Controlled release of TGF-beta 1 from RADA self-assembling peptide hydrogel scaffolds

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Abstract: Bioactive mediators, cytokines, and chemokines have an important role in regulating and optimizing the synergistic action of materials, cells, and cellular microenvironments for tissue engineering. RADA self-assembling peptide hydrogels have been proved to have an excellent ability to promote cell proliferation, wound healing, tissue repair, and drug delivery. Here, we report that D-RADA16 and L-RADA16-RGD self-assembling peptides can form stable second structure and hydrogel scaffolds, affording the slow release of growth factor (transforming growth factor cytokine-beta 1 [TGF-beta 1]). In vitro tests demonstrated that the plateau release amount can be obtained till 72 hours. Moreover, L-RADA16, D-RADA16, and L-RADA16-RGD self-assembling peptide hydrogels containing TGF-beta 1 were used for 3D cell culture of bone mesenchymal stem cells of rats for 2 weeks. The results revealed that these three RADA16 peptide hydrogels had a significantly favorable influence on proliferation of bone mesenchymal stem cells and hold some promise in slow and sustained release of growth factor.

Keywords: molecular self-assembly, self-assembling peptide, hydrogel scaffolds, controlled release, TGF-beta 1, bone mesenchymal stem cells

Introduction

Ideal hydrogel scaffolds consisting of natural or synthetic materials are important for regenerative medicine and tissue engineering. Materials such as alginate, agarose, and collagen have been widely applied in medical and drug research. However, there are some problems such as poor biocompatibility, immune rejection, and degradation product toxicity. Therefore, it is crucial and urgent to develop a novel and ideal biomaterial hydrogel.¹⁻³

Recent studies have shown that self-assembling peptide nanofiber hydrogel scaffolds have been found to exert a significantly favorable influence on bone and cartilage regeneration, wound healing, as well as drug delivery.¹⁻⁵ These ionic self-complementary peptides can spontaneously self-assemble into beta sheet secondary structure and peptide nanofiber after stimulation with electrolyte solutions. In addition, these peptide nanofibers with a diameter of 10–20 nm can then form highly hydrous hydrogels with pore sizes ranging from 5 nm to 200 nm and water content >99.5% w/v.⁶⁻¹⁰ Self-assembling peptide hydrogels have been proved to have good biocompatibility, and they can completely degrade without any toxicity of degradation products.^{9,11,12} They also have the ability to deliver signal molecules and functional proteins.¹³⁻¹⁵ For example, RADA16 peptide hydrogel scaffolds afforded the controlled release of various functional proteins, including fibroblast growth factor, vascular endothelial growth factor, brain-derived neurotrophic factor, lysozyme, trypsin inhibitor, BSA, and IgG. Further studies have demonstrated that these released functional proteins can maintain their original protein structure and function based on conformation analysis

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and bioactivity detection.^{9,14} Many studies have demonstrated that self-assembling peptide hydrogel scaffolds have been widely applied for drug and protein delivery, tissue regeneration, and cell culture.^{1–5,9}

Previous studies have demonstrated that L-RADA16 peptide hydrogels can mimic cell microenvironment and promote cell proliferation, whereas D-form peptide hydrogels have the ability to induce cell growth and rapid hemostasis.^{10,16} In addition, peptide hydrogels could be functionalized with active motifs such as IKVAV and RGD, which benefited to regulate cell behaviors, including differentiation, attachment, and migration.^{17,18} This study focused on the secondary structure of D-RADA16 peptide and D-RADA16-RGD peptide hydrogel scaffold using circular dichroism spectroscopy, their microstructural feature using transmission electron microscopy (TEM) and rheometry assays, as well as the capability of promoting the proliferation of bone mesenchymal stem cells (BMSC) of rats compared with L-RADA16 peptide hydrogel scaffolds. It was known that transforming growth factor beta 1 (TGF-beta 1) cytokine can promote osteogenesis and chondrogenesis and can be used for cell function assays or blood vessel regeneration.^{19–21} Peptide hydrogel scaffolds containing TGF-beta 1 were used to explore their potential in controlled release of TGF-beta 1 for the functions of BMSC.

