Development of Magnetic Nanocomposite Hydrogel with Potential Cartilage Tissue Engineering

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ABSTRACT: Magnetic nanocomposite hydrogels show high potential to improve tissue engineering. In this study, a magnetic nanocomposite hydrogel was prepared from poly(vinyl alcohol), nano-hydroxyapatite (n-HA), and magnetic nanoparticles (Fe₂O₃) using the ultrasonic dispersion method and freeze–thaw cross-linking molding. The water content and crystallinity of the magnetic nanocomposite hydrogel were tested. Microscopic morphology assessment, mechanical testing, and characterization were performed. Additionally, the magnetic nanocomposite hydrogel was co-cultured with bone mesenchymal stem cells (BMSCs) to determine its cell compatibility. We found that the magnetic nanocomposite hydrogel had good mechanical properties and that its mechanical properties were enhanced by the addition of n-HA. The BMSCs showed uniform growth on the surface of the magnetic nanocomposite hydrogel and high rates of proliferation. BMSC growth was also enhanced by the addition of Fe₂O₃ and also significant stimulated chondrocyte-related gene expression. Thus, the magnetic nanocomposite hydrogel scaffold material we describe here could have broad applications in cartilage tissue engineering.

1. INTRODUCTION

Articular cartilage defects are one of the most common forms of orthopedic disease and are challenging to treat in the clinic. Articular cartilage must be replaced after injury or lesion because it has limited ability to self-repair. The emergence of orthopedic disease and are challenging to treat in the clinic. Articular cartilage must be replaced after injury or lesion because it has limited ability to self-repair. The emergence of orthopedic disease and are challenging to treat in the clinic. Articular cartilage must be replaced after injury or lesion because it has limited ability to self-repair. The emergence of orthopedic disease and are challenging to treat in the clinic. Articular cartilage must be replaced after injury or lesion because it has limited ability to self-repair. The emergence of orthopedic disease and are challenging to treat in the clinic. Articular cartilage must be replaced after injury or lesion because it has limited ability to self-repair. The emergence of orthopedic disease and are challenging to treat in the clinical environment for seed cells. The chemical composition and crystal structure are similar to the inorganic component of the natural bone. The addition of HA could enhance the mechanical properties of scaffolds, which provide a template for a covering chondrocyte suspension and render the scaffolds sufficient integration with the host tissue to protected subchondral bone regeneration. Magnetic nanoparticles (Fe₂O₃) can promote the proliferation and differentiation of bone mesenchymal stem cells (BMSCs). Previous studies have integrated Fe₂O₃ into a porous scaffold to create a magnetic scaffold and found enhanced proliferation and adhesion of BMSCs in vitro. It is reported that labeling BMSCs with Fe₂O₃ causes cells to transfer from the

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extracellular matrix to the sites of cartilage damage, which promotes the differentiation of BMSCs into cartilages and stimulates chondrogenesis. Previous findings by Ngadiman et al. and Sheikh et al. have shown the biocompatibilities of Fe₂O₃/PVA and n-HA/PVA, and they are recommended to execute a side-by-side study comparing the proliferation and the differentiation of cells when they are seeded on PVA, Fe₂O₃/PVA, n-HA/PVA, and n-HA/Fe₂O₃/PVA. In this study, we prepared a magnetic nanocomposite hydrogel using PVA, n-HA, and Fe₂O₃. We found that the Fe₂O₃ regulated cell behavior within the scaffold material and had good cell compatibility; thus, they were used to construct tissue-engineered cartilage in vitro.

