

# microRNA-25 promotes osteosarcoma cell proliferation by targeting the cell-cycle inhibitor p27

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Abstract. An increasing body of evidence indicates that microRNAs (miRNAs), a class of small non-coding RNAs, are often aberrantly expressed in human osteosarcoma. This study aimed to investigate the effects of miR-25 and to identify its potential target genes in osteosarcoma (OS) cells. First, the expression of miR-25 was detected by reverse transcription-quantitative polymerase chain reaction (RT-qPCR), which revealed a significant upregulation of miR-25 in osteosarcoma tissues compared to the adjacent healthy tissues. To investigate the role of miR-25 in osteosarcoma cell proliferation, the miR-25 precursor was next transfected into Saos-2 and U2OS cells. Overexpression of miR-25 promoted cell proliferation in vitro and tumor growth in a xenograft mouse model. In addition, our results revealed that the protein expression of p27, a cell-cycle inhibitor, is negatively regulated by miR-25. Restoring the p27 level in miR-25-overexpressing cells reversed the enhancing effect of miR-25 on cell proliferation. Therefore, miR-25 may act as an onco-miRNA in osteosarcoma, which provides new perspectives in cancer treatment strategies based on molecular targeting.

# Introduction

microRNAs (miRNAs), a class of short non-coding RNA molecules, function as transcriptional or post-transcriptional regulators of gene expression (1,2). Deregulation of miRNAs is tightly associated with human disorders, including obesity, cardiovascular diseases and tumorigenesis (3,4). Recent studies have also shown that a number of miRNAs are upregulated or downregulated and play critical roles in osteosarcoma development (5,6). For instance, miR-376c was shown to inhibit cell proliferation and invasion in osteosarcoma by targeting

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transforming growth factor- $\alpha$  (7), whereas miR-221 induces osteosarcoma cell survival through enhancing the AKT signaling pathway (8).

Previous studies have shown that miR-25 expression and function is deregulated in several types of tumor (9-12). It was shown that miR-25 is consistently highly expressed in the serum of patients with breast cancer or hepatocellular carcinoma, indicating that this miR may be used as a biomarker in diagnosis and treatment (9,10). In addition, the miR-25 level was significantly increased in human gastric cancer (GC) tissues and cell lines (11). Overexpression of miR-25 markedly enhanced cell proliferation, migration and invasion in GC cells, whereas inhibition of miR-25 caused a significant reduction in proliferation rates and a significant increase in apoptosis (11). However, miR-25 was found to be downregulated in human colon cancer tissues when compared to matched, non-neoplastic mucosa tissues (12). Functional studies revealed that restoration of the miR-25 expression inhibits cell proliferation and migration (12). By contrast, miR-25 inhibition promoted cell proliferation and migration (12). Therefore, miR-25 appears to act as either an onco-miRNA or a tumor suppressor in different types of cancer, and plays a crucial role in cancer biology. However, its biological functions in osteosarcoma remain unexplored to date.

## Materials and methods

Cell culture and tissue samples. Osteosarcoma cell lines (Saos-2 and U2OS) were obtained from the American Type Culture Collection (Rockville, MD, USA). Cells were cultured in Gibco<sup>®</sup> RPMI-1640 (Thermo Fisher scientific, Inc., Beijing, China) supplemented with 10% Gibco® fetal bovine serum (Thermo Fisher Scientific, Inc.). Tumor tissues and adjacent noncancerous normal tissues were collected from routine therapeutic surgery at the Department of Orthopaedics, Pudong New Area Zhoupu Hospital. A total of 25 samples (male; median age, 57 years; range, 48-65 years) were obtained with informed consent from patients with osteosarcoma, and the procedures were approved by the Institutional Review Board of the hospital. Subjects were excluded if they had other diseases, including biliary obstructive diseases, and acute or chronic virus hepatitis. The individuals with an alcohol consumption of  $\geq 120$  g/week for men at the time of the study or in the prior 6 months were also excluded from the study.

