Abstract. In recent years, mesenchymal stem cells (MSCs), which possess the ability to specifically home to tumor sites, with the potential of multi-directional differentiation and low immunogenicity, have been reported to inhibit the growth of various types of tumors. In the present study, we isolated MSCs from the rib perichondrium (PMSCs). By comparing PMSCs with bone marrow-derived mesenchymal stem cells (BMSCs), we demonstrated that PMSCs present biological characteristics similar to those of BMSCs. Furthermore, we explored the effect and antitumor mechanism of PMSCs in rat SHZ-88 breast cancer cells. The growth, migration and invasion of the SHZ-88 cells were significantly inhibited, and the Wnt/β-catenin pathway and its target genes were down-regulated in the SHZ-88 cells by PMSC-conditioned medium. The expression level of dickkopf-1 (DKK-1) was higher in the PMSCs than that noted in the SHZ-88 cells. Neutralization of DKK-1 in the PMSC-conditioned medium attenuated the inhibitory effects of PMSCs on SHZ-88 cells. Therefore, PMSC-secreted DKK-1 is involved in the inhibition of SHZ-88 cell growth, migration and invasion, via the Wnt/β-catenin signaling pathway. In addition, we demonstrated that PMSCs inhibited the growth of breast cancer in vivo and prolonged the survival time of tumor-bearing rats. PMSCs inhibited the growth of transplanted breast tumors through the Wnt/β-catenin signaling pathway. In conclusion, our data confirmed that MSCs derived from the perichondrium present biological characteristics similar to those of BMSCs and inhibit the growth of breast cancer cells through the Wnt/β-catenin signaling pathway in vitro and in vivo. DKK-1 secreted by PMSCs played a vital role in controlling the Wnt/β-catenin signaling pathway in breast cancer.

Introduction

Mesenchymal stem cells (MSCs), known as pluripotent stem cells, were initially isolated from the bone marrow and named bone marrow-derived mesenchymal stem cells (BMSCs) (1-3). In-depth analysis has been performed for BMSCs (4,5), which exist in a wide range of tissues (6-8). Many researchers (6,8,9) believe that MSCs originate from adult stem cells. MSCs possess the ability of self-renewal, multiple differentiation potential, specific homing to tumors, and low immunogenicity (3,10,11). MSCs are being used increasingly for cancer treatment (12-14). Various reports state that MSCs can promote the progression of breast cancer and colon cancer (15,16). Other reports have demonstrated that MSCs can inhibit the growth of pancreatic cancer, Kaposi's sarcoma and breast cancer (13,17,18). In breast cancer, the effect of MSCs is controversial. Studies reported that the regulated self-renewal of stem cells mediated by the Wnt/β-catenin signaling pathway might be subverted in cancer cells to allow malignant proliferation (19,20). Likewise, the stem cell microenvirom-
ment plays an essential role in preventing carcinogenesis by inhibiting proliferation (21). MSCs were found to inhibit tumor proliferation through secretion of dickkopf-1 (DKK-1), which acts as an inhibitor of the Wnt/β-catenin signaling pathway (22).

The rat rib perichondrium contains osteoprogenitor cells, a type of adult stem cells. In the present study, we first determined whether MSCs could be isolated from the perichondrium and whether perichondrium MSCs (PMSCs) present biological characteristics similar to those of BMSCs. We next explored the effect of PMSCs on rat SHZ-88 breast cancer cells in vitro and in vivo. We also investigated whether PMSCs affect breast cancer cells through the DKK-1/Wnt/β-catenin signaling pathway. Our findings demonstrated that MSCs could be derived from the perichondrium and that they inhibited the growth of SHZ-88 breast cancer cells through the DKK-1/Wnt/β-catenin signaling pathway.

