

# Osteogenic ability using porous hydroxyapatite scaffold-based delivery of human placenta-derived mesenchymal stem cells

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**Abstract.** Previous preliminary studies have suggested that hydroxyapatite with a grooved structure (HAG) scaffold has good osteogenic potential. This type of scaffold may aid osteogenesis during the repair of large maxillofacial bony defects. The ectopic osteogenic effect and underlying mechanism were further studied using porous HAG scaffold-based delivery of human placenta-derived mesenchymal stem cells (hPMSCs). A total of 18 dogs were randomly allocated into a HAG scaffold group and a HAG scaffold-based hPMSC (HAG/hPMSC) group, and three scaffolds were implanted into the dorsal muscle of each dog. Samples were taken for subsequent analysis and tested 4, 8 and 12 weeks following heterotopic implantation. H&E staining was used to study the osteogenic effect in dog dorsal muscles, and RNA sequencing (RNA-seq) was used for exploring the underlying osteogenic mechanism. The osteogenic ability and effector of the HAG/hPMSC group were significantly greater than those of the HAG scaffold group at 4 weeks after implantation. After 12 weeks, a mature bone plate structure was seen in the HAG/hPMSC group. RNA-seq demonstrated that various osteogenesis-related pathways participated at different stages of metabolism, and that the expression of collagen-1 and runt-related transcription factor 2 increased with implantation time. The present study preliminarily focused on the ectopic osteogenic effect of the

porous HAG scaffold-based delivery of hPMSCs *in vivo*, which may be helpful for the improved application of HAG scaffolds in the future.

## Introduction

Maxillary defect or deficiency are the most common problems in dental implantation, especially in the treatment of patients with extensive jawbone defects caused by trauma or tumor resection (1). Due to the rapid development of material science, bone tissue engineering has attracted wide attention, and extensive research has been carried out on the subject. Bone tissue engineering strikes a balance between the advantages and disadvantages of autogenous bone and allogeneic bone (2). Artificial bone substitute materials are based on bone tissue engineering using scaffolds, growth factors and stem cells; these have good histocompatibility and reproducibility, and are expected to be good substitutes for bone defect repair (3).

Hydroxyapatite (HA), as an inorganic mineral, is the main component of natural bone inorganic salts, and an indispensable inorganic component of the human skeleton (4). Research demonstrates that 50% of the human skeleton is comprised of homogeneous inorganic HA (4). The physicochemical properties, chemical composition and crystal structure of HA are very similar to human bone and are highly safe for humans. The results of our previous study demonstrated that porous HA with a grooved structure (HAG) scaffolds enhanced osteogenesis *in vivo* and *in vitro* (5).

Transcriptome sequencing is a technique for determining the expression level of the transcriptome, and is widely used in research (6). RNA sequencing (RNA-seq) is a high-throughput sequencing assay that can provide detailed information regarding the working mechanisms underlying the target tissue or the molecular pathobiology of a disease (7-11). Thus, RNA-seq plays an important role in studying the mechanisms underlying osteogenesis. High-throughput transcriptome profiling was used to further understand the molecular mechanism underlying lithium in regulating the osteogenic fate of human mesenchymal stem cells (hMSCs), when stimulated with lithium for 7 days (12).

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According to the International Society for Cellular Therapy (ISCT) (13), mesenchymal stem cells (MSCs) must satisfy three requirements: i) MSCs must be plastic-adherent when maintained in standard culture conditions; ii) MSCs must express CD105, CD73 and CD90, and lack the expression of CD45, CD34, CD14 and human leukocyte antigen (HLA)-DR surface molecules and either CD14 or CD11b, and either CD79a or CD19; and iii) MSCs must differentiate to osteoblasts, adipocytes and chondroblasts *in vitro* (13). The criteria do not support the purification of homogenous MSC populations. The isolation of MSCs according to ISCT criteria produces heterogeneous, non-clonal cultures of stromal cells containing stem cells with different multipotent properties, committed progenitors and differentiated cells. Currently, MSCs can be isolated from multiple tissues, such as placenta (14,15).

In the present study, the RNA-seq technique was used to study the osteogenic mechanism underlying porous HA scaffold-based delivery of human placenta-derived (hP) MSCs in the dorsal muscles of dogs. Samples were taken for sequencing 4, 8 and 12 weeks after heterotopic implantation. The association between the expression and time of different genes was identified by analyzing differentially expressed genes, and the osteogenesis mechanism underlying the porous HAG scaffold-based delivery of hMSCs was investigated using results of transcriptome sequencing. The novelty of the present study was demonstrated by the analysis of the biological properties of the porous HAG scaffold-based delivery of hMSCs via transcriptomics, which provided novel applications of this cell type in osteogenesis research.