2. MATERIALS AND METHODS

2.1. Reagents and Instruments. The following reagents and instruments were used in this study: PVA, n-HA, Fe₂O₃ (Shenzhen Xincheng Biotechnology Co., Ltd.), Dulbecco’s modified Eagle’s medium (DMEM), trypsin, fetal bovine serum (FBS), 500 U/mL penicillin and 500 μg/mL streptomycin (Gibco USA), phosphate buffer solution (PBS), Cell Counting Kit-8 (CCK-8) (Keygen Tec), ultrasonic oscillator, weight meter, magnetic stirrer (Central Laboratory of The Second People’s Hospital of Shenzhen), differential scanning calorimeter (DSC) (Shenzhen Polytechnic), mechanical tester CTM4000 (Research Institute of Tsinghua University in Shenzhen), and electronic microscope (MIRA3 TESCAN, Research Institute of Tsinghua University in Shenzhen).

2.2. Preparation of Magnetic Nanocomposite Hydrogel. Figure 1 presents the schematic illustration of the fabrication of n-HA/Fe₂O₃/PVA hybrid magnetic nanocomposite hydrogel. Briefly, 1:0:0.5:10 were mixed and dissolved in deionized distilled water to produce Fe₂O₃. The n-HA/Fe₂O₃/PVA composite hydrogel was successfully prepared using the ultrasonic dispersion method and freeze-thawing cross-linking molding process. To create PVA aqueous solutions, varying amounts of PVA were weighed and placed in distilled water and heated and stirred at 90 °C for 1 h, to fully dissolve the PVA. Varying quantities of n-HA particles and Fe₂O₃ were added into the PVA aqueous solution; after ultrasonic treatment, the solution was stirred at 60 °C until evenly mixed. Next, after 30 min of heat preservation, all the bubbles in the solution were removed with a needle tube. The solution was poured into a mold, and then the mold was placed in the freezer (−20 °C) overnight for 16 h. Next, the solution was removed and thawed at room temperature for 4 h. Repeated freezing and thawing were performed seven times to obtain the n-HA/Fe₂O₃/PVA composite hydrogel (Figure 2A). The prepared n-HA/Fe₂O₃/PVA composite hydrogel was immersed in distilled water for one day to remove non-cross-linked monomers. As a control, we also concurrently prepared a pure PVA hydrogel (Figure 2B).

2.3. Water Content and Crystallinity of the Magnetic Nanocomposite Hydrogel. 2.3.1. Determination of Water Content of the Magnetic Nanocomposite Hydrogel. The surface water of the magnetic nanocomposite hydrogel was gently dried with a piece of filter paper and placed into a dry beaker of known mass (m₁). By weighing with a scale, the total mass of the hydrogel with beaker was m₂, and the mass of the magnetic nanocomposite hydrogel was calculated as m₃ = (m₂ − m₁). The magnetic nanocomposite hydrogel was placed in a vacuum-drying oven and was dried at 60 °C for 36 h, until it reached a constant weight. The mass of the dried magnetic nanocomposite hydrogel was m₄ and the water content of the dried magnetic nanocomposite hydrogel was determined as W = ([m₄ − m₃]/m₃).

2.3.2. Test of Magnetic Nanocomposite Hydrogel Crystallinity. The magnetic nanocomposite hydrogel was placed in a vacuum-drying oven and then dried at 60 °C for 24 h until a dried magnetic nanocomposite hydrogel was obtained. About 5 mg of the dried magnetic nanocomposite hydrogel (precision: 0.01 mg) was accurately weighed with a microbalance and placed in an aluminum crucible with a hole in the cover and then was measured with a DSC. An empty crucible of the same type was used as a control. High-purity nitrogen (volume fraction: 0.99999) was used as a protective gas and a purge gas, with a respective flow rate of 20 and 70 mL·min⁻¹. The temperature of the DSC was elevated from 10 °C/min to 250 °C using the programmed temperature to obtain the melting enthalpy of the dried magnetic nanocomposite hydrogel. This value was then used to calculate the crystallinity of the dried magnetic nanocomposite hydrogel.

