

# MicroRNA-124-3p inhibits the differentiation of precartilaginous stem cells into nucleus pulposus-like cells via targeting FSTL1

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**Abstract.** MicroRNA (miRNA/miR)-124-3p has been extensively studied in tumor biology and stem cells. However, little is known regarding its functional roles in the differentiation of precartilaginous stem cells (PSCs) into nucleus pulposus-like cells (NPLCs). In the present study, using miRNA microarray screening, it was demonstrated that the miRNA expression profiles differed between rat primary PSCs and TGF- $\beta$ 1-induced differentiated NPLCs, and that miR-124-3p was significantly differentially expressed during the differentiation of PSCs to NPLCs. Furthermore, RT-qPCR analysis verified that miR-124-3p expression was decreased during PSC differentiation, with the lowest levels being detected at the later stages. Subsequent experiments revealed that miR-124-3p overexpression significantly decreased the expression of the extracellular matrix proteins, aggrecan and collagen type II, which was accompanied by a significant decrease in follistatin-related protein 1 (FSTL1) expression levels. Moreover, bioinformatics analysis indicated that FSTL1 was a potential target of miR-124-3p, which was additionally verified using luciferase reporter assays. Taken together, these data revealed a specific regulatory pathway of miR-124-3p, which negatively regulated its target gene, FSTL1, during the differentiation of PSCs to NPLCs, and suggested a functional role for miR-124-3p in the differentiation of PSCs.

## Introduction

Degeneration of the intervertebral discs is characterized by alterations in the morphology of the discs and the composition of the extracellular matrix, including a reduction in the number of the intervertebral disc cells (1,2). It has been reported that when the intervertebral disc degenerates, the extracellular

matrix is reduced, resulting in a decrease in the water content of the intervertebral disc, thereby resulting in the degeneration of the intervertebral disc structure (1-4).

Tissue engineering has been increasingly employed in the medical field as a treatment for degenerative discs, with cell therapy becoming a potential treatment for intervertebral disc degeneration (5). Previous studies have reported that various types of cells, including nucleus pulposus cells, chondrocytes and mesenchymal stem cells (MSCs), can be used for cell therapy (6-8). With further progress, stem cell therapy has been applied in the clinic, with several studies reporting these results. For example, Centeno *et al* (9) reported that patients with degenerative disc disease (DDD) who are treated with MSCs to counteract lower back pain with radicular symptoms, exhibit minor adverse effects and considerable improvements in the degree of pain, function and overall quality of life during a 6 year follow-up. Li *et al* (10) investigated the characteristics of stem cells, including nucleus pulposus-derived stem cells (NPSCs), annulus fibrosus-derived stem cells (AFSCs) and cartilage endplate-derived stem cells (CESCs), in human degenerative intervertebral discs to determine the best stem cell-like characteristics. No significant differences in cell morphology among NPSCs, AFSCs and CESCs have been revealed; however AFSCs have been indicated to exhibit the best stem cell-like characteristics in the human degenerative intervertebral disc (10).

The present study focused on another type of stem cell, namely precartilaginous stem cells (PSCs). It has been indicated that PSCs are localized in the perichondrium surrounding the epiphysis (11). PSCs are adult stem cells with a multi-directional differentiation potential, which have been reported to differentiate into chondrocytes and osteoblasts to promote the growth of animal limbs (12). PSCs have been isolated and purified via immunomagnetic bead sorting, and used as seed cells in several tissue engineering studies (12). The discovery of PSCs has provided a novel tool for cell transplantation to repair degenerative intervertebral discs (13). PSCs have been indicated to serve an important role in cartilage growth and internalization, as well as articular cartilage injury repair (14). PSCs are precursor cells of chondrocytes and exhibit a homology with intervertebral disc nucleus cells. The alterations in nucleus pulposus cells have been reported to serve an important role in the degeneration of intervertebral discs (15). The nucleus pulposus is an avascular

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tissue with a limited number of cells and it is difficult to repair its structure following the occurrence of a lesion (16). Therefore, various studies have investigated the use of seed cells to repair degenerated nucleus pulposus cells (17-19). The nucleus pulposus cells and chondrocytes of the intervertebral disc have been reported to exhibit numerous common characteristics, such as production of collagen II and aggrecan as principal components of the extracellular matrix (18,19). Moreover, nucleus pulposus-like cells (NPLCs), which are derived from the degenerated intervertebral disc, have been indicated to undergo osteogenesis and promote cartilage repair owing to the fact that these cells secrete factors (TGF-1, IL-1, TNF, prostaglandin E2, IL-10, and granulocyte-macrophage colony stimulating factor) which promote the proliferation and regulate the differentiation of chondrocytes, as well as promote the synthesis of extracellular matrix components (20).