## Materials and methods

**Preparation of cell-adhered HAG scaffolds for transplantation.** Cell culture conditions were the same as in our previous study (5): High glucose DMEM (Thermo Fisher Scientific, Inc.) containing 10% FBS (Thermo Fisher Scientific, Inc.), 20 mmol/l  $\beta$ -glycerophosphate, 50 g/ml vitamin C and 10 mol/l dexamethasone at 37°C in 5% CO<sub>2</sub>. The immortalized hPMSCs used in this study originated from the State Key Laboratory of Biological Therapy at Sichuan University (provided by the National Experimental Cell Resource Sharing Platform, <http://www.cellresource.cn/>). The fifth generation of hPMSCs in the exponential growth period were counted. According to the counting results, high glucose DMEM was added until the cell density reached 2x10<sup>5</sup>/ml. According to our previous study (5), cells were analyzed using a flow cytometer (FACS Aria III; BD Biosciences) with related antibodies (all dilutions, 1:100; all from Abcam) for the following surface markers: HLA-DR (cat. no. ab92511), CD11b (cat. no. ab133357), CD14 (cat. no. ab183322), CD19 (cat. no. ab134114), CD34 (cat. no. ab81289), CD44 (cat. no. ab189524), CD45 (cat. no. ab40763), CD73 (cat. no. ab202122), CD79 (cat. no. ab134147), CD90 (cat. no. ab23894) and CD105 (cat. no. ab231774) (16,17). A total of 5x10<sup>5</sup> cells were cultured in a flow tube at 37°C, after which cells were re-suspended into monocytes using 100 IX PBS and antibodies (1  $\mu$ l) were added. Samples were incubated at 4°C for 30 min in the dark and analyzed using FlowJo 10.6.2 software (BD Biosciences).

The cells were directly dripped onto the surface of the porous HAG scaffold until the material was just infiltrated by the cell suspension. Following incubation for 3 h, the HAG scaffold was turned over, cell suspension droplets were added to the other side of the scaffold and culture medium (high glucose DMEM) was added following a further 3 h in the incubator. Tissue engineered bone was cultured in a constant temperature incubator at 37°C on a six-well plate for 7 days, and the solution was changed once every 1-2 days. hPMSCs were cultured in low-glucose DMEM supplemented with 10% FBS and 1% penicillin-streptomycin in a 5% CO<sub>2</sub> atmosphere at 37°C.

**Preparation and characterization of the scaffolds.** The HAG scaffolds were used in this study as described by our previous work (5).

**Experimental animals and implantation.** All animal experiments were conducted according to the protocols approved by the Animal Care and Use Committee of Sichuan Provincial People's Hospital [Sichuan, P.R. China; approval no. 2019NSF(98)]. A total of 18 male beagles (age, 12 months old; weight range, 10-12 kg) were selected as the experimental samples. Beagles were housed at 16-28°C (softened animal feed with milk and water were obtained freely from 8 a.m. to 5 p.m.) and under relative humidity of 40-80%. They were exposed to a 12 h light/dark cycle and ammonia concentrations of <14 mg/m<sup>3</sup>, which was administered to avoid the occurrence of disease. Beagles were provided by the Experimental Animal Center of Sichuan Provincial People's Hospital (Sichuan, P.R. China). All animal surgery was approved by the Institutional Animal Care and Use Committee and implemented according to relevant requirements. The beagles were used for a duration of 4 months in the present study. Pentobarbital sodium was injected intravenously into the forelimb for general anesthesia at a concentration of 30 mg/kg. The animals were euthanized 1 month after the experiment, using pentobarbital sodium injected intravenously into the forelimb at a concentration of 100 mg/kg. The death of all animals was confirmed by cardiac arrest, respiratory arrest and loss of reflex. A total of 18 beagles were randomly allocated into one of the following groups, with 3 beagles/group: i) HAG group with stem cell implantation (HAG/hPMSC group) and ii) HAG group without stem cell implantation. The scaffolds were implanted separately into the dorsal muscle of each dog as described by the heterotopic implantation discussion in our previous study (5). The scaffold attached by muscle from each beagle was obtained, and RNA and protein were extracted for analysis by crushing with a grinder. A total of three scaffolds from the dorsal muscle of each dog were pooled together for RNA-seq. Western blotting analysis and qPCR were repeated three times.

**H&E staining and western blot analysis.** At 4, 8 and 12 weeks after surgery, the surface of the scaffold was washed with normal saline and placed into 10% neutral formaldehyde fixing solution at 4°C overnight. New bone was identified using optical microscopy of H&E-stained implanted materials. H&E staining can detect the formation of bone matrix, which can indicate osteogenic effects (18,19). Images were captured using an optical microscope (CX23; Carl Zeiss AG).

Following SDS-PAGE and membrane transfer, the membrane was incubated overnight at 4°C with IgG (Abcam; 1:300). Following incubation with the secondary antibody, buffer from the membrane was discarded and developing reagent was added. The PVDF membrane was rocked until development of bands was apparent. The membrane was washed three times using distilled water for 30 min. The membrane was visualized using ChemiDocMP (Bio-Rad Laboratories, Inc.).

**Isolation and purification of RNA.** For mRNA deep sequencing, total RNA was extracted from the control and treated frozen samples using TRIzol® (Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer's instructions. The quality of total RNA was checked using an Agilent 2100 Bioanalyzer (Agilent Technologies, Inc.). Equal amounts of RNA samples from three control individuals were pooled to generate a mixed sample for the control subgroup, and the same was repeated for the treated subgroup.