2.4. Characterization of Hydrolytic Degradation. The weight loss method was used for the characterization of the hydrogel degradation performance in PBS. In this experiment, the same qualities of n-HA/Fe₂O₃/PVA hydrogel and pure PVA hydrogel were suspended in a PBS (pH 7.5) incubator at 37 °C for degradation experiments. At the selected time points (0, 2, 4, 6, 8, 10, 12, 14, 16, 18, 20 days), the qualities of the hydrogels in each group were measured.
Table 1. Primer Sequences for Real-Time PCR Analysis

| genes  | forward primer (5’→3’) | reverse primer (5’→3’) |
|--------|------------------------|------------------------|
| SOX-9  | GACGTGCAAGCTTGGGAAA    | CCGCAGGTATTTTGCAACCTC  |
| Col2A1 | GCCACGGCTCCTACATGTCC   | GTCACCCTCGGTGCTTGCAC   |
| AGG    | GCTACACCTAAGGCGCTGCT   | CGTAGTGTCCCTAGGTCATC   |
| GAPDH  | GGGCAGTCAAGGGCTGGAAATG | ATGGTGTGTAAGCCAGCT    |

2.5. Mechanical Testing and Microscopic Morphology Analysis. 2.5.1. Mechanical Testing. The elastic modulus measurement of the n-HA/Fe3O4/PVA composite hydrogel was characterized using an electronic universal testing machine CTM4000 in the mechanical testing laboratory. The diameter, thickness, and loading rate of the n-HA/Fe3O4/PVA composite hydrogel were 5 mm, 3 mm, and 5.00 mm/min, respectively.

2.5.2. Microscopic Morphology Analysis. The n-HA/Fe3O4/PVA composite hydrogel was frozen and dried. Next, after metal spraying, scanning analysis of the surface and cross section of the n-HA/Fe3O4/PVA composite hydrogel were conducted. The microscopic morphology of the n-HA/Fe3O4/PVA composite hydrogel was observed and analyzed using a TESCAN (Czech Republic) field emission scanning electron microscope. We acquired secondary electron images and backscattered electron images.

2.6. Cell Culture and Cell Compatibility Study. A New Zealand white rabbit (aged 3 months) was anesthetized and then underwent skin disinfection and preparation. The rabbit’s bone marrow was punctured at the left femur trochanter, and about 3 mL of bone marrow was extracted using a syringe containing 2 mL heparin. The extracted bone marrow was placed into a 10 mL centrifuge tube, and the PBS was added to reach 5 mL. The bone marrow solution was centrifuged at 1200 rpm for 10 min, and the supernatant was discarded. The pellet was washed twice with PBS, followed by centrifugation at 1000 rpm. High-glucose DMEM culture solution (containing 10% FBS, 1% double-antibody penicillin/streptomycin) (5 mL) was added to the precipitated cell suspension, and then the mixed solution was cultured in a CO2 incubator at 5% CO2 and 37 °C. Three days after culture, fresh medium was added to the cultures.

2.6.1. Cell Proliferation Assay. Cell proliferation was measured using the CCK-8 method. This experiment analyzed three groups: BMSCs co-cultured with n-HA/Fe3O4/PVA composite hydrogel (group A), BMSCs co-cultured with pure PVA hydrogel (group B), and BMSCs cultured alone (group C), with five repeat samples in each group. Cells at the exponential phase were seeded in 96-well plates at a density of 1 × 10^3 cells per well. On the first, third, fifth, seventh, and ninth days of cell culture, 50 μL CCK-8 solution was added to each complex hole and then incubated at 37 °C for 4 h. Next, the solution in each complex hole was transferred to a 96-well plate, and the amount of solution in each well was equalized.

2.6.2. Stereoscopic Optical Microscopy. This experiment examined two groups: BMSCs cultured with n-HA/Fe3O4/PVA composite hydrogel (group A) and BMSCs cultured with pure PVA hydrogel (group B). The culture plate was uncovered after one week of culture and placed under a stereotropical microscope. A 100X magnification was used to observe cell growth.