Materials and methods

Cell culture. Specific pathogen-free 4-week-old female Sprague-Dawley (SD) rats were obtained from the Experimental Animal Center of Xi'an Jiaotong University School of Medicine. All animals were cared for in accordance with the institutional guidelines for the use of experimental animals. The perichondrium was separated under a dissecting microscope and treated with 5 ml of 0.2% collagenase II (Sigma-Aldrich St. Louis, MO, USA) for 3 h in a 37°C incubator shaker. We then collected the supernatant via 200 mesh sieves and added 1 ml fetal bovine serum (FBS; HszSjq Co., Ltd., Hangzhou, China) to stop the digestion. After centrifugation at 1,200 x g for 10 min, the perichondrium cells were cultured in Dulbecco's modified Eagle's medium (DMEM)/F12 (HyClone Co., Logan, UT, USA) with 10% FBS. Bone marrow cells were isolated from femurs and tibias as previously described (8), and cultured in DMEM/F12 with 10% FBS. SHZ-88 cells were obtained from Shanghai Cell Research Institute (Shanghai, China) and cultured in Roswell Park Memorial Institute (RPMI)-1640 medium (HyClone) with 10% newborn calf serum (NBCS; HszSjq Co. Ltd.) and 100 µg/ml penicillin/streptomycin (Gibco Life Technologies, Rockville, MD, USA). All cell lines were maintained in a humidified atmosphere at 37°C with 5% CO₂ and routinely passaged using trypsin (Invitrogen, Camarillo, CA, USA) when nearly confluent.

Immunofluorescence staining. The primary to 3rd-generation perichondrium cells and BMSCs were seeded at 4x10⁴/ml in 24-well plates for 48 h. Migration and invasion assays were conducted using Transwell plates with 8-µm pore size membranes (Millipore Inc., Billerica, MA, USA) as previously described (29,30). After incubation for 16 h (for migration assays) or 24 h (for invasion assays), the slides were removed and the cells were fixed for 30 min with 4% paraformaldehyde. The lipids droplets were stained by Oil Red O (Sigma-Aldrich). The mineralization nodes were stained by 5% silver nitrate (Sigma-Aldrich) and exposed to an ultraviolet lamp within 60 min. Hematoxylin was used to stain the cell nuclei.

Cell proliferation assessment. Colorimetric 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assay was used to assess the proliferation of cells. SHZ-88 cells (100 µl) were seeded at a density of 4x10⁴ cells/ml in 500-µl wells of 96-well plates and treated with PMSC-conditioned medium (control without PMSC-conditioned medium) for 24, 48 and 72 h. The 3rd-generation PMSCs were cultured as indicated above. When the cells grew to full confluency, the medium was replaced by serum-free medium for 10% FBS. Bone marrow cells were isolated from femurs and tibias as previously described (8), and cultured in DMEM/F12 with 10% FBS. SHZ-88 cells were obtained from Shanghai Cell Research Institute (Shanghai, China) and cultured in Roswell Park Memorial Institute (RPMI)-1640 medium (HyClone) with 10% newborn calf serum (NBCS; HszSjq Co., Ltd.) and 100 µg/ml penicillin/streptomycin (Gibco Life Technologies, Rockville, MD, USA). All cell lines were maintained in a humidified atmosphere at 37°C with 5% CO₂ and routinely passaged using trypsin (Invitrogen, Camarillo, CA, USA) when nearly confluent.

Cell migration and invasion assays. After treatment of SHZ-88 cells with 50% PMSC-conditioned medium for 48 h, migration and invasion assays were conducted using Transwell plates with 8-µm pore size membranes (Millipore Inc., Billerica, MA, USA) as previously described (29,30). After incubation for 16 h (for migration assays) or 24 h (for invasion assays), the cells remaining on the upper side of the filter were removed with cotton swabs. The cells that attached to the lower surface were fixed, stained using crystal violet and washed with PBS. Cells were counted in five high-power fields/membrane. The results are presented as the mean number of cells that migrated per field per membrane. All experiments were conducted in triplicate.