**cDNA synthesis and library construction for mRNA deep sequencing.** For mRNA sequencing, a cDNA library was constructed with a NEBNext® Ultra™ RNA Library Prep kit (Illumina, Inc.). The concentration of the library was >2 nM, indicating that the library was effective. Samples were diluted to 1 ng/μl and detected using a DNA Quantification kit (Illumina, Inc.). Briefly, mRNA was purified from total RNA using poly-T oligo-attached magnetic beads. Fragmentation was carried out using divalent cations under elevated temperature in NEBNext First Strand Synthesis Reaction Buffer (5X) (Illumina, Inc.). Index codes were added to attribute sequences to each sample and 150 bp paired-end reads were generated. The obtained library was used for sequencing with Illumina HiSeq™ 2500 (Illumina, Inc.).

Reference genome and gene model annotation files were downloaded from genome websites (NCBI, <https://www.ncbi.nlm.nih.gov/>) directly. An index of the reference genome was built using Bowtie v2.2.3 (<http://bowtie-bio.sourceforge.net/bowtie2/index.shtml>), and paired-end clean reads were aligned to the reference genome using TopHat v2.0.12 (<http://ccb.jhu.edu/software/tophat/manual.shtml>). TopHat was selected as the mapping tool as it can generate a database of splice junctions based on the gene model annotation files. We used KOBAS software (<http://kobas.cbi.pku.edu.cn/>) to test the statistical enrichment of differential expression genes in KEGG pathway.

**Reverse transcription-quantitative PCR (RT-qPCR).** To validate the RNA-seq data, differentially expressed genes were selected for RT-qPCR analysis. All primers were designed by the Premier5.0 software (Premier Biosoft International; Table I). RNA was extracted using the TRIzol kit (Thermo Fisher Scientific, Inc.). cDNA was synthesized at 37°C for 15 min and 85°C for 5 sec using the Prime Script® RT reagent kit with the gDNA Eraser kit (all from Takara Biotechnology Co., Ltd.). RT-qPCR was carried out with SsoFast™ EvaGreen® Supermix (Bio-Rad Laboratories, Inc.). Two mRNAs were selected at random from upregulated, downregulated and non-altered genes for qPCR analysis using β-actin as a reference gene. The fold change from the qPCR was determined using the  $2^{-\Delta\Delta C_q}$  method (20). The RNAs of samples were reverse transcribed into cDNA in a

Table I. Primers for reverse transcription-quantitative PCR.

Name	Sequence (5'-3')	Fragment, bp
<i>COL-1</i> -F	AGGGGTCTCCATGGTGAGTT	119
<i>COL-1</i> -R	GAAGGACCTCGGCTTCCAAT	
<i>Runx2</i> -F	TTCCAGAATGCTTCCGCCAT	110
<i>Runx2</i> -R	AACTGCTGTGGCTTCCATCA	
<i>β-actin</i> -F	CAATACAACCTCTCCACAACC	281
<i>β-actin</i> -R	CAGATAGCACCTTCAGCAC	

*Col-1*, collagen-1; *Runx2*, runt-related transcription factor 2; F, forward; R, reverse.

Table II. Statistical analysis of the ratio of total area occupied by the newborn.

Sample	Mean, %	Standard deviation, %
HAG-4w	3.7	0.3
HAG/hPMSC-4w	5.2	0.5
HAG-8w	6.5	1.2
HAG/hPMSC-8w	5.1	0.3
HAG-12w	11.2	0.3
HAG/hPMSC-12w	12.3	0.4

HAG, hydroxyapatite with a grooved structure; hPMSC, human placenta-derived mesenchymal stem cells; HAG/hPMSCs, HAG scaffold-based delivery of hPMSCs; w, weeks.

20 μl reaction containing 13 μl RNA template, 2 μl Oligo dT primer or microRNA sequence-specific primers (10 μM), 4 μl of 5X PrimeScript Buffer and 1 μl PrimeScript RT Enzyme Mix I (Takara Bio, Inc.). For subsequent qPCR reactions using gene-specific primers, 1 μl cDNA sample was amplified under the following cycle conditions: 94°C for 30 sec, followed by 39 cycles of 55°C for 20 sec and 72°C for 20 sec.

**Statistical analysis.** All the statistical data were analyzed using Bonferroni ANOVA using SPSS 20.0 statistics software (IBM Corp.). All data are presented as the mean ± SD.  $P < 0.05$  was considered to indicate a statistically significant difference. All analyses were repeated three times. Gene expression used double-screening by q-value (adjusted P-value,  $\leq 0.05$ ) and normalized log2ratio ( $\log_2\text{ratio} \geq 1$ ; Tables SI-SIII) according previous sequencing methods (21).