2.6.3. Scanning Electron Microscopy. This experiment examined two groups: BMSCs cultured with n-HA/Fe3O4/PVA composite hydrogel (group A) and BMSCs cultured with pure PVA hydrogel (group B). After one week of culture, the hydrogels in each group were washed twice with PBS, fixed with 2.5% glutaraldehyde fixative at 4 °C for 2 h, and then dehydrated progressively with ethanol. After critical-point drying with CO2 and metal spraying, the adhesion of cells on the scaffold was observed with a scanning electron microscope.

2.7. Histological and Immunohistochemical Analyses to Detect the Collage Protein. For histology, the cell/scaffold constructs were fixed in 2.5% glutaraldehyde for 16 h, through a series of gradient ethanol to dehydrate, embedded in paraffin wax, and cut into 5 μm thick sections, and then the samples were stained with hematoxylin and eosin (H&E). For immunofluorescent staining to detect the collage II protein expression levels, the cell/scaffold constructs were fixed with 4% parafomaldehyde for 10 min, permeabilized with 0.25% Triton X-100 for 10 min, and then blocked with 3% bovine serum albumin for 60 min. Subsequently, these constructs were incubated with a collagen type II antibody (ab34712, 1:100, Abcam, Cambridge, USA) and then were subsequently exposed to Alexa Fluor 594-conjugated secondary antibody overnight. Nuclei were counterstained with 4′,6-diamidino-2-phenylindole (DAPI) and are shown in blue. The constructs were cut into thick sections and were visualized using laser-scanning confocal microscopy (ZEISS, Germany).

2.8. RNA Extraction and Real-Time Polymerase Chain Reaction Analysis. Total RNA was isolated using Trizol and reverse-transcribed into complementary DNA using a DNA synthesis kit (TakaRa, Shiga, Japan) according to the manufacturer’s protocol. Real-time polymerase chain reactions (PCRs) of glyceraldehyde phosphate dehydrogenase (GAPDH), collagen II, and aggrecan were performed using the SYBR Green QPCR Master Mix. Primer sequences are listed in Table 1. Real-time PCRs were performed using a 7500 real-time PCR detection system (ABI, Foster City, CA) at 95 °C for 15 min followed by 40 cycles of denaturation at 95 °C for 10 s, extension at 60 °C for 15 s, and annealing at 72 °C for 15 s. Gene expression levels were normalized to the housekeeping gene GAPDH, and the relative gene expression was calculated using the ΔΔCT method.
3. RESULTS AND DISCUSSION

3.1. Water Content and Crystallinity of the Magnetic Nanocomposite Hydrogel.

3.1.1. Water Content of the Magnetic Nanocomposite Hydrogel. The water content of the novel magnetic nanocomposite hydrogel we prepared was 82.6%, lower than the water content of the pure PVA hydrogel (85.3%, Figure 3A). This phenomenon could be explained as follows: PVA was closely linked with n-HA and Fe₂O₃, producing stress shrinkage, causing molecules to become densely packed, forcing water in the pores to be discharged, and decreasing the volume of the magnetic nanocomposite hydrogel. Consequently, the hydroxyl of water molecule-producing hydrogen bond left little space to accommodate water molecules, which resulted in lower water content.

3.1.2. Crystallinity of the Magnetic Nanocomposite Hydrogel. Scaffolds with high crystallinity can have a high elastic modulus and elongation at break. Changes to the crystallinity of the dried magnetic nanocomposite hydrogel could indirectly suggest that its structure has changed. It was believed that a polymer’s crystallinity is proportional to its melting heat when melted. The melting heat can be measured using DSC atlas. Figure 3B shows the DSC atlas of the dried n-HA/Fe₂O₃/PVA composite hydrogel and the DSC atlas of the dried pure PVA hydrogel. As shown in Figure 3B, the dried n-HA/Fe₂O₃/PVA composite hydrogel showed a sharp endothermic peak around 217 °C, corresponding to its melting heat, and showed a wide endothermic peak at 231 °C, corresponding to its decomposition. The dried pure PVA hydrogel showed a sharp endothermic peak around 221 °C, corresponding to its melting heat, and showed a wide endothermic peak at 236 °C, corresponding to its decomposition. By comparing to previously reported values of the melting heat of 100% crystallized PVA, the crystallinity of the hydrogel samples was obtained using the following equation