Materials and methods

Peptide synthesis and purification

D-RADA16 peptide sequence is Ac-RADARADARADA RADA-CONH₂ (containing all D-amino acids). L-RADA16 sequence containing all L-amino acids is Ac-RADARADARADARADA-CONH₂, and L-RADA16-RGD sequence containing all L-amino acids is Ac-RADARADARADARADARGDS-CONH₂. These peptides were commercially custom-synthesized by solid-phase peptide synthesis (Shanghai Biotech Bioscience and Technology Co, Ltd, Shanghai, People's Republic of China). The N-terminus and C-terminus of these peptides were acetylated and amidated, respectively, and the peptides were purified by high-performance liquid chromatography and characterized by mass spectroscopy. The final purity of the D-RADA16, L-RADA16, and L-RADA16-RGD was 97.78%, 97.45%, and 96.68%, respectively.

Circular dichroism spectroscopy

The sample was composed of RADA peptide (20 mg/mL) dissolved in NaCl solution, and the concentration was regulated to 50 μ M. The sample was prepared at 25°C and equilibrated for 30 seconds. A total of 500 μ L of sample solution was added in a circular dichroism colorimetric cuvette. Tests were conducted using a J-810 CD spectrometer (J-810;

JASCO Corporation, Tokyo, Japan). We measured the entire 190–310 nm wavelength range, and the wavelength ranging from 190 to 260 nm was used for the analysis.

Transmission electron microscopy

TEM peptide solution samples (5 mg/mL) were prepared at 25°C by dissolving lyophilized peptide powder in phosphate-buffered saline (PBS, pH 7.4). After 48 hours, a micropipette was used to load 5 mL of this peptide solution to a carbon-coated copper grid. The excess solution was removed by a piece of filter paper. Then, the peptide solution sample was dyed using 10 mL of uranyl acetate for 30 seconds and dried overnight in a desiccator, and then bright-field images were taken using a transmission electron microscope (Philips Tecnai G2 F20, Koninklijke Philips N.V., Amsterdam, the Netherlands) operating at 200 kV.

Rheometry

Rheology assays were carried out using a 1° stainless steel cone-controlled rheometer with a diameter of 20 mm (Thermo Fisher Scientific, Waltham, MA, USA). Samples were dissolved in PBS (pH 7.4) at a concentration of 10.0 mg/mL. The peptide solution was mixed with PBS (pH 7.4) at a volume ratio of 1:1 or 1:3 and stored at 4°C in a refrigerator over night, and 150–200 μ L of samples were used for analysis at 25°C.

Cell culture

The BMSC were obtained from the femurs of 4–5 weeks old female Sprague Dawley rats from the experimental animal center of Chongqing Medical University. Conventional reagents were used, and cells prepared for 2D cell culture and a modified 3D culture were grown in 10% fetal bovine serum (PAN Biotech UK Ltd., Dorset, UK). All cells were subcultured to third to fifth generation for follow-up experiments. Our study followed the principles of Medical Ethics and Declaration of Helsinki, and judged by the Ethics Committee of the First Affiliated Hospital of Chongqing Medical University. This experimental protocol was approved by the Institutional Animal Care and Use Committee of Chongqing Medical University of China, and Chinese national guidelines for the care and use of laboratory animals were applied.

TGF-beta 1 release

RADA self-assembling peptide hydrogels with or without BMSC were used to release TGF-beta 1 at 37°C. The TGF-beta 1 solution was mixed with 1% v/w RADA self-assembling peptide solutions (20 mg/mL), and the final mixed peptide solution consisted of 10 ng of TGF-beta 1 and 40 μ L of peptide solution. A total of 20 μ L of mixed solution was added

into a 24-multiwell plate (NEST Biotechnology Co. Ltd., Rahway, NJ, USA) and then was stimulated to self-assemble after adding 180 μ L of PBS (7.4 pH) in each well for at least 30 minutes. Empty wells filled with PBS were used as the control group. BMSC of adult Sprague Dawley rat were cultured in the 24-multiwell plate, and the concentration was 2×10^4 cells per well. Standard DMEM/F12 culture medium without growth factors was used for the test, and the dose was 500 μ L per well. Cell proliferation was assessed in 3 days, 7 days, and 14 days using a VersaMax microplate reader. Enzyme-linked immunosorbent assay (ELISA) kit (eBioscience, San Diego, CA, USA) was used to measure the amount of TGF. A total of 40 μ L of supernatant of the mixed solution in each well was taken away at the time point of 1 hour, 2 hours, 4 hours, 8 hours, 24 hours, 72 hours, and 168 hours, while the same dose of fresh PBS solution was added into each well slowly and gently. These supernatant samples were stored in a -80°C refrigerator for the following ELISA determination. The amounts of TGF-beta 1 in each group were then obtained and the diffusion coefficients were calculated according to the 1-D unsteady state form of Fick's second law.^{9,22}