## Results

**Osteogenesis effects of the HAG-based delivery of hPMSCs.** The beagles in the HAG/hPMSC experimental group had HAG scaffolds implanted with hPMSCs implanted in their dorsal muscles, while the HAG control group was implanted with the HAG scaffolds alone. At a total of 4, 8 and 12 weeks after the operation, two groups of scaffolds were obtained from the dorsal muscles of the beagles and labelled as follows:



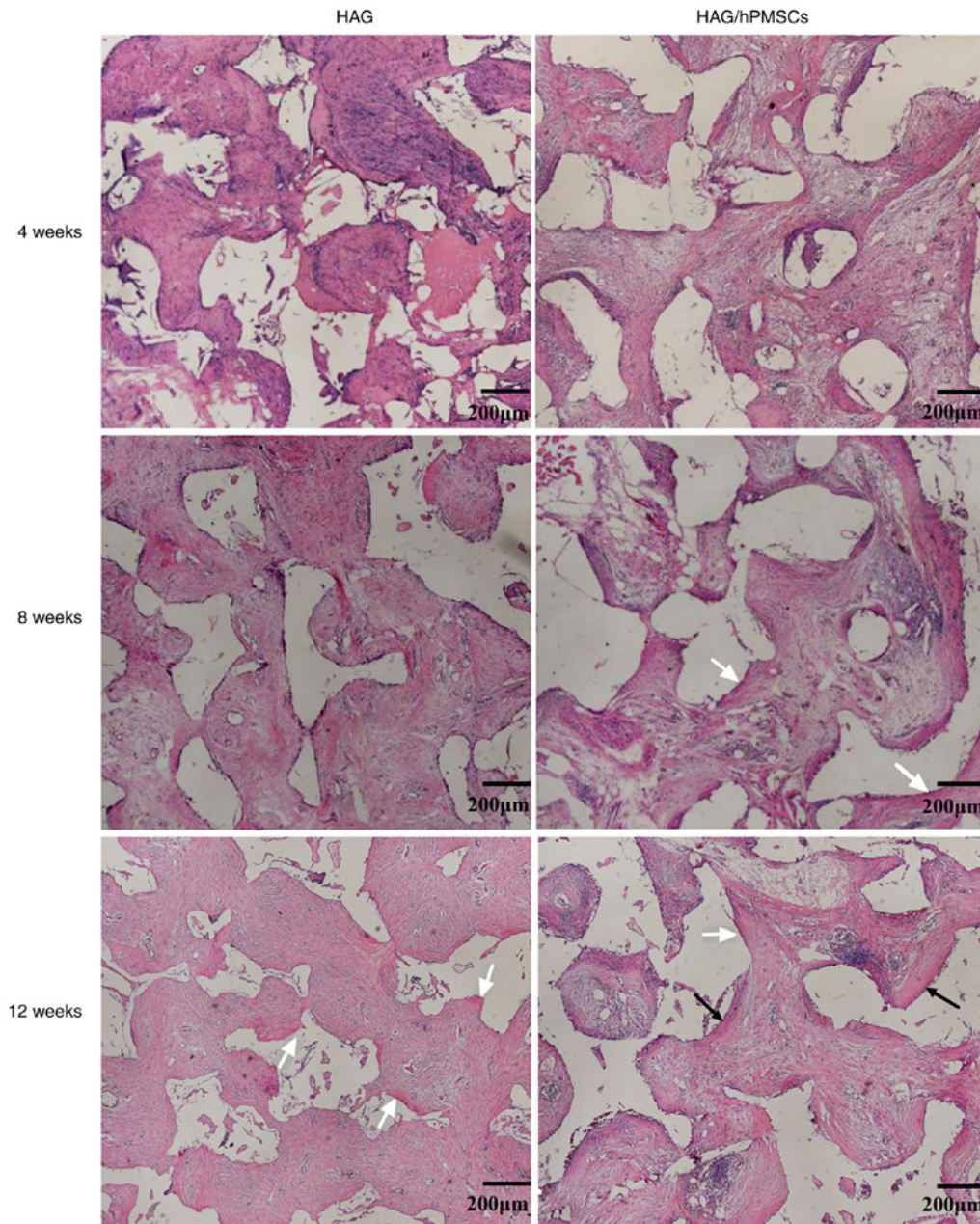


Figure 1. Bone formation of the scaffold analyzed using H&E staining. Magnification, x100. White arrow, bone matrix; black arrow, mature bone plate structure. HAG, hydroxyapatite with a grooved structure; hPMSC, human placenta-derived mesenchymal stem cells; HAG/hPMSCs, HAG scaffold-based delivery of hPMSCs; w, weeks.

HAG-4 weeks (w), HAG-8w, HAG-12w and HAG/hPMSC-4w, HAG/hPMSC-8w and HAG/hPMSC-12w.

H&E staining demonstrated that a large number of fibroblasts grew and fibrous tissue formed in both groups at 4 weeks after implantation, while osteoid formation was found on the surface of scaffolds in the HAG/hPMSC-4w compared with the HAG-4w group (Fig. 1). After 8 weeks, the area of bone matrix in the HAG/hPMSC-8w group was larger than that in the HAG group (Fig. 1). After 12 weeks, a large amount of bone matrix was formed in both groups, but mature bone plate structure was observed in the HAG/hPMSC-12w group (Fig. 1). The statistical results indicated that the osteogenic ability and effects of the HAG/hPMSC group were markedly greater than those of the HAG group (Table II).