Our previous study indicated that PSCs can be induced to differentiate to NPLCs to repair degenerative intervertebral discs with unique therapeutic advantages (21). However, the detailed mechanisms of action behind the differentiation are still unclear. MicroRNAs (miRNAs/miRs) are a class of small noncoding RNA molecules that negatively regulate gene expression at the post-transcriptional level (22). It has been reported that miRNAs not only regulate several normal physiological processes but are also associated with the development of DDD (23). In the present study, miRNA microarray screening was employed to identify differences in miRNA expression profiles between TGF- $\beta$ 1-induced rat primary PSCs and differentiated NPLCs. miR-124-3p was verified as a significantly differentially expressed miRNA during the differentiation of PSCs to NPLCs. Moreover, the mechanisms of action through which miR-124-3p regulates the fate of PSCs and its role in the differentiation process were also investigated.

## Materials and methods

**Chemicals, reagents and antibodies.** TGF- $\beta$ 1 was purchased from Selleck Chemicals. miR-124-3p and negative control (NC/nc) agomirs (cat. no. miR40000422-4-5) were obtained from Guangzhou RiboBio Co., Ltd. Primary antibodies against fibroblast growth factor receptor 3 (FGFR-3; cat. no. ab133644), follistatin-related protein 1 (FSTL1; cat. no. ab232777), collagen II (cat. no. ab188570), aggrecan (cat. no. ab3778) and HRP-conjugated secondary antibody GAPDH (cat. no. ab181602) were obtained from Abcam. FSTL1 small interfering (si)RNA and siRNanc sequences were designed, synthesized and validated by Shanghai GenePharma Co., Ltd. The siRNA sequences used were as follows: FSTL1, 5'-GCAAUACUUACGGACUUU-3'; and siRNanc, 5'-GCAUCUUGAGAUUUAUCA-3'. FSTL1 (cat. no. HP100806), TPST2 (cat. no. HP101143) and enhancer of zeste homolog 2 (EZH2; cat. no. HP101222) quantitative PCR (qPCR) primers were purchased from Sino Biological Inc. (Shanghai, China). COS7 and 293 cells were purchased from the Cell Bank of the Chinese Academy of Sciences.

**Primary PSCs culture.** PSCs were generated as previously described (11). Briefly, the perichondrial mesenchyme (the rings of LaCroix) were isolated from Sprague Dawley rats (Trophic Animal Feed High-Tech Co., Ltd.) and subsequently digested

in 0.25% Trypsin Solution (cat. no. 25200072; Thermo Fisher Scientific, Inc.) supplemented with 0.05% collagenase type I (Sigma-Aldrich; Merck KGaA) for 3-5 min at 37°C. The digestion mixture was resuspended in FBS. Subsequently, the cells were dispersed and resuspended as a single cell suspension in 0.1 M PBS with centrifugation speeds of 167.7 x g for 5 min at room temperature. The cell suspension was incubated with the FGFR-3 antibody (1:500; cat. no. ab133644; Abcam) for 15 min at 4°C, and subsequently, an immunomagnetic separation system (Miltenyi Biotec GmbH) was used to purify the FGFR-3 expressing cells. The isolated PSCs were cultured in complete DMEM/F12 medium supplemented with 20% FBS (Nanjing KeyGen Biotech Co., Ltd.) at 37°C with 5% CO<sub>2</sub>. The phenotype identification of the rat PSCs was based on the FGFR-3 protein expression, which was detected via western blotting. The present study was approved by the Institutional Animal Care and Use Committee of Wuxi People's Hospital (approval no. WXP20190103c0600105). In the current study, 60 rats were used between January and March 2019. Animal health and behavior were monitored daily. All operations were performed according to international guidelines for the care and treatment of experimental animals. The rats were male Sprague-Dawley rats of 5-7 weeks of age weighing 150-200 g. Cages were cleaned and enrichment items were renewed weekly. The animal room had a controlled 12-h light/dark cycle (lights on at 6:00 AM), temperature (22±2°C) and relative humidity (45-65%). Daily food and water were supplied by laboratory staff. The rats were maintained in an ordinary housing facility, which is in accordance with the national standard 'Laboratory Animal-Requirements of Environment and Housing Facilities' (standard no. GB 14925-2010). The animals were euthanized in accordance with the requirements of the 'Guidelines for the Examination of the Scientific Research of Experimental Animal Welfare' (standard no. GB/T 35892-2018) using intravenous administration of pentobarbital (100-150 mg/kg) and death was verified via examining the animals' heartbeat for complete cessation and pupil dilation.