SHZ-88 cell transplantation in SD rats. SHZ-88 cells in the log-phase were diluted in normal saline (NS) and were subcu-
taneously injected into the right armpit of 10-day-old SD female rats. Every rat was injected with 5x10^5 cells/0.2 ml. When subcutaneous tumors reached 5-8 mm in size, SD rats were randomly divided into PMSC and NS groups (n=16 rats/group). Animals in the PMSC group were injected with 3x10^6/0.2 ml PMSMCs (diluted in NS) via the tail vein every 3 days. Animals in the NS group were injected with 0.2 ml NS via the tail vein every 3 day as a control group. We measured the size of subcutaneous tumors every 6 days. The tumor volume was calculated using the following formula: Tumor volume = \frac{1}{2} (long tumor diameter × short tumor diameter). The tumor growth curve was drawn by plotting tumor volumes according to the time in each group. Finally, 6 rats in each group were euthanized after 30 days and tumor growth inhibitory rate was calculated using the following formula: Tumor inhibition rate = \frac{[(average volume of control group - average volume of experimental group)/average volume of control group] × 100\%}{\text{A piece of the tumor tissue from each animal was fixed in 4\% paraformaldehyde while the remaining tumor tissue was stored at 80\°C. Surviving animals (the remaining 10 animals in each group) were observed daily for 60 days.}

Western blot analyses. The proteins were extracted from the SHZ-88 cells treated with MSC-conditioned medium for 72 h and frozen tumors from the tumor-bearing rats according to standard procedures. Proteins were examined with specific primary antibodies: anti-DKK-1 rabbit polyclonal antibody (BS7731), anti-Bcl-2 rabbit polyclonal antibody (BS1031), anti-PCNA rabbit polyclonal antibody (BS6438), anti-survivin rabbit polyclonal antibody (BS8456) (1:1,000 dilution; Bioworld Technology), anti-β-catenin rabbit polyclonal antibody (S1067-2-AP) (1:500 dilution; Proteintech, Huhan, China), anti-c-Myc rabbit polyclonal antibody (10057-1-AP) (1:100 dilution; Proteintech), and anti-β-actin rabbit polyclonal antibody (bs-0061R; 1:1,000 dilution; Bioss), followed by a conjugated secondary antibody: goat-anti-rabbit IgG antibody (1:3,000 dilution; Bioss). The reactions were visualized using the enhanced chemiluminescence (ECL) reagent (Millipore). The band intensity of western blotting was measured by densitometry using the Quantity One software (Bio-Rad Laboratories, Hercules, CA, USA). The protein levels were normalized to the protein level of β-actin which was used as a loading control.

Immunohistochemical (IHC) staining. Tumor tissues fixed in 4\% paraformaldehyde were processed for paraffin embedding and sectioned (5-μm thickness). Formalin-fixed paraffin-embedded tumor sections from NS and PMSC-treated animals were analyzed by IHC staining using an anti-β-catenin rabbit polyclonal antibody (1:200; Proteintech) and a biotin-conjugated secondary antibody. The staining was performed following the SP kit procedure (Golden Bridge International, Beijing, China). As a control, the primary antibody was replaced by PBS. IHC staining results were assessed independently by two pathologists in a semi-quantitative manner, scored by a semi-quantitative immunoreactivity scoring (IRS) system, and divided into high and low expression specimens.

Statistical analysis. All data are expressed as means with standard error (SE). One-way ANOVA and Student’s t-test were used to test differences between the groups. The Kaplan-Meier method was used to test the survival time of tumor-bearing rats in different groups. P<0.05 was considered to indicate a significant result. All analyses were performed using SPSS software version 18.0 (IBM, Corp., New York, NY, USA).