*Illumina sequencing and mapping to reference the genome.* cDNA libraries were constructed for different RNA samples at different times of implantation, with one treatment and one control for Illumina sequencing. From these libraries, 49,398,934, 53,012,698, 51,462,894, 60,674,628, 53,302,412 and 48,165,136 raw reads were obtained from HAG/hPMSC-4w, HAG/hPMSC-8w, HAG/hPMSC-12w, HAG-4w, HAG-8w and HAG-12w, respectively, which generated 7.05, 7.59, 7.36, 8.65, 7.61 and 6.89 Gb of cleaned data. The schematic of Illumina deep sequencing and analysis is demonstrated in Table III.

High-quality reads were mapped to the coding sequences from the reference genome by Bowtie software, with the default parameters for the HAG group set as 48,759,095 (84.59%), 43,082,474 (84.9%) and 38,758,173 (84.42%). For the HAG/hPMSC group, default parameters were set

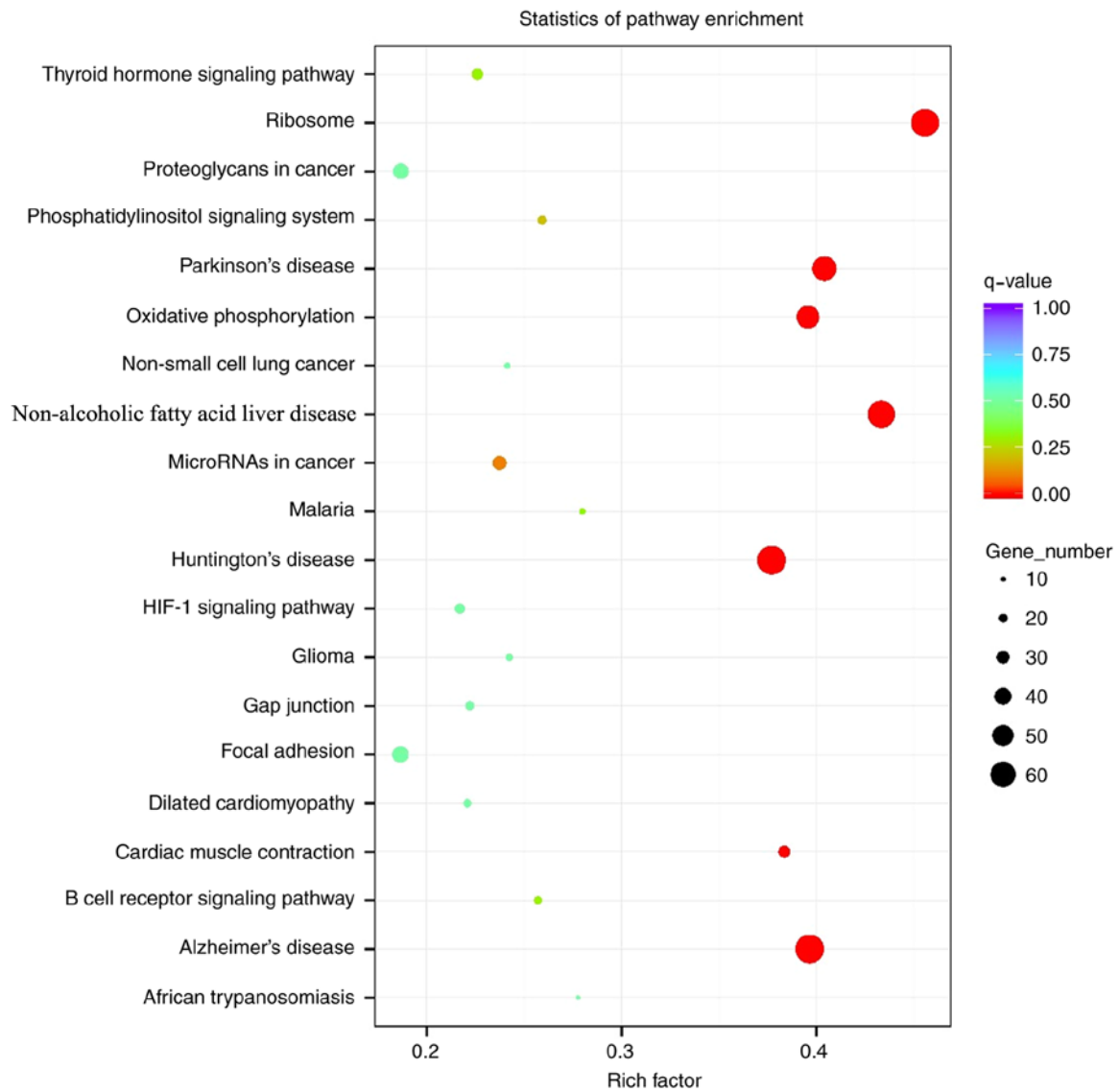


Figure 2. Kyoto Encyclopedia of Genes and Genomes enrichment pathways involved in the different stages of osteogenesis. The size of the solid circle represents the number of genes and the depth of the color represents the size of the q-value. HIF-1, hypoxia-inducible factor-1.

as 38,443,778 (81.81%), 42,956,903 (84.9%) and 41,701,352 (84.94%; Table IV).