**PSCs cell differentiation.** PSCs were differentiated according to a previously published protocol (24). Briefly, PSCs at passage three were trypsinized and transferred to six-well plates at a concentration of 1x10<sup>5</sup> cells/ml. Differentiation of PSCs was induced by culturing the cells in DMEM/F12 medium supplemented with 10% FBS, 10 ng/ml TGF- $\beta$ 1, 100 nM dexamethasone, 50  $\mu$ g/ml L-ascorbic acid 2-phosphate, 100  $\mu$ g/ml sodium pyruvate, 40  $\mu$ g/ml proline and 6.25 mg/l insulin at 37°C with 5% CO<sub>2</sub> for 7 days.

**Cell lines and culturing conditions.** COS-7 and 293 cells were cultured in DMEM (Gibco; Thermo Fisher Scientific, Inc.) supplemented with 10% fetal bovine serum (FBS; Gibco; Thermo Fisher Scientific, Inc.) and maintained in an incubator at 37°C and 5% CO<sub>2</sub>. The medium was replaced 2-3 times a week. When the cell density reached 70-80%, cells were digested by 0.25% trypsin (Gibco; Thermo Fisher Scientific, Inc.) and passaged.

**Cell transfection.** Small interfering RNA targeting FSTL1 (siFSTL1) was synthesized by Guangzhou RiboBio Co., Ltd., and a scrambled siRNA (siNC; Guangzhou RiboBio Co., Ltd.)

was used as the negative control. One day before transfection, cells were digested with 0.25% trypsin (cat. no. C0202; Beyotime Institute of Biotechnology). Then, 50  $\mu$ l of Opti-MEM<sup>®</sup> medium (cat. no. 31985062; Invitrogen; Thermo Fisher Scientific, Inc.) was mixed with siFSTL1 (cat. no. siB14715103107-1-5; Guangzhou RiboBio Co., Ltd.) or siNC. And 3  $\mu$ l Lipofectamine<sup>®</sup> 3000 reagent (cat. no. L3000015; Thermo Fisher Scientific, Inc.) was diluted with 50  $\mu$ l OptiMEM. Then, the above two mixtures were mixed. Then cells were transfected with the mixture according to the Lipofectamine<sup>®</sup> 3000 kit (Thermo Fisher Scientific, Inc.).

miR-124-3p agomir (miR40000422-4-5 Guangzhou RiboBio Co., Ltd.) and negative control (agomir NC) were synthesized by Guangzhou RiboBio Co., Ltd. Cells were seeded in six-well plates ( $1 \times 10^5$  cells/ml) and transfected with 50 nM miR-124-3p agomir or 50 nM agomir NC using Lipofectamine<sup>®</sup> 3000 kit (Thermo Fisher Scientific, Inc.). The transfection efficiency was observed by a fluorescence microscope after 24 h.

**miRNA microarray analysis.** Total RNA from rat primary PSCs and TGF- $\beta$ 1-differentiated NPLCs was isolated using the TRIzol<sup>®</sup> reagent (Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer's instructions. Following quantification of total RNA using NanoDrop ND-1000 Ultraviolet Spectrophotometer (NanoDrop Technologies; Thermo Fisher Scientific, Inc.), a miRNA profiling system (Agilent Technologies, Inc.) was used to detect the differences in miRNA expression between primary PSCs and NPLCs. The slides were scanned using a microarray scanner (GenePix 4100A) and the data were quantified using Feature Extraction software v10.7 (both from Agilent Technologies, Inc.). The GeneSpring GX software v12.6 (Agilent Technologies, Inc.) was additionally used to analyze the miRNA expression data that were detected using the microarray system.

**Reverse transcription-quantitative (RT-q) PCR.** Total RNA of PSCs was extracted using TRIzol<sup>®</sup> reagent (Invitrogen; Thermo Fisher Scientific, Inc.). The primers used for the amplification of miR-124-3p (cat. no. miRA1001726-1-100) and U6 endogenous control (cat. no. miRAN0002-1-100) were purchased from Guangzhou RiboBio Co., Ltd. Subsequently, qPCR was performed using the Mir-X<sup>™</sup> miRNA RT-qPCR TB Green<sup>®</sup> kit according to the manufacturer's instructions (Takara Biotechnology). Briefly, Total RNA was reverse transcribed to obtain cDNA. The reverse transcription process of miR-124-3p used the Mir-X miRNA First-Strand Synthesis Kit (cat. no. 638313; Takara Biotechnology, Co., Ltd.), the tube was incubated for 1 h at 37°C, and the reaction terminated at 85°C for 5 min to inactivate the enzymes. The PCR reactions were carried out using TB Green Advantage qPCR Premix (cat. no. 639676; Takara Biotechnology, Co., Ltd.) under the following conditions: Denaturation, 95°C 10 sec; qPCR x 40 cycles of 95°C 5 sec and 60°C 20 sec; dissociation curve, 95°C 60 sec, 55°C 30 sec and 95°C 30 sec. The relative miRNA expression was calculated using the  $2^{-\Delta\Delta C_t}$  method (25).