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Analysis of miRNA expression. miRNA from tissue samples and cell lines was harvested using the Ambion® mirVana miRNA Isolation kit (Thermo Fisher Scientific, Inc., Grand Island, NY, USA), following the manufacturer's instructions. All RNA samples were examined as to their concentration and purity. RNA purity was measured using the NanoDrop ND-1000 spectrophotometer (Thermo Fisher Scientific, Waltham, MA, USA). Based on the absorbance ratio at 260/280 nm (mean $\pm$ standard deviation = 1.86 $\pm$ 0.03), all RNA samples were pure and protein free. Expression of mature miRNAs was assessed with the Applied Biosystems® TaqMan® microRNA assay (Thermo Fisher Scientific, Inc.) with probes specific to the human hsa-miR25 (5'-aggcggagacuugggcaauug-3') and the small nuclear U6 snRNA, used as an internal control (5'-cauugaccauggacauacgacug-3'). The quantitative (q)PCR reaction was performed using a TaqMan Universal PCR Master mix on an Applied Biosystems® 7900HT Real-Time PCR system (all from Thermo Fisher Scientific, Inc.). Briefly, PCR conditions included an initial holding period at 95°C for 5 sec and 60°C for 30 sec for 45 cycles. Relative quantitation analysis of the gene expression data was conducted according to the  $2^{-\Delta\Delta Ct}$  method.

*Plasmid construction and transfection.* To construct the miR-25 expression plasmid, the precursor sequence of the human miR-25 was cloned into Ambion<sup>®</sup> pSilencer<sup>™</sup>, while the negative control (NC) plasmid contained a scrambled sequence (both from Thermo Fisher Scientific, Inc.). For transfection, Lipofectamine<sup>®</sup> 2000 (Thermo Fisher Scientific, Inc.) was employed following the manufacturer's instructions.

Bromodeoxyuridine (BrdU) assay. To assess cell proliferation, an enzyme-linked immunosorbent assay (ELISA) was used, based on the incorporation of BrdU during DNA synthesis, (BrdU kit; Beyotime Institute of Biotechnology, Shanghai, China), following the manufacturer's protocols. All experiments were performed in triplicate. The absorbance of the samples at 450 nm ( $A_{450}$ ) was measured on a SpectraMax 190 ELISA reader (Molecular Devices, Sunnyvale, CA, USA).

Western blot analysis. Cells were harvested and lysed with ice-cold lysis buffer containing 50 mM Tris-HCl, pH 6.8, 100 mM 2-Mercaptoethanol, 2% w/v sodium dodecyl sulfate (SDS) and 10% glycerol. Following centrifugation at 4°C, proteins in the supernatants were quantified by a BCA quantification kit (Beyotime Institute of Biotechnology), separated by 10% SDS-polyacrylamide gel electrophoresis (PAGE), and transferred onto nitrocellulose membranes (Amersham Biosciences, Buckinghamshire, UK). Anti-p27 antibody (Cell Signaling Technology, Inc., Danvers, MA, USA) was used at 1:2,000 overnight at 4°C, and anti-glyceraldehyde 3-phosphate dehydrogenase (GAPDH) antibody (Cell Signaling Technology, Inc.) was used at 1:5,000 overnight at 4°C. The membranes were then incubated with the HRP-linked secondary antibodies (Cell Signaling Technology, Inc.). The signals were detected by SuperSignal West Pico Chemiluminescent Substrate kit (Pierce Biotechnology, Inc., Rockford, IL, USA) according to manufacturer's instructions. The images were visualized by a LAS-4000 Luminescent Image analyzer (Fujifilm, Tokyo, Japan). The p27 protein level was quantified using Quantity One software (Bio-Rad, Hercules, CA, USA) and was normalized to that of GAPDH.



Figure 1. Expression level of miR-25 in osteosarcoma tissues. miR-25 expression was determined by reverse transcription-quantitative polymerase chain reaction assays in human osteosarcoma tissues and adjacent noncancerous tissues (normal). \*\*\*P<0.001.

