹

In addition to sufficient mechanical properties, excellent cytocompatibility and sound bioactivity are also indispensable. *In vitro* apatite mineralization of an implanted material in SBF is known as the initial evaluation of osteoinductive ability for bone regeneration.^{2,33,34} Our study showed that 13-93 BG incorporation significantly improved the apatite forming ability of SA scaffolds in SBF (Fig. 5). The improved apatite-mineralization ability of SA scaffolds might benefit the rBMSC response to scaffolds because improved cell attachment and higher specific ALP activity were observed for 13-93 BG/SA compared with pure SA scaffolds. 13-93 BG incorporation also led to the release of Mg^{2+} and SiO_4^{4-} ions (Fig. 6), which were generally recognized as enhancers of bone regeneration. Previous investigations have confirmed that Mg^{2+} and SiO_4^{4-} ions not only improve cell attachment and proliferation, but also simulate osteogenic differentiation of rBMSCs.^{35,36} In addition, environment pH also plays an important role in cell growth, proliferation and mineralization. For bone regeneration, it has been reported that a relatively high local pH is necessary during the bone formation process because the optimum ALP occurs at pH 8.5, in comparison with the nominal physiological value of 7.4.³⁷ Furthermore, 13-93 BG was also shown to degrade the release of alkaline ions, which would exchange with hydrogen ions (H^+) in water to generate a weakly alkaline pH. This pH would promote the nucleation of calcium phosphates and facilitate the proliferation of osteoblasts. The results of this study showed that 13-93 BG incorporation into scaffolds immersed in PBS increased the environment pH (Fig. 6). Therefore, the rBMSC activities, comprising attachment (Fig. 7), proliferation (Fig. 7) and mineralization (Fig. 8), on 13-93 BG/SA 2 : 4 were the highest, and were simultaneously regulated by three factors, namely apatite mineralization, the release of bioactive Mg^{2+} and SiO_4^{4-} ions, and the environment pH.

Conclusions

In this study, we fully utilized the crosslinking property of SA to fabricate 13-93 BG/SA scaffolds by 3D printing under mild conditions. All the obtained scaffolds were completely interconnected, with pore sizes, porosities and shrinkage ratios that were regulated by the amount of 13-93 BG incorporated. Moreover, in comparison with the pure SA scaffold, 13-93 BG incorporation significantly improved the mechanical properties, increased apatite mineralization and released bioactive Mg^{2+} and SiO_4^{4-} ions. Meanwhile, rBMSC preferred to spread, adhere and osteogenically differentiate on the 13-93 BG/SA scaffolds, especially BG/SA 2 : 4. Therefore, the 3D printed 13-93 BG/SA scaffolds, a novel biodegradable organic–inorganic composite scaffold with suitable porosity, tailored mechanical strength and sound bioactivity, could be a potential candidate for bone regeneration.

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