\[ f(DSC) = \left( \frac{\Delta H}{\Delta H^0} \right) \times 100\% \]

where \( f(DSC) \) is the crystallinity of the hydrogel samples measured through differential thermal analysis, \( \Delta H \) is the melting heat enthalpy of the hydrogel samples, and \( \Delta H^0 \) is the melting heat enthalpy of the PVA hydrogel with 100% crystallinity, reported to be 138.6 J/g. Table 2 shows the melting heat enthalpies of the dried n-HA/Fe₂O₃/PVA composite hydrogel and the dried pure PVA hydrogel and their crystallinities as calculated using the formula above.

3.2. Microscopic Morphology Analysis. The scanning electron microscopy (SEM) showed that many pores of different sizes were distributed on the surface of the magnetic nanocomposite hydrogel and that magnetic particles and n-HA particles were uniformly distributed in the magnetic nanocomposite hydrogel, forming rough bumps on the surface. These pores and bumps were beneficial to the adhesion and growth of cells on the surface of the magnetic nanocomposite hydrogel. In contrast, relatively few pores were distributed on the surface of the pure PVA hydrogel (Figure 3C,D).

3.3. Mechanical Property and the Degradation Testing. The mechanical properties of the hydrogel samples were detected by a three-point bending resistance experiment. The results showed that the n-HA/Fe₂O₃/PVA composite hydrogel had better mechanical properties than the pure PVA hydrogel, including greater tensile strength, greater bending strength, and a higher bending modulus (as shown in Table 3). The in vitro
degradation of magnetic nanocomposite hydrogel at different time points has shown that the magnetic hydrogel formulations have relatively intermediate and slow degradation rates (Figure 4).

### 3.4. Cell Compatibility

#### 3.4.1. Identification of the BMSCs

To confirm the stem cell characterization for the next study, we used multilineage differentiation potential of BMSCs. We successfully characterized the differentiation capacities of rabbit (rb)-BMSCs into various lineages like chondrogenic, adipogenic, and osteogenic cells (Figure 5C–E).

#### 3.4.2. Cell Proliferation Assay

We measured BMSC proliferation on the n-HA/Fe2O3/PVA composite hydrogel using the CCK-8 method (Figure 6). There was no significant difference in cell proliferation between the BMSCs cultured with n-HA/Fe2O3/PVA composite hydrogel and either of the control groups (i.e. BMSCs cultured with pure PVA hydrogel or BMSC culture alone, \( p > 0.05 \)).

#### 3.4.3. Stereoscopic Optical Microscopy

After one week of BMSC culture with different forms of hydrogel, we performed stereoscopic optical microscopy. When the BMSCs were cultured with the n-HA/Fe2O3/PVA composite hydrogel (group A), cells had a polygon shape, contacted each other through projections, proliferated and differentiated well, and showed a regular stacking growth with time. When the BMSCs were combined with the pure PVA hydrogel, they had a polygon shape and proliferated well but we observed relatively few cells. This result shows that the magnetic nanocomposite hydrogel had better biocompatibility than pure PVA hydrogel alone.