Luciferase reporter assay. Total cDNA from Saos-2 cells was synthesized from total RNA using random hexamers with the Superscript III Reverse Transcriptase kit (Invitrogen, Carlsbad, CA, USA), according to the manufacturer's instructions. The reactions were incubated in a thermal cycler for 30 min at 16°C, 30 min at 42°C, 5 min at 85°C and then held at 4°C. The cDNA was used to amplify the 3' untranslated region (3' UTR) of p27 by PCR. Mutations were introduced in potential miR-25-binding sites (5'-GCUAUUACGAAUACAUCCGUUAAC-3' was changed into 5'-GCUAUUACGAAUACAUCCGAAUUC-3') using the QuickChange® Site-Directed Mutagenesis kit (Stratagene, La Jolla, CA, USA). The pRL-SV40 vector (Promega Corp., Madison, WI, USA) carrying the Renilla luciferase gene was used as an internal control of transfection efficiency. Luciferase values were determined using the Dual-Luciferase® Reporter Assay system (Promega Corp.).

*Tumor growth assay.* Male BALB/c nude mice, 4 weeks-old, were purchased from the Shanghai Laboratory Animal Center (Shanghai, China). Saos-2 cells  $(2x10^5)$  were subcutaneously injected into the skin under the legs of the mice. The mice were observed for 5 weeks for tumor formation. Following sacrifice, the tumors were recovered and the wet weights of each tumor were measured.

Statistical analysis. Data were expressed as the mean  $\pm$  SEM from at least three independent repetitions of the experiment. Differences between groups were analyzed using Student's t-tests or one way analysis of variance (ANOVA). P<0.05 was considered to indicate a statistically significant difference.

## Results

The miR-25 expression level is increased in osteosarcoma tissues. First, in order to examine whether the miR-25 is differentially expressed in human osteosarcoma compared to noncancerous tissues, its expression level was determined using reverse transcription (RT)-qPCR in 25 pairs of human osteosarcoma tissues and pair-matched adjacent noncancerous tissues. The results demonstrated that the expression level of miR-25 was significantly increased in osteosarcoma tissues compared to the adjacent noncancerous tissues (Fig. 1).





Figure 2. Overexpression of miR-25 promotes osteosarcoma cell proliferation. Expression of miR-25 was determined in (A) Saos-2 and (B) U2OS cells following transfection with the plasmid bearing the miR-25 precursor or the negative control (NC) plasmid. The growth curve of (C) Saos-2 and (D) U2OS cells following miR-25 precursor transfection as compared to that a transfection with NC. Cell proliferation was determined by the bromodeoxyuridine (BrdU) assay in (E) Saos-2 and (F) U2OS cells transfected with the miR-25 precursor-bearing plasmid or the NC. Absorption at 450 nm ( $A_{450}$ ) was measured 24 h after the transfection. \*\*P<0.05; \*\*\*P<0.01.



Figure 3. miR-25 promotes osteosarcoma cell proliferation *in vivo*. Saos-2 cells mock-transfected (NC) or stably transfected with miR-25 were injected into nude mice (n=6 for each group), which were monitored for tumorigenesis. (A) Tumor weights and (B) representative pictures of tumors 4 weeks following the injection.

*miR-25 overexpression promotes cell proliferation in vitro*. In order to assess the effects of miR-25 on osteosarcoma cell growth, Saos-2 and U2OS cells were transfected with a plasmid bearing the miR-25 precursor or the NC plasmid, and cell growth was

subsequently examined. Transfection with the miR-25 precursorbearing plasmid increased miR-25 expression compared to the NC transfection (Fig. 2A and B), and significantly increased the cell number and proliferation in both cell lines (Fig. 2C-F).



Figure 4. miR-25 negatively regulates p27 expression in osteosarcoma cells. (A) Predicted miR-25-binding sites (highlighted in bold) in the 3' untranslated region (3' UTR) of the human gene *p27*. (B) Luciferase reporter assays in Saos-2 cells. Cells were transfected with 200 ng of the wild-type 3' UTR reporter or mutant constructs together with 20 nM of the miR-25 precursor or a scrambled control. Representative western blots showing the relative expression of the protein p27 in (C) Saos-2 and (D) U2OS cells transfected with the miR-25 precursor or negative control (NC), (E) tumors from nude mice and (F) two patients with osteosarcoma. The glyceraldehyde 3-phosphate dehydrogenase (GAPDH) level was used as the loading control.