**Functions of differentially expressed genes.** The sequencing data revealed that 1,539 (HAG-4w vs. HAG/hPMSC-4w), 1,187 (HAG-8w vs. HAG/hPMSC-8w) and 30 (HAG-12w vs. HAG/hPMSC-12w) genes that were expressed differently between the control and treated groups were related to osteogenesis responses. This was due to HAG scaffold-based delivery of hPMSCs under a double-screening using q-value (adjusted P-value,  $\leq 0.05$ ) and normalized log2ratio ( $|\log_2 \text{ratio}| \geq 1$ ; Tables SI-SIII) according previous sequencing methods (22).

Analysis of Kyoto Encyclopedia of Genes and Genomes (KEGG) enrichment demonstrated that the signal pathways enriched with differentially expressed genes differed during the all periods after implantation (Fig. 2). Fig. 2 summarizes all of the KEGG pathways, rather than the differences at each particular stage. At 4 weeks following implantation, the involved signaling pathways included 'extracellular matrix

(ECM)-receptor interaction' and 'PI3K-AKT signaling', compared with 'hematopoietic cell lineage', 'focal adhesion', 'cell adhesion molecules' (CAMs), 'protein digestion and absorption', 'platelet activation', 'B cell receptor signaling pathway' and 'osteoclast differentiation', which were involved at 8 weeks following implantation. At 12 weeks following implantation, the signaling pathways involved mainly included 'protein digestion and absorption', 'ECM-receptor interaction', 'AMP-activated protein kinase (AMPK) signaling' and 'osteoclast differentiation'.

**Detection and verification of osteogenic genes.** To validate the results of the Illumina sequencing, the osteogenic genes *COL-1* and *RUNX2* were selected for RT-qPCR analysis. The results revealed that *COL-1* (Fig. 3A) and *RUNX2* (Fig. 3B) genes were expressed in HAG and HAG/hPMSC groups during heterotopic osteogenesis and that the expression increased with time. The gene expression in the HAG/hPMSC group was markedly upregulated compared with in the HAG group at all time points (Fig. 3). The results of western blot

Table III. Statistics of mRNA sequencing reads.

Sample	Raw reads	Clean reads	Clean bases, G	Error rate, %	Q20, %	Q30, %	GC content, %
HAG-4w	60,674,628	57,642,840	8.65	0.02	96.65	91.59	50.29
HAG/hPMSC-4w	49,398,934	46,993,014	7.05	0.02	96.88	92.03	50.79
HAG-8w	53,302,412	50,746,542	7.61	0.02	96.96	92.26	51.79
HAG/hPMSC-8w	53,012,698	50,596,524	7.59	0.02	96.83	91.94	50.55
HAG-12w	48,165,136	45,911,572	6.89	0.02	96.55	91.37	51.01
HAG/hPMSC-12w	51,462,894	49,097,376	7.36	0.02	96.81	91.88	50.58

HAG, hydroxyapatite with a grooved structure; hPMSC, human placenta-derived mesenchymal stem cells; HAG/hPMSCs, HAG scaffold-based delivery of hPMSCs; w, weeks; Q20, the error probability for the identified bases during sequencing is 1%; Q30, the error probability for the identified bases during sequencing is 0.1%.

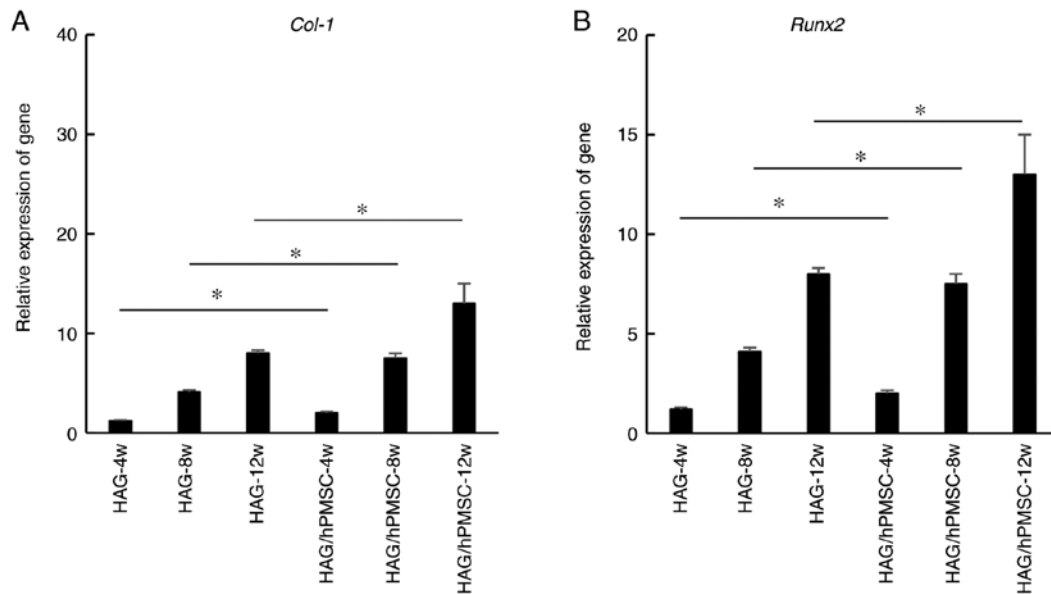


Figure 3. Relative expression of osteogenesis-associated genes. (A) Relative gene expression of *Col-1* at 4, 8 and 12 weeks after implantation of two different scaffolds. (B) Relative gene expression of *Runx2* at different times after implantation of two different scaffolds. \* $P < 0.05$ . *Col-1*, collagen-1; *Runx2*, runt-related transcription factor 2; HAG, hydroxyapatite with a grooved structure; hPMSC, human placenta-derived mesenchymal stem cells; HAG/hPMSCs, HAG scaffold-based delivery of hPMSCs; w, weeks.