Results

Biological characteristics of the PMSCs and BMSCs

Growth and morphology. The primary perichondrium cells began to adhere to the plate after 12 h of culture. They were passaged after 5-6 days of culture when they presented a round or polygon shape (Fig. 1A). The primary bone marrow cells began to adhere to the plate after 4-5 h of culture and were passaged after 8-10 days of culture when they presented a spindle or polygon shape (Fig. 1A). As the generations increased, from 3rd-generation, two types of cells showed spindle-shape with whirlpool arrangement (Fig. 1A) and the generation times of perichondrium cells and BMSCs were 1-2 and 4-5 days, respectively. The perichondrium cells were passaged for 30 generations and retained their shape (data not shown).

Cell surface marker expression. Immunofluorescence staining indicated that type II collagen, CD34, CD90 and CD105 proteins were not expressed in primary and first generation perichondrium cells. From the 2nd-generation, cells began to express CD90 and CD105 proteins. The primary bone marrow cells were weakly positive for CD34, CD90 and CD105 proteins, but not for type II collagen. Subsequently, the cells only expressed CD90 and CD105 proteins. From the 3rd-generation, the two cell types did not expressed type II collagen and CD34 proteins, but strongly expressed CD90 and CD105 proteins (Fig. 1B). The 30th generation of perichondrium cells still strongly expressed CD90 and CD105, but did not expressed CD34 and type II collagen (data not shown).

Multipotent differentiation ability. No change in term of the induction of differentiation was observed in the control group (Fig. 1C). After 14 days of adipogenic induction of the 3rd-generation cells, perichondrium cells and BMSCs became round. Lipid droplets were observed in the cytoplasm of these cells after Oil Red O staining (Fig. 1C). After 21 days of osteogenic induction, the perichondrium cells grew as a stratified layer, some black mineralized nodules stained by silver nitrate were observed on the cell surface (Fig. 1C). BMSCs grew as a simple layer, but mineralized nodules gathered as bone nodules (Fig. 1C). The 30th generation perichondrium cells still presented lipid droplets and black mineralized nodules after induction (data not shown).

PMSC-conditioned media inhibits SHZ-88 cell growth, migration and invasion. Human MSCs can inhibit tumor growth (22,28). However, no study has reported such information regarding PMSCs. In the present study, we explored the effect of PMSCs on breast cancer. The stem cell microenvironment plays an essential role in preventing carcinogenesis by inhibiting proliferation (21). Thus, we investigated the inhibitory effect of 50\% PMSC-conditioned medium on SHZ-88 cells in vitro. MTT assay showed that PMSC-conditioned medium inhibited the proliferation of SHZ-88 cells in a time-dependent manner.
The Wnt/β-catenin signaling pathway is suppressed in the SHZ-88 cells treated with PMSC-conditioned medium. Studies have indicated that the self-renewal of stem cells by the Wnt/β-catenin signaling pathway was subverted in cancer cells, resulting in tumor progression (19,20). The Bcl-2, c-Myc, PCNA and survivin genes are all targets of the Wnt/β-catenin signaling pathway. Western blot analysis showed that β-catenin, Bcl-2, c-Myc, PCNA and survivin proteins were downregulated in SHZ-88 cells treated with 50% PMSC-conditioned medium (P<0.001; Fig. 3). These results indicated that some soluble exocrine factors in the MSC-conditioned medium are involved in the inhibition of the Wnt/β-catenin signaling pathway in SHZ-88 cells.

DKK-1 derived from PMSCs contributes to the inhibition of SHZ-88 cells. MSCs have been shown to inhibit tumor proliferation by secreting DKK-1, which acts as an inhibitor of the Wnt/β-catenin signaling pathway (28,31). Thus, we determined the expression levels of DKK-1 in the SHZ-88 cells and PMSCs. Western blot analysis indicated that the
DKK-1 expression level was higher in the PMSCs than that in the SHZ-88 cells (P<0.001; Fig. 4A). DDK-1 neutralization by the rabbit antibody against the rat DKK-1 in the PMSC-conditioned medium abolished the inhibitory effect of the PMSC-conditioned medium on breast cancer cell migration (B) and invasion (C). The results are presented as the mean ± SE of cell numbers obtained in three independent experiments (**P<0.001).