**Target gene prediction.** To predict the potential target genes of miRNA-124-3p, three bioinformatics analysis tools, TargetScan v7.2 ([www.targetscan.org](http://www.targetscan.org)), Pictar (<https://pictar.mdc-berlin.de/>)

and miRanda (<http://www.microrna.org/microrna/home.do>) were used to predict its potential target genes.

**Luciferase reporter assay.** For the luciferase reporter assay, a construct containing the 3'-untranslated region (UTR) of the FSTL1 mRNA carrying the putative miRNA-124-3p binding site or a respective mutant, which was used as a control, were cloned into the pGL3 Luciferase Reporter plasmid (Promega Corporation). Cells ( $1 \times 10^5$ ) were co-transfected with a 50 ng/well reporter plasmid and 50 nM miR-124-3p agomir or 50 nM agomir NC in a 6-well plate for 48 h. Firefly and *Renilla* luciferase activities were detected using Dual-Luciferase<sup>®</sup> Reporter Assay System (cat. no. E1910; Promega Corporation). Briefly, 500  $\mu$ l of PLB was added and agitated for 15 min. LAR II (100  $\mu$ l) and cells (20  $\mu$ l) were sequentially added to 96-well plate. Then, 100  $\mu$ l of 1X Stop & Glo reagent was added to detect the luminescence intensity of *Renilla* luciferase.

**Western blot analysis.** Following miR-124-3p agomir or FSTL1 siRNA transfection, PSCs were collected and solubilized in RIPA lysis buffer (cat. no. KGP702; Nanjing KeyGen Biotech Co., Ltd.). Following centrifugation for 10 min ( $1,600 \times g$ , 4°C), the protein concentration was measured using a BCA assay kit (Sigma-Aldrich; Merck KGaA). The samples (20  $\mu$ g/well) were separated on 10% SDS-PAGE (Nanjing Genscript Biotechnology Co., Ltd.) at 140 V for 50 min. Subsequently, the separated proteins were transferred onto PVDF membranes at 300 mA for 60 min (Nanjing Genscript Biotechnology Co., Ltd.). The membranes were subsequently blocked for 1 h at room temperature with 5% fat-free dried milk diluted in TBS-0.1% Tween-20, followed by incubation with primary antibodies against FSTL1 (1:1,000), collagen II (1:1,000), with GAPDH being used as an internal control (1:80,000), at 4°C overnight. The membranes were subsequently incubated with HRP-conjugated secondary antibodies (anti-rabbit; cat. no. ab6721; 1:10,000; Abcam) for 1 h at room temperature. The protein bands were detected using an ECL chemiluminescence kit (Nanjing KeyGen Biotech Co., Ltd., China) and semi-quantified using ImageQuant TL analysis software v8.1 (Cytiva).

**Statistical analysis.** All statistical analyses were performed using SPSS v19 software (IBM Corp.). Each experiment was repeated three times and data were presented as the mean  $\pm$  SD. Unpaired Student's t-test or one-way ANOVA followed by Bonferroni's post hoc test were applied to identify the statistical significance.  $P < 0.05$  was considered to indicate a statistically significant difference.

## Results

**miR-124-3p is associated with the differentiation of PSCs to NPLCs.** To identify miRNAs that are associated with the regulation of the differentiation of PSCs to NPLCs, miRNA microarray technology was used to detect differences in miRNA expression profiles between TGF- $\beta$ 1-treated rat primary PSCs (day 0) and differentiated NPLCs (day 15). The efficient induction of PSC differentiation by TGF- $\beta$ 1 has been previously verified in our previous study (16). The results of the present study indicated that when compared