Figure 5. Re-introduction of p27 reverses the promoting effect of miR-25 on cell proliferation. (A) p27 protein expression was determined by western blot analysis in Saos-2 cells. Cells were pre-transfected with the miR-25 precursor or the negative control (NC) for 24 h and then transfected with expression plasmids for p27 or the empty vector (EV) for another 24 h. (B) The growth curve and (C) cell proliferation rate was determined in Saos-2 cells. Absorption at 450 nm ( $A_{450}$ ) was measured 30 h. after the transfection with p27 or EV.

*miR-25 overexpression promotes cell proliferation in vivo*. To study the role of miR-25 overexpression in tumorigenesis *in vivo*, we generated Saos-2 cells stably overexpressing miR-25 and injected the cells into a xenograft mouse model. miR-25 overexpression markedly promoted tumorigenesis in comparison with the NC, as evidenced by tumor weights and sizes (Fig. 3A and B).

miR-25 targets the p27 in osteosarcoma cells. Since miR-25 overexpression promoted cell proliferation and tumor growth, we examined the underlying mechanisms. Through a stringent bioinformatics approach (miRWalk software; http://www.umm. uni-heidelberg.de/apps/zmf/mirwalk/), we identified the gene p27, encoding a cell-cycle inhibitor, as a candidate target of miR-25, since p27 bears a potential miR-25-binding site (Fig. 4A). The 3' UTR of p27 was cloned into a reporter luciferase system. When the reporter construct contained the p37 3' UTR, overexpression of miR-25 led to a reduction in luciferase activity (Fig. 4B). By contrast, mutation of the conserved miR-25-binding motif abrogated the reduced luciferase expression (Fig. 4B). In addition, overexpression of miR-25 in osteosarcoma cells led to a reduction in the protein level of p27 (Fig. 4C and D). In addition, the p27 protein level was reduced in miR-25-overexpressing tumors and human osteosarcoma tissues (Fig. 4E and F), further supporting that p27 may be a target of miR-25 in osteosarcoma cells.

p27 restoration attenuates the promoting effect of miR-25 overexpression on cell proliferation. In order to confirm the functional connection between miR-25 and p27, Saos-2 cells were transfected with p27 expression plasmids following transfection with miR-25 (Fig. 5A). As shown in Fig. 5B and C, the re-introduction of p27 reversed miR-25-induced cell proliferation, indicating that the interaction between miR-25 and p27 is involved in this process. Taken together, our results suggest that the gene p27 is an important target of miR-25 in osteosarcoma cells.

## Discussion

In this study, we demonstrate that miR-25 expression is increased in osteosarcoma tissues. Overexpression of miR-25 by means of cell transfection promoted cell proliferation in Saos-2 and U2OS cells. Therefore, our study provided evidence, for the first time to the best of our knowledge, that miR-25 may act as an onco-miRNA, promoting the progression of osteosarcoma. It is notable that miR-25 was shown to enhance cell proliferation, migration and invasion in GC cells, but inhibit cell proliferation and migration in colon cancer cells (11,12). Although the reasons for this inconsistence remain unexplored, we hypothesize that the biological functions of miR-25 in tumorigenesis may be cell- or tissue-specific. Therefore, the efficiency and safety associated with the use of miR-25 in gene therapy should be seriously considered in future studies.

Our study explored the mechanisms underlying the miR-25 effects, and revealed that p27 maybe a target of miR-25 in osteosarcoma cells. p27 encodes an enzyme inhibitor that belongs to the Cip/Kip family of cyclin-dependent kinase (CDK) inhibitor proteins (13). Through binding to the CDK2 and CDK4 complexes, it prevents their activation, and controls the cell-cycle progression at the G1 phase (14,15). Transcriptional or post-transcriptional downregulation of the gene has been observed in a number of human malignancies,

including osteosarcoma (16,17). For instance, p27 was shown to be negatively regulated by miR-24, miR-200 and miR-222 in human cancer (18-20). Therefore, our results provide a novel mechanism for the downregulation of p27 in osteosarcoma.

Taken together, the key finding of the present study is that miR-25 can promote osteosarcoma cell proliferation *in vitro* and *in vivo* by targeting p27, suggesting that miR-25 may be used as a molecular target for the treatment of osteosarcoma. However, the roles of this miRNA in osteosarcoma need to be further investigated.

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