Figure 3. Effect of PMSC-conditioned medium on the Wnt/β-catenin signaling pathway of tumor cells in vitro. Western blot analyses showed that β-catenin, Bcl-2, c-Myc, PCNA and survivin were downregulated in the SHZ-88 cells treated with 50% PMSC-conditioned medium (**P<0.001).

Figure 2. Inhibitory effect of PMSC-conditioned medium on SHZ-88 cells. (A) MTT assay showed that the growth of SHZ-88 cells was reduced in the presence of 50% PMSC-conditioned medium for 72 h (P<0.001). (B and C) Cell Transwell assays were conducted to investigate the effect of PMSC-conditioned medium on breast cancer cell migration (B) and invasion (C). The results are presented as the mean ± SE of cell numbers obtained in three independent experiments (**P<0.001).

Migration and invasion of SHZ-88 cells was also lost (P>0.05; Fig. 4C and D). Moreover, western blot analysis indicated that neutralization of DKK-1 in the PMSC-conditioned medium abrogated the downregulating effect of the conditioned medium on the expression of β-catenin, Bcl-2, c-Myc, PCNA and survivin in SHZ-88 cells (P>0.05; Fig. 4E). These data indicated that the inhibition of SHZ-88 cell growth, migra-
PMSCs inhibit the growth of breast cancer in vivo

PMSCs inhibit the growth of transplanted tumors. To determine the effect of PMSCs on transplanted tumors in vivo, we established a subcutaneous tumor model of breast cancer in SD rats. When the tumor nodules reached 5-8 mm in length, the rats were injected with either PMSCs or NS (control group). Tumor volume gradually increased and the tumor growth rate was faster in the NS group than that determined in the PMSC group. On day 30, no necrosis was observed in the tumors from rats treated with PMSCs, while partial necrosis and a hemorrhagic tendency on the surface of tumors were observed in the tumors from rats treated with NS. Additionally, the size of the tumors in the NS group was larger than that of the tumors from the PMSC group (Fig. 5A). We measured the size of the subcutaneous tumors every 6 days and constructed the tumor growth curve based on tumor volumes (Fig. 5A). The tumor inhibition rate in the PMSC group reached 62.8%. These data indicated that treatment of the rats with PMSCs resulted in a significant growth inhibition in vivo when compared with the control group (P<0.001). The mean survival time of the rats in the NS group was 39.7±1.2 days, while that of the rats in the PMSC group was 53.4±2.0 days. The survival time of the rats in the NS group was significantly shorter than that of the rats treated with PMSCs (P<0.001; Fig. 5B).

PMSCs inhibit the growth of transplanted tumors via the Wnt/β-catenin signaling pathway. We further investigated the molecular mechanisms underlying the inhibitory effect of PMSCs on breast cancer. Immunohistochemistry staining indicated that the β-catenin protein in the tumors from rats treated with PMSCs was expressed at low levels, while it was highly expressed in the tumors of rats treated with NS when using the IRS system (Fig. 5C). Western blot analyses showed that β-catenin, Bcl-2, c-Myc, PCNA and survivin expression in SHZ-88 cells was attenuated by DKK-1 neutralization.
transplanted tumors via the Wnt/β-catenin signaling pathway in vivo.

**Discussion**

Many researchers (8,10,14) believe that MSCs originate from adult stem cells. The rib perichondrium contains osteo-progenitor cells, a type of adult stem cells. In the present study, we investigated whether MSCs could be isolated from the perichondrium. Our results showed that, from the 3rd-generation, the perichondrium cells presented a typical spindle-shape with whirlpool arrangement and strongly expressed CD90 and CD105 proteins (32). Additionally, adipogenic and osteogenic differentiation could be induced in these cells. These data demonstrated that perichondrium cells presented biological characteristics similar to those of BMSCs. These findings strongly suggest that we successfully derived stem cells from the rib perichondrium, which were named PMSCs.