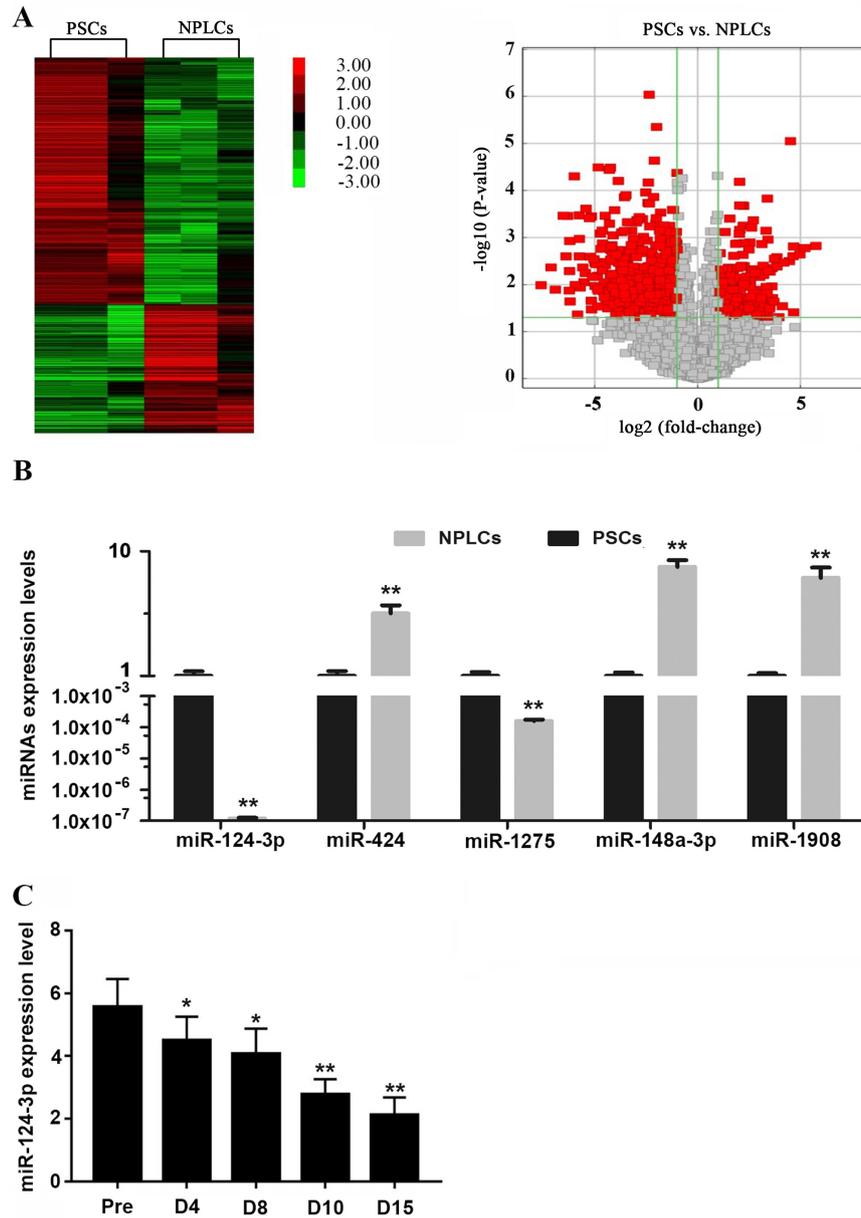


Figure 1. miR-124-3p participates in the differentiation of PSCs to NPLCs. (A) The differences in miRNA expression profiles were detected using a miRNA microarray. (B) The indicated miRNAs were validated using RT-qPCR. \*\* $P < 0.01$  vs. PSCs. (C) The expression of miR-124-3p during the differentiation of PSCs was detected using RT-qPCR. \* $P < 0.05$  and \*\* $P < 0.01$  vs. Pre (D0) group. miRNA/miR, microRNA; PSCs, precartilaginous stem cells; NPLCs, nucleus pulposus-like cells; RT-qPCR, reverse transcription-quantitative PCR; D, day.

with primary PSCs, 50 miRNAs in the differentiated NPLCs were considerably upregulated ( $>3$ -fold), while 36 miRNAs were downregulated  $<3$ -fold (Fig. 1A). A total of five miRNAs (miR-124-3p, miR-424, miR-1275, miR-148a-3p and miR-1908) with a  $>5$ -fold expression difference and a small within group variation were selected for subsequent validation using RT-qPCR. As indicated in Fig. 1B, the expression tendency of the five miRNAs, was consistent with the results of the microarray screening. Among them, the difference in the expression levels of miR-124-3p was the lowest (Fig. 1B).

The alterations in miR-124-3p expression during the differentiation of PSCs to NPLCs were subsequently analyzed (between days 0 to 15). PSCs were induced to differentiate into NPLCs via treatment with TGF- $\beta$ 1 and the expression levels of miR-124-3p at various time points of differentiation were

detected using RT-qPCR (days 0, 4, 8, 10 and 15). The results verified that the expression of miR-124-3p decreased along with the differentiation of PSCs and exhibited the lowest levels during the later stages of differentiation (Fig. 1C).

