

MicroRNA-107 Promotes Proliferation, Migration, and Invasion of Osteosarcoma Cells by Targeting Tropomyosin 1

Rui Jiang,* Chao Zhang,† Guangyao Liu,* Rui Gu,* and Han Wu*

*Department of Orthopedics, China–Japan Union Hospital of Jilin University, Changchun, Jilin, P.R. China

†Department of Ophthalmology, The Second Hospital of Jilin University, Changchun, Jilin, P.R. China

Osteosarcoma is the most common primary bone malignancy manifested predominantly in children and young adults. Studies indicate that miR-107 is involved in the pathogenesis of osteosarcoma and that tropomyosin 1 (TPM1) acts as a tumor suppressor in many types of cancer. In this study, we analyzed the effect of miR-107 on human osteosarcoma cells and investigated the mechanism in which TPM1 is involved. miR-107 expression in human osteosarcoma tissues and cells was analyzed in quantitative real-time PCR (qRT-PCR). Human osteosarcoma (U2OS) cells were transfected with miR-107 mimic, inhibitor, or scramble controls to evaluate the effect of miR-107 on cellular migration and invasion, cell viability, and apoptosis. Cells were cotransfected with the miR-107 mimic and TPM1 3'-UTR wild-type (wt) recombinant vector or mutant type (mt) as a negative control. The binding effect of miR-107 on TPM1 3'-UTR was determined by dual-luciferase reporter assay. The expression of TPM1, apoptosis-related proteins, and signaling molecules was determined by qRT-PCR and Western blotting. The results showed that miR-107 expression was upregulated in osteosarcoma tissues and cell lines. miR-107 overexpression promoted U2OS cell viability, migration, and invasion whereas it inhibited apoptosis. miR-107 inhibitor transfection ameliorated or abolished these effects after miR-107 binding to TPM1 3'-UTR-wt regulated TPM1 expression. miR-107 in U2OS cells activated MEK/ERK and NF- κ B signaling pathways via TPM1. In conclusion, miR-107 overexpression promoted U2OS cell viability, migration, and invasion via down-regulation of TPM1 and might be through activating the MEK/ERK and NF- κ B signaling pathways.

Key words: Osteosarcoma; miR-107; Tropomyosin 1 (TPM1); MEK/ERK; NF- κ B

INTRODUCTION

Osteosarcoma is the most common primary bone malignancy, representing about 20% of all bone tumors. Osteosarcoma manifests predominantly in children and young adults and accounts for 5% of all pediatric tumors^{1–3}. Furthermore, this complex and heterogeneous disease, characterized by the direct formation of immature bone or osteoid tissue by tumor cells, is associated with high rates of recurrence and early metastasis. Despite improvements in surgical techniques and chemotherapeutic approaches, the treatment of osteosarcoma remains a clinical challenge^{4,5}. Thus, a comprehensive understanding of the pathogenesis of osteosarcoma is critical for developing novel and improved therapeutic strategies.

MicroRNAs (miRNAs) represent a class of highly conserved, small, noncoding RNA molecules that are widely expressed in eukaryotic cells. These molecules regulate protein expression by binding to specific sequences in mRNA molecules, most commonly in the 3'-UTR^{6,7}. Studies have confirmed the existence of differences in

miRNA expression profiles between cancerous and matched noncancerous tissues⁸, and reports have indicated that miRNAs play an important role in the growth and metastasis of human cancers, including osteosarcoma^{9,10}.

miR-107 plays a variety of roles in different tissues and cells, including cell division, metabolism, stress responses, and tumor angiogenesis. Furthermore