MSCs possess the ability to specifically home to tumors and to self-renew, and present low immunogenicity (3,10,11,16). Thus, they have been used for cancer treatment (13,33,34). However, the effects of MSCs on tumors are controversial. Various reports state that MSCs promote the progression of breast cancer and colon cancer cells (15,16). Other reports indicate that MSCs can inhibit the growth of pancreatic cancer, Kaposi's sarcoma, and breast cancer cells (13,17,18). In the present study, using MTT, cell migration and invasion assays, we found that PMSC-conditioned medium inhibited the growth, migration and invasion of SHZ-88 cells (P<0.001). Our findings are consistent with several reports (18,35). The Wnt/β-catenin signaling pathway plays an important role in the growth and invasion of cancer cells.
in the self-renewal of stem cells (19), but abnormal activation of the Wnt/β-catenin signaling pathway can lead to human cancer (20). The Wnt/β-catenin pathway is related to the efficacy of MSCs in suppressing hepatoma and breast cancer (18,28,35). The Bcl-2, c-Myc, PCNA and survivin genes are all targets of the Wnt/β-catenin signaling pathway (36,37).

In the present study, western blot analyses confirmed that the Wnt/β-catenin signaling pathway in SHZ-88 cells was blocked by PMSC-conditioned medium. These results are consistent with the above report. These data suggest that various soluble factors in PMSC-conditioned medium are responsible for the inhibition of the growth, migration, invasion and Wnt/β-catenin signaling pathway in SHZ-88 cells.

Next, we attempted to identify the inhibitors of the Wnt/β-catenin signaling pathway in the PMSC-conditioned medium. It was previously reported that MSCs inhibit tumor proliferation via secretion of DKK-1, which acts as a negative regulator of the Wnt/β-catenin signaling pathway (22,28). Accordingly, we speculated that DKK-1 secreted by PMSCs was involved in the inhibition of the Wnt/β-catenin signaling pathway in SHZ-88 cells. Western blot analyses showed that DKK-1 expression level in PMSCs was higher than that in SHZ-88 cells, which provided evidence that DKK-1 from PMSCs may play a vital role in controlling the Wnt/β-catenin signaling pathway in SHZ-88 cells. These findings are consistent with various reports (18,22,28). In order to further confirm the role of DKK-1 secreted by PMSCs in the inhibition of the Wnt/β-catenin signaling pathway in SHZ-88 cells, we neutralized DKK-1 in the PMSC-conditioned medium using an antibody against DKK-1 and demonstrated that the inhibition of SHZ-88 cell growth, migration and invasion mediated by PMSC-conditioned medium was reduced by DKK-1 neutralization. Additionally, the treated conditioned medium lost the ability to downregulate the expression of β-catenin, Bcl-2, c-Myc, PCNA and survivin in SHZ-88 cells. This provided evidence that DKK-1 secreted by PMSCs contributed to the inhibition of the Wnt/β-catenin signaling pathway in SHZ-88 cells. Moreover, we showed that PMSCs inhibited the growth of transplanted tumors in vivo. When compared with the NS group, injection of PMSCs significantly inhibited tumor growth and prolonged the survival time of tumor-bearing rats. Wnt/β-catenin and its target genes were downregulated in the tumors from rats treated with PMSCs. These results indicate that PMSCs inhibit the growth of transplanted breast tumors through the Wnt/β-catenin signaling pathway as observed in vivo.

In summary, the present study is the first to report the isolation of mesenchymal stem cells from the perichondrium (PMSCs) of rat. Furthermore, PMSCs were found to inhibit breast cancer cell growth through the DKK-1/Wnt/β-catenin signaling pathway. The present study provided novel information, which can be useful for the development of new therapeutic agents for breast cancer treatment.

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