analysis revealed that the expression levels of proteins COL-1 (Fig. 4A and B) and RUNX2 (Fig. 4A and C) were similar to those of the respective genes, but that the difference was not marked between each of the experimental groups.

## Discussion

Tissue-engineered bone consists of seed cells, scaffolds and growth factors. Of the seed cells, bone marrow MSCs, embryonic stem cells and adipose stem cells are the most widely studied. Biomaterials with co-cultured stem cells exhibit good biocompatibility for increasing osteogenesis and thus offer potential as bone graft substitutes (23,24). The seed cells used in the present study are MSCs derived from human placenta, which is an ideal source of seed cells for humans (25). To provide the growth factors required for tissue engineering, seed cells in the present study were co-cultured with porous scaffolds to explore whether this method effectively improved osteogenesis. The experimental

results revealed that the scaffolds in the HAG/hPMSC group demonstrated improved heterotopic osteogenesis in beagle dorsal muscles. However, in contrast to other engineered cells, the stem cells used in the present study are derived from humans, allowing for more valuable application in future clinical use. The human-derived stem cells used in the present study provide the necessary growth factors for osteogenesis.

At 4 weeks following implantation, bone-like structure was also observed on the surface of the scaffold. Compared with canine stem cell-seeded collagen-hydroxyapatite scaffolds, the HAG/hPMSC scaffold used in the present study exhibited an advantage in promoting osteogenesis at 4 weeks compared with 3 months (26). Analysis of KEGG enrichment revealed that ECM-receptor interactions (27) were involved in the process of osteogenesis. Furthermore, the PI3K-AKT pathway, which inhibits apoptosis and increases cell survival rate (28), may have improved the surface compatibility between the scaffold and the muscle cells. At 8 and 12 weeks

Table IV. Results of reads compared with the reference genome.

Sample	Total reads	Total mapped (%)	Multiple mapped (%)
HAG-4w	57,642,840	48,759,095 (84.59)	1,043,484 (1.81)
HAG/hPMSC-4w	46,993,014	38,443,778 (81.81)	885,270 (1.88)
HAG-8w	50,746,542	43,082,474 (84.9)	866,880 (1.71)
HAG/hPMSC-8w	50,596,524	42,956,903 (84.9)	844,023 (1.67)
HAG-12w	45,911,572	38,758,173 (84.42)	701,480 (1.53)
HAG/hPMSC-12w	49,097,376	41,701,352 (84.94)	742,653 (1.51)

HAG, hydroxyapatite with a grooved structure; hPMSC, human placenta-derived mesenchymal stem cells; HAG/hPMSCs, HAG scaffold-based delivery of hPMSCs; w, weeks.