Cell imaging and proliferation assay

We observed cells through an inverted microscope (Olympus Corporation, Tokyo, Japan) after coculturing for 2 weeks. CCK-8 assay was applied to investigate the effect of TGF-beta 1 released from RADA peptide hydrogel scaffolds on cell viability. CCK-8 kit (Dojindo Laboratories, Kumamoto, Japan) was added into the culture medium of wells in the ratio of 1:10, and then the 24-multiwell plate was incubated at 37°C for 1 hour. The absorbance was measured using a VersaMax microplate reader (Thermo Fisher Scientific) at an absorption wavelength of 450 nm. Data were expressed as arbitrary absorbance units.

Statistical analysis

SPSS software was used for statistical analysis. An unpaired *t*-test was used to compare the two groups. Comparisons between multiple groups were performed by one-way analysis of variance followed by least significant difference or Student–Newman–Keuls as post hoc tests. Data were expressed as mean $\pm$ standard error of the mean, and a *P*-value of <0.05 was defined as statistically significant.

Results

Secondary structure of L-RADA16-RGD

The β -sheet structure of L-RADA16-RGD is shown in Figure 1, indicating that L-RADA16-RGD peptide could

Figure 1 Circular dichroism spectra of the chiral peptides.

Notes: Peptide L-RADA16-RGD of 50 μ M has stable secondary structures in 20 mM NaCl at 25°C , 193.5 nm (13.5827), 193 nm (12.4508), and 198 nm (−15.8484).

Abbreviation: deg, degree.

produce stable secondary structures in 20 mM NaCl at 25°C . This result indicates that the 3D cell culture should be handled quickly at room temperature.

Nanostructures of RADA16 self-assembling peptide nanofiber scaffolds

TEM was applied to observe the microstructure of the RADA16 peptide hydrogels. TEM images showed that the RADA16 peptides were able to self-assemble into interwoven nanofibers at 25°C (Figure 2). These nanofibers were ~ 10 nm in diameter and formed pores <200 nm, whereas the diameter of nanofibers of L-RADA16-RGD was wider than that of the other peptides. The physical network and these structures coincided with the mechanism of interwoven formation in these peptide hydrogels.

Rheological tests of L-RADA16 and D-RADA16

These peptides can self-assemble into nanofibers and further hydrogelation occurs. The earliest few minutes of hydrogelation are crucial to keep cells in the truly 3D microenvironment. However, gelation nanomechanical properties in the cell media are poorly clear, especially in the initial phase. We chose typical ions (Na^+/K^+) in the cell media to investigate the gelation nanomechanical properties. Rheological experiments measure a storage modulus (G') and a loss modulus (G'') by evaluating elastic response (rigidity) and viscous response of material, respectively, by varying frequencies of applied oscillatory stress. Upon comparing the rheological property of D-RADA16 and L-RADA16 (1 mg/mL) self-assembly in water or salt solution (0.01 M PBS, Na^+/K^+), D-RADA16 peptide is more sensitive to

Figure 2 TEM images of microstructure of (A) L-RADA16, (B) D-RADA16, and (C) L-RADA16-RGD (200 nm).

Note: Magnification: (A) $\times 75,000$; (B) $\times 60,000$; and (C) $\times 75,000$.

Abbreviation: TEM, transmission electron microscopy.

ions than L-RADA16, and the frequency G' was greater than G'' (Figure 3A). High concentration of peptides was revealed to result in higher frequency G' (20.0 mg/mL), both D- and L-RADA16 > 12 Pa (Figure 3A and B), and D-form

peptide showed a little bit lower frequency G' (Figure 3B). High concentration of D-RADA16 and L-RADA16 demonstrated similar mechanical properties and Na^+/K^+ can affect them immediately. A reasonable explanation is that

Figure 3 Rheological assays: 2.5, 5.0 mg/mL peptide hydrogels in PBS (pH 7.4), 25°C.

Notes: (A) The rheological properties of 2.5 and 5.0 mg/mL L-RADA16 hydrogels in 10.0 mg/mL PBS (pH 7.4) at 25°C. (B) The rheological properties of 2.5 and 5.0 mg/mL D-RADA16 hydrogels in 10.0 mg/mL PBS (pH 7.4) at 25°C.