#### 3.4.4. SEM

According to our SEM results, when BMSCs were cultured with n-HA/Fe2O3/PVA composite hydrogel (group A), a large number of spindle or polygonal cells adhered to and aggregated in the pores on the gel surface. In contrast, only a small number of cells adhered to the pores on the surface of the pure PVA hydrogel (group B). This demonstrates that

### Table 3. Mechanical Properties of the n-HA/Fe2O3/PVA Composite Hydrogel and the Pure PVA Hydrogel (\( \bar{x} \pm s, n = 10 \))

| name of the hydrogel          | tensile strength (MPa) | bending strength (MPa) | bending modulus (GPa) |
|------------------------------|------------------------|------------------------|-----------------------|
| the n-HA/Fe2O3/PVA composite hydrogel | 28.7 ± 0.6*a          | 83.5 ± 0.2*a           | 1.7 ± 0.1*a           |
| the pure PVA hydrogel        | 26.3 ± 0.3             | 79.7 ± 0.7             | 1.3 ± 0.1             |

*aStatistical analysis of variance, \( p < 0.05 \).
the magnetic nanocomposite hydrogel had good biocompatibility with the BMSCs (Figure 7A,B).

Figure 7. Hydrogel material was co-cultured with BMSCs for one week. (A) A large number of spindle or polygonal cells adhered to and aggregated in the pores on the surface of the n-HA/Fe$_2$O$_3$/PVA composite hydrogel. (B) A small number of cells adhered to the pores on the surface of the pure PVA hydrogel.

3.4.5. Photomicrographs of the Histological Findings. The BMSCs formed a capsule around the construct in the central region of the cell/scaffold constructs (Figure 8A,B). By contrast, at the peripheral edge of the cell/scaffold constructs, clumps of cells were found and some cells were well-dispersed. Also the cells at the surface were larger compared with those in the central region (Figure 8C,D).

The levels to enhance the immunofluorescent signal of the chondrogenic differentiation marker were obviously observed in the n-HA/Fe$_2$O$_3$/PVA composite hydrogel group (Figure 9).

3.4.6. Chondrogenic Differentiation Potential of Hydrogel Material. BMSCs cultured with the n-HA/Fe$_2$O$_3$/PVA composite hydrogel exhibited significantly higher SOX9, aggrecan, and type II collagen expression (Figure 10). This material significantly enhances the expression of cartilage-specific gene markers such as type II collagen, aggrecan, and SOX9 relative to pure PVA hydrogel during the culture period, thereby inducing differentiation of BMSCs into hyaline chondrogenic cells. In conclusion, we demonstrated that our magnetic nanocomposite hydrogel scaffold material promotes chondrogenic differentiation. Thus, the magnetic nanocomposite hydrogel may be a reliable scaffold for cartilage tissue engineering. In addition, this study will advance the development of magnetic strategies in tissue engineering to repair the damaged cartilage tissue.

Figure 8. Photomicrographs of the histological findings in the central (AB) and surface regions (C,D) of the cell/scaffold constructs, in static culture for 14 days (20× magnification).

Figure 9. Confocal images of cells immunostaining of collagen type II antibody; cell nuclei were labeled with DAPI (blue) (scale bar is 20 μm).

4. DISCUSSION AND CONCLUSION

At present, the biomaterials studied in cartilage tissue engineering mainly include natural or synthetic polymers, injectable hydrogels, or other scaffold materials. A clinically useful scaffold material for cartilage tissue engineering will have good biocompatibility, biodegradability, low cytotoxicity, and certain mechanical properties. The scaffold material acts as an exogenous cell carrier, namely, it promotes the migration, adhesion, proliferation, and differentiation of BMSCs, or provides structural support. In this study, we describe a novel magnetic nanocomposite hydrogel prepared using the repeated freeze–thaw approach. The aqueous solution of the composite material formed physical cross-linking points in the form of tiny crystalline regions via intermolecular H-bonds in the process of repeated freezing and thawing, which then formed a hydrogel with a three-dimensional network structure with considerable mechanical strength and ability to swell. No harmful chemicals were added throughout the process, and the biocompatibility of the composite material was not reduced. Our novel hydrogel also has high water content and good elasticity and can be prepared quickly and easily.