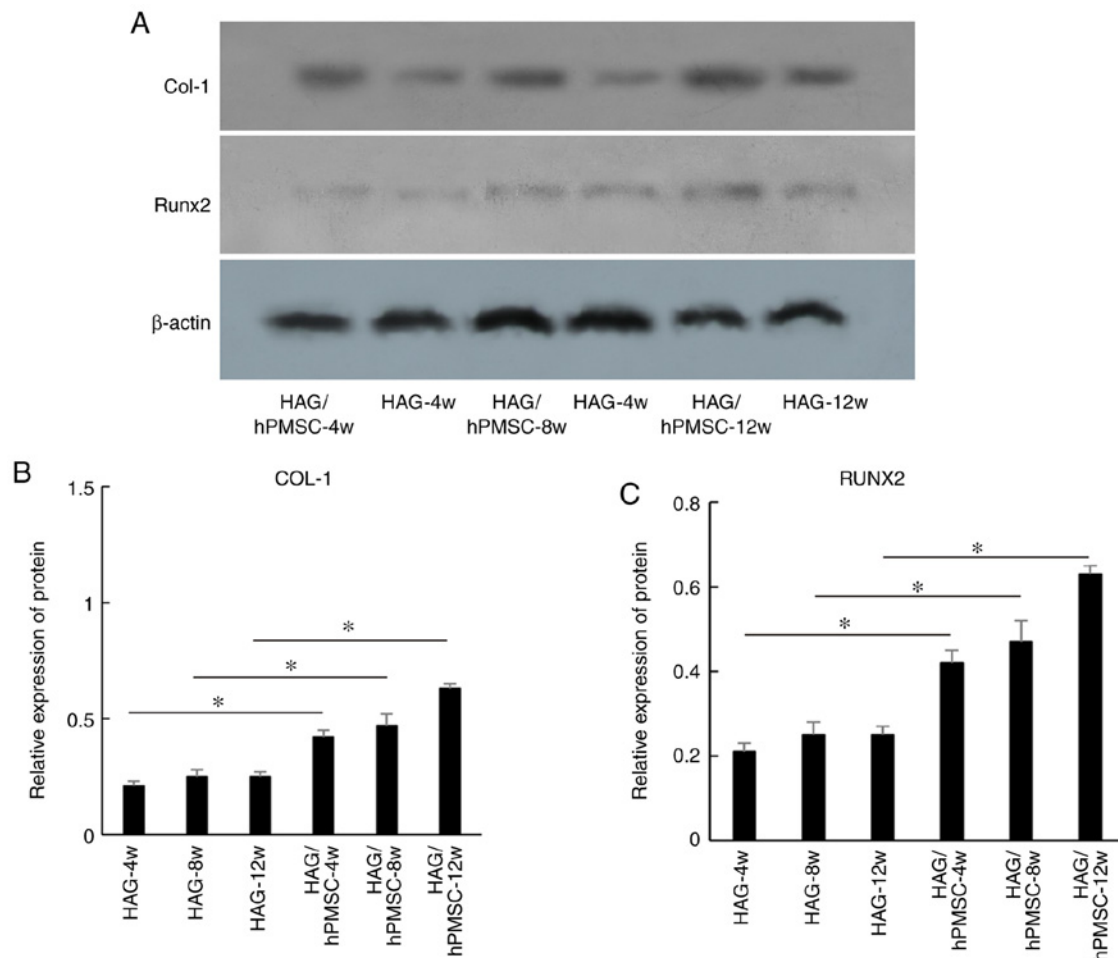


Figure 4. Relative expression of osteogenesis-associated proteins. (A) Relative protein expression of Col-1 and RUNX2 proteins at 4, 8 and 12 weeks after implantation of two different scaffolds. (B) Statistical analysis of the relative protein expression of Col-1. (C) Statistical analysis of the relative protein expression of Runx2. \*P<0.05. Col-1, collagen-1; Runx2, runt-related transcription factor 2; HAG, hydroxyapatite with a grooved structure; hPMSC, human placenta-derived mesenchymal stem cells; HAG/hPMSCs, HAG scaffold-based delivery of hPMSCs; w, weeks.

following implantation, the areas of bone matrix and mature bone plate in the HAG/hPMSC group were larger than those in the control group, suggesting that hPMSCs may continue to play an important role in the whole process of osteogenesis. At 8 weeks following implantation, pathways associated with cell adhesion, hematopoietic cell lineage, focal adhesion and CAMs played important roles, and osteoclast differentiation was activated. At 12 weeks following implantation,

osteogenesis was closely associated with protein digestion and absorption, ECM-receptor interactions, the AMPK signaling pathway and osteoclast differentiation. The AMPK signaling pathway plays a vital role in osteoclast differentiation and aids in improving osteogenesis (21,29). In the present study, differences in the signaling pathways involved in the different stages of osteogenesis were revealed, which are novel findings.



COL-1 is the main type of collagen secreted by osteoblasts, accounting for ~90% of the components of the ECM (30). The expression of COL-1 is upregulated following the differentiation of osteoblasts (30). *COL-1* is a marker gene for the phenotypic identification and osteogenic differentiation of osteoblasts (31). The results of the present study revealed that the expression of COL-1 increased with time, indicating that bone formation also increased with time. RUNX2, a transcription factor belonging to the Runt family, plays a key role in bone development (32). RUNX2 regulates osteoblast differentiation and osteogenesis during osteogenic differentiation, and the expression level of RUNX2 can also indicate the function and differentiation of osteoblasts (33). The results of the current study are consistent with the qPCR results of a previous study (5). In the present study, the expression level of RUNX2 increased with time, suggesting that osteoblasts continued to differentiate and drive bone formation. However, the present study is limited by the use of only two markers and an increase in the number of marker genes is required for further studies.

In conclusion, the present study preliminarily explored the effect of ectopic osteogenesis of hPMSCs combined with porous HAG scaffolds *in vivo*. The results of the present study provide the potential for novel applications of this cell type in osteogenesis research.

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### Availability of data and materials

The datasets generated and/or analyzed during the current study are available in the NCBI SRA repository (<https://ncbi-insights.ncbi.nlm.nih.gov/tag/sra/>) with accession numbers SRR13236657, SRR13236658, SRR13236659, SRR13236660, SRR13236661 and SRR13236662.

### Authors' contributions

YM conceived and designed the study. XR, QW, CL, QZ, JZ and HX performed the experiments. QW wrote the paper. XR, KT and YM revised the manuscript and gave final approval of the version to be published. KT reanalyzed the data. YM, QW and KT confirmed the authenticity of all the raw data. All authors have read and approved the final manuscript.

### Ethics approval and consent to participate

All animal experiments were conducted according to the protocols approved by the Animal Care and Use Committee of Sichuan Provincial People's Hospital [Sichuan, P.R. China; approval no. 2019NSF(98)].

### Patient consent for publication

Not applicable.

### Competing interests

The authors declare that they have no competing interests.

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