Abbreviation: PBS, phosphate-buffered saline.

Na^+/K^+ can induce the peptides to self-assemble into short nanofibers rapidly. Our result is useful in a process of 3D cell culture.

In vitro TGF-beta 1 release from three types of RADA16 hydrogels

The release experiments were observed for 14 days. The release profiles of TGF-beta 1 from the RADA peptide hydrogels are shown in Figures 4 and 5. Results revealed that ~18% (Figure 4) and 31% (Figure 5) of incorporated TGF were released from 1% w/v L-RADA16-RGD hydrogel within the first 24 hours. In contrast, only 17% (Figure 4) and 20% (Figure 5) of TGF-beta 1 were released from the D-RADA16 hydrogel within 24 hours. These results demonstrated that protein release rate in different types of peptide hydrogel was dissimilar. Compared with the D-RADA16 hydrogel, the initial burst effect of the L-RADA16-RGD hydrogel was higher.

Released TGF-beta 1 activity assays

TGF-beta 1 has been proven to have the ability to stimulate and promote proliferation of BMSC by adjusting mitotic process. To assess the activity of the TGF-beta 1 released from the RADA peptide hydrogel scaffolds CCK-8 assay was used (Figure 6), which indicated that there was enhanced proliferation of BMSC on L-RADA16, D-RADA16, and L-RADA16-RGD scaffolds, especially on L-RADA16 scaffolds incorporating TGF-beta 1, whereas the proliferation of BMSC cultured on RADA16 scaffolds was not significantly different from the control group. Collectively, asymptotical delivery of TGF-beta 1 from these peptide scaffolds was shown within the first 8 hours, and after that initial burst

Figure 4 Release profiles for TGF-beta 1 through three types of self-assembling peptide hydrogels (data points represent the average of three samples).

Abbreviations: h, hours; TGF, transforming growth factor.

Figure 5 Release profiles for TGF-beta 1 through three types of self-assembling peptide hydrogels 3D cells coculture (data points represent the average of three samples).

Abbreviations: h, hours; TGF, transforming growth factor; 3D, three-dimensional.

and similar release continued for the next days on condition that the whole release systems should not be disturbed and the diffused and released TGF-beta 1 should be kept in balance. The initial burst release might also be affected by the covered TGF-beta 1 on the surface peptide scaffolds. This result suggested that the release effect of active TGF-beta 1 was possibly able to continue for at least 2 weeks till the storage of the diffused growth factor was exhausted, and the electrostatic charges in the self-assembling nanofibers may interact with the TGF. These findings may be helpful for tissue regeneration strategies with sustained release of TGF-beta 1 to stimulate and promote the chondrogenesis or osteogenesis.

Discussion

Physicochemical properties of self-assembling peptide nanofiber hydrogel

Self-assembling peptide nanofiber hydrogel provides the possibility of slow and sustained release of growth factors mainly due to its stable network structures. Circular dichroism spectroscopy image showed that L-RADA16-RGD had stable β -sheet structure in NaCl solution at 25°C (Figure 1). But the secondary structure of this peptide can be affected by high temperature. CD spectrum also showed the L-RADA16 and D-RADA16 had a stable β -sheet secondary structure at 4°C. The change in the secondary structure of D-RADA16 after the stimulation of PBS is much stronger than L-RADA16 suggesting that D-RADA16 is more easily affected by ions than L-RADA16 (unpublished data). Rheometry assays showed that L-RADA16 and D-RADA16 nanofiber hydrogels (5.0 mg/mL) had stable mechanical features. The increase in concentration can improve the stability of hydrogels.

Figure 6 Proliferation effect of released TGF-beta 1 over bone mesenchymal stem cells.

Notes: The cells and RADA16 hydrogels contained TGF were co-cultured for 3 days (A), 7 days (B), and 14 days (C). CCK-8 assay for proliferated progeny (n=5). In the case of L-RADA16, a long-term effect of released TGF-beta 1 is evidenced by significantly higher total cell population. The TGF-beta 1 mitogenic activity can be appreciated till 2 weeks after being mixed with the self-assembling scaffolds. *P<0.05.

Abbreviations: TGF, transforming growth factor; OD, optical density.