Water content is an important property of hydrogels. In the hydrogel system, water content has a direct effect on the viscoelasticity and mechanical properties of the material. A hydrogel’s compressive strength, tensile strength, and modulus increase with decreasing water content. When the water content is 80%, the tensile strength is similar to natural cartilage. When the water content is less than 80%, the homogeneity of the hydrogel becomes problematic. The water content of the magnetic nanocomposite hydrogel prepared in this study was 82.6%, slightly less than the water content of the pure PVA hydrogel (85.3%, Figure 3). During hydrogel formation, the molecular chains in the amorphous region unwrap and the lamellar structure in the crystalline region begins to expand. This expansion is accompanied by the turnover and sliding of molecular chains, a process that absorbs the energy generated by the viscoelastic deformation. Therefore, high crystallinity can be compatible with high elastic modulus and elongation at break. Changes to the crystallinity of a hydrogel can indirectly suggest that its structure has changed. Table 2 shows that the crystallinity of the n-HA/Fe$_2$O$_3$/PVA composite hydrogel is lower than that of the pure PVA hydrogel, indicating that adding n-HA and Fe$_2$O$_3$ reduced...
the entanglement points of interchain and intrachain hydrogen bonds formed in the PVA hydrogel. While crystallinity decreased, the structure of amorphous region increased and the network structure of the PVA hydrogel became loose, thus the network gap in unit volume increased, resulting in lower crystallinity of the PVA hydrogel.20

Mechanical testing showed that the mechanical properties of the n-HA/Fe$_2$O$_3$/PVA composite hydrogel were superior to the pure PVA hydrogel. This may be caused by the addition of n-HA and Fe$_2$O$_3$ to the PVA hydrogel, as n-HA enhanced the mechanical properties of the composite hydrogel while promoting the repair of subchondral bone. SEM showed that many pores of different sizes were distributed on the surface of the n-HA/Fe$_2$O$_3$/PVA composite hydrogel, forming rough bumps on the surface. These structures were beneficial to the adhesion and growth of cells on the surface of the composite hydrogel, and the pores may have improved the transmission and metabolism of nutrients. When the third-generation BMSCs were co-cultured with the n-HA/Fe$_2$O$_3$/PVA composite hydrogel, we found that the cells were polygon-shaped, contacted each other through projections, and showed regular stacking growth with time. Additionally, proliferation, survival, and differentiation rates were high in BMSCs co-cultured with the composite hydrogel. A large number of spindle or polygonal cells adhered to and aggregated in the pores on the surface of the n-HA/Fe$_2$O$_3$/PVA composite hydrogel, showing that the n-HA/Fe$_2$O$_3$/PVA composite hydrogel had good biocompatibility. Moreover, the prepared n-HA/Fe$_2$O$_3$/PVA composite hydrogel showed an improvement in the differentiation of the BMSCs.

Cartilage tissue engineers have always aimed to develop novel scaffold materials with adequate mechanical strength, high biocompatibility, and hydrophilicity, which are easy to degrade and absorb. The scaffold material we describe in this study not only had all of the above characteristics but also had featured superparamagnetic nanoparticle Fe$_2$O$_3$, which can promote the proliferation and differentiation of the BMSCs and closely bind to the cell surface.

In this experiment, a novel magnetic nanocomposite hydrogel scaffold material was prepared. This hydrogel had similar water content as natural cartilage and low crystallinity. This scaffold material had good mechanical properties, and SEM demonstrated that micropores on the hydrogel surface were beneficial to the growth and proliferation of cells. In addition, experiments where BMSCs were cultured with the magnetic nanocomposite hydrogel scaffold material demonstrated that the scaffold material had good cell compatibility and induced BMSC differentiation into chondrocytes. Thus, this scaffold material has the potential for a broad range of therapeutic applications and can be considered as a promising material for cartilage tissue engineering.

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