*miR-124-3p inhibits the differentiation of PSCs.* As specific markers for nucleus pulposus cells have not been identified, collagen II and aggrecan are used to detect differentiated nucleus pulposus cells (26). Therefore, the expression of collagen II and aggrecan during the differentiation of PSCs into NPLCs was detected using western blotting. It was demonstrated that collagen II and aggrecan protein were expressed at a significantly higher levels in NPLCs compared with PSCs, which suggested that PSCs were efficiently differentiated into NPLCs (Fig. 2A). Furthermore, to identify

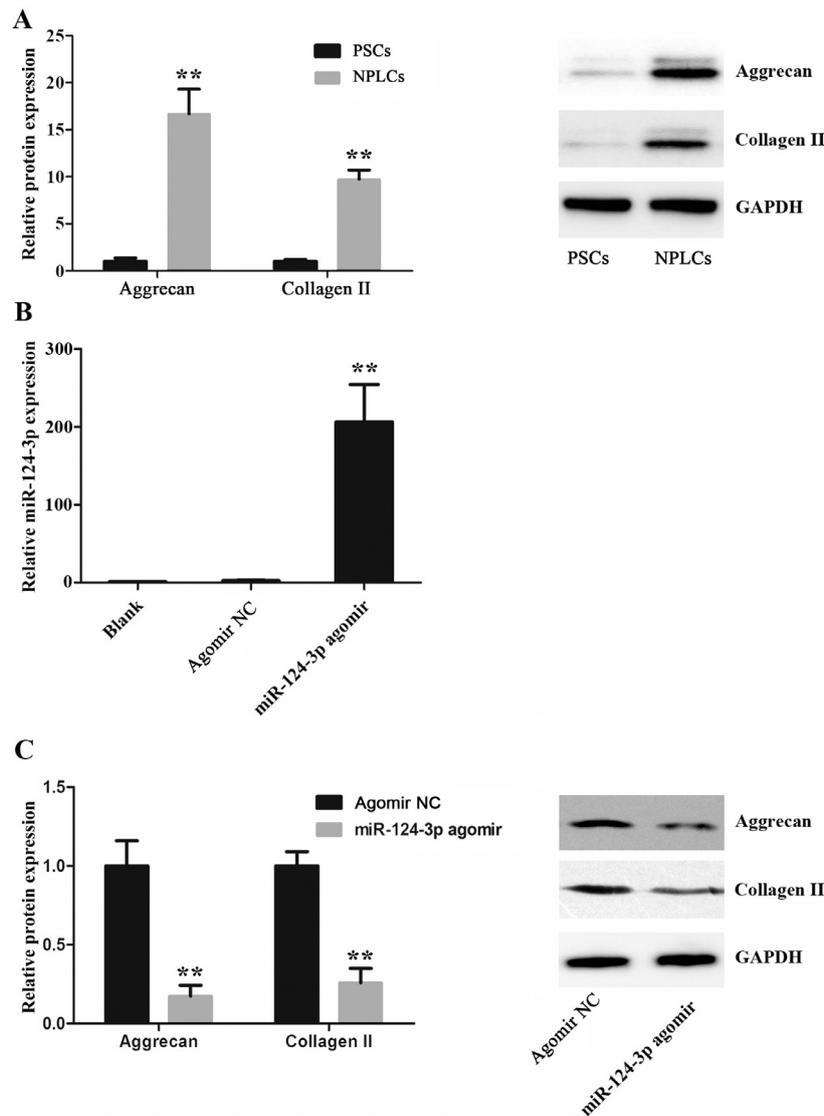


Figure 2. Effect of miR-124-3p in the differentiation of PSCs to NPLCs. (A) The expression of collagen II and aggrecan in PSCs and NPLCs was analyzed via western blotting. \*\*P<0.01 vs. PSCs. (B) The expression of miR-124-3p in PSCs transfected with miR-124-3p agomir was detected via reverse transcription-quantitative PCR. \*\*P<0.01 vs. agomir NC or Blank. (C) The expression of collagen II and aggrecan following transfection with miR-124-3p agomir was examined via western blotting. \*\*P<0.01 vs. agomir NC. miR, microRNA; PSCs, precartilaginous stem cells; NPLCs, nucleus pulposus-like cells; NC, negative control.

the role of miR-124-3p in PSCs differentiation, miR-124-3p was overexpressed in PSCs via transfection with miR-124-3p agomir (Fig. 2B). As presented in Fig. 2C, collagen II and aggrecan protein expression levels in the miR-124-3p agomir group were significantly decreased when compared with the NC group on day 18 post-transfection. These results suggested that miR-124-3p overexpression inhibited the differentiation of PSCs into NPLCs.

*miR-124-3p targets FSTL1 during PSCs differentiation.* Using TargetScan software, 7-9 binding sites were predicted in the 3'-UTR region of the FSTL1 mRNA. The results indicated that FSTL1, EZH2 and TPST2 may represent potential target genes of miR-124-3p (Fig. 3A). The alterations in the expression levels of the aforementioned three genes during the differentiation process of PSCs (0, 4, 8 and 15 days) were subsequently determined. As indicated in Fig. 3B, only FSTL1 was demonstrated to be gradually increased during the differentiation

process, in contrast to the expression levels of miR-124-3p, which were indicated to decrease during PSCs differentiation, as aforementioned. However, the expression of EZH2 and TPST2 showed neither gradual increase nor decrease. Notably, the majority of miRNAs have been indicated not to affect the mRNA expression of their target gene. However, certain miRNAs have been reported to result in mRNA degradation of their target genes (27). In the present study, it was demonstrated that miR-124-3p overexpression resulted in decreased FSTL1 mRNA levels in PSCs (Fig. 3C).

Moreover, in order to verify whether miR-124-3p can directly bind to and regulate the mRNA expression levels of FSTL1, a dual-luciferase reporter assay was used in the present study. Luciferase reporter plasmids containing the predicted GUGCCUU binding site of the 3'-UTR of FSTL1 and a mutant site (UGCUUGC) were constructed and co-transfected with miR-124-3p into COS7 and 293 cells to detect the alterations in luciferase activity. As indicated in Fig. 4A, the results