Growth factor release through the self-assembling peptide hydrogel

It is critical to apply biomaterials for regulating cellular microenvironment and tissue regeneration.^{1,2,23} RADA self-assembling peptide nanofiber hydrogels formed 3D pores with diameter similar to spherical protein and provided an ideal network structure for cell growth. They also provided a natural physical barrier for controlled release of functional proteins. Figure 2 shows that the nanofiber and pore size of D-RADA16 hydrogel was smaller than that of L-RADA16 and L-RADA16-RGD, which may account for the slow release rate of TGF-beta 1 from D-RADA16 hydrogel than other

hydrogels (Figures 4 and 5). However, the secondary structure of D-RADA16 was found to be more easily affected by high temperature and ions.^{24,25} An initial burst of TGF-beta 1 that occurred in the first 8 hours maybe due to the surface location of protein molecules (Figures 4 and 5). Molecular motion might exist in the highly interwoven nanofiber network of the hydrogel.^{9,14,22} In addition, the release rate of TGF-beta 1 from the peptide hydrogel (1% w/v) was slowed down for 24 hours, after that a plateau value occurred asymptotically. However, a 100% molecule release is seldom observed in the hydrogels. Protein molecules in the interwoven nanofiber network structure of the hydrogel scaffold are physically obstructed and held

back and most diffusate cannot move freely, which might be the cause for that.^{7,26} Besides, our study also demonstrated that RADA16 self-assembling peptide would not affect the activity of TGF-beta 1 in cell proliferation.

Cell proliferation

Self-assembling peptide hydrogels are composed of amino acids, which have the function of biological recognition and good compatibility. RADA16 self-assembling peptide hydrogel can highly simulate the extracellular matrix and form suitable microenvironment for cell growth and functional protein action. During the release process, protein structure and bioactivity are maintained. TGF-beta 1 can promote the proliferation of mesenchymal stem cells.^{19,20} In Figure 6, all experiment groups had better cell proliferation than the control group. Besides, cell proliferation on the L-RADA16 + TGF-beta 1 group was significantly better than that on the L-RADA16 group, indicating the promotion of released TGF-beta 1

to proliferation of mesenchymal stem cells. In Figures 4 and 5, the release rate and release amount of TGF-beta 1 of L-RADA16 + TGF-beta 1 group were higher compared with those of the D-RADA16 + TGF-beta 1 group on 7 days. Previous studies showed that D-form self-assembling peptide hydrogels had the ability of being resistant to natural proteases degradation and promoting cell proliferation, which were beneficial to tissue regeneration and repair.^{10,27} The L-RADA16 + TGF-beta 1 group also showed a better effect on cell proliferation than that in the L-RADA16-RGD + TGF-beta 1 group. But the effect of L-RADA16-RGD on promoting cell differentiation, migration, and adhesion is greater than that of L-RADA16.^{28–32} These results indicated that the sustained release of TGF-beta 1 from L-RADA16 hydrogel had better promotion to cell proliferation than L-RADA16-RGD hydrogel, which was consistent with the release profiles of TGF-beta 1 from peptide hydrogel. In Figure 7, compared with the control group, BMSC in RADA16 peptide

Figure 7 The morphological results of BMSC in each group under a light microscope ($\times 100$).

Notes: (A) L-RADA16 group, (B) L-RADA16 + TGF-beta 1 group, (C) D-RADA16 + TGF-beta 1 group, (D) L-RADA16-RGD + TGF-beta 1 group, and (E) negative control.

Abbreviations: BMSC, bone mesenchymal stem cells; TGF, transforming growth factor.

hydrogel showed no obvious change in morphology, cell apoptosis, and degenerescence, which also demonstrated that the RADA16 peptide hydrogels were atoxic and safe for cells. Thus, RADA16 hydrogel can be used as an ideal tissue engineering material for the regeneration and repair of tissues or organs.

Conclusion

Our studies focused on controlled release of TGF- β 1 from RADA16 functional self-assembling peptide hydrogels as well as the promotion to cell proliferation. With 3D cell culture to the microenvironment formed by peptide hydrogels, there was a significant difference in high cell viability and low-level cell apoptosis between the L-form self-assembling peptide and D-form peptide scaffolds. This study using two chiral self-assembling peptides and functional motifs benefited to stimulate and promote the development of new biomaterials for tissue engineering at amino acids level and may further enhance the application of designer self-assembling peptides in clinical medicine and nanomedicine.

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Disclosure

The authors report no conflicts of interest in this work.

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