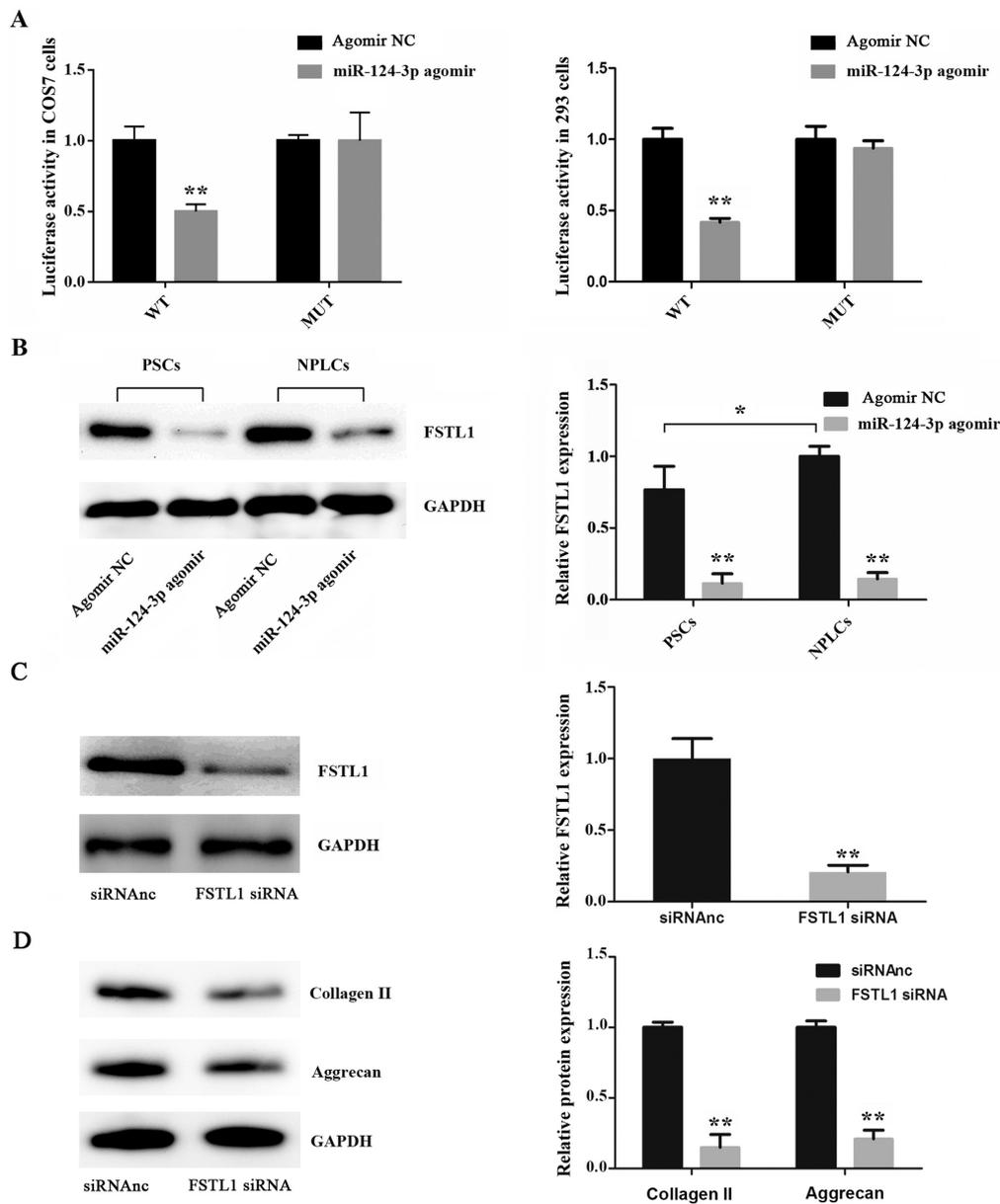


Figure 4. miR-124-3p directly regulates FSTL1 function. (A) Luciferase reporter plasmids containing the predicted GUGCCUU binding site of the 3'-untranslated region of FSTL1 and its mutant homolog (UGCUUGC) were constructed and co-transfected with miR-124-3p into COS7 and 293 cells to detect the alterations in luciferase activity. \*\*P<0.01 vs. agomir NC. (B) The effects of miR-124-3p overexpression on FSTL1 expression during the differentiation of PSCs to NPLCs was analyzed via western blotting. \*P<0.05 vs. PSCs; \*\*P<0.01 vs. agomir NC. (C) The efficiency of FSTL1 siRNA was verified via western blotting. \*\*P<0.01 vs. siRNanc. (D) The expression of collagen II and aggrecan following FSTL1 knockdown was detected via western blotting. \*\*P<0.01 vs. siRNanc. miR, microRNA; MUT, mutant; PSCs, precartilaginous stem cells; NPLCs, nucleus pulposus-like cells; NC/nc, negative control; FSTL1, follistatin-related protein 1; si, small interfering; WT, wild-type.

only FSTL1 levels increased gradually during the differentiation of PSCs to NPLCs. Considering that miRNAs recognize their target genes via complementary base pairing and subsequently guide the silencing complex to degrade or repress the translation of the target mRNA according to the degree of complementarity (27), it was speculated that FSTL1 may be a potential target gene of miR-124-3p. FSTL1 was firstly identified in mouse osteoblasts and as TGF- $\beta$ 1 has been indicated to induce its upregulation, FSTL1 is also known as TGF- $\beta$ -stimulated clone-36 (36). FSTL1 is an extracellular matrix protein, which is widely expressed in all eukaryotic cells and has been associated with cell differentiation, metabolism, cell proliferation and the immune response (37-39). A previous

study indicated that FSTL1 also participates in the regulation of Lumbar disc herniation (40). It was demonstrated that FSTL1 expression is increased during the progression of intervertebral disc disease and promotes inflammatory reactions in the nucleus pulposus via the MAPK and NF- $\kappa$ B signaling pathways (40). The present study revealed that FSTL1 was also associated with the differentiation of PSCs to NPLCs, a process which was indicated to be regulated by miR-124-3p. Overexpression of miR-124-3p resulted in decreased expression levels of FSTL1 during the differentiation of PSCs to NPLCs and dual-luciferase reporter assays identified FSTL1 as a direct target of miR-124-3p. A previous study reported that PSCs can be induced to differentiate to

NPLCs to repair degenerative intervertebral discs, which can provide unique therapeutic advantages (24). The present study elucidated the function and molecular mechanism underlying the role of miR-124-3p in the differentiation of PSCs to NPLCs. Therefore, it was hypothesized that miR-124-3p may be a target for the induction of PSC differentiation and the subsequent repair of degenerated intervertebral discs.

In conclusion, the present study demonstrated a novel role for miR-124-3p in the differentiation of PSCs to NPLCs, and provided evidence that miR-124-3p negatively regulated its target gene, FSTL1, during PSC differentiation. However, the limitations of the present study lie in the lack of effective *in vivo* experiments to demonstrate the regulatory effect of miR-124-3p on PSCs. Moreover, whether miR-124-3p can alleviate DDD remains unclear. Therefore, further investigations into the function of miR-124-3p *in vivo* should be an aim of future studies.

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### Availability of data and materials

All data generated and/or analyzed during the present study are included in the published article.

### Authors' contributions

QW designed the current study and acquired funding. JW, XG, DF, DL and TJ performed the experiments, collected the data and wrote the manuscript. QW and JW confirmed the authenticity of all the raw data. All authors have read and approved the final manuscript.

### Ethics approval and consent to participate

The present study was approved by Institutional Animal Care and Use Committee of Wuxi People's Hospital (approval no. WXP20190103c0600105).

### Patient consent for publication

Not applicable.

### Competing interests

The authors declare that they have no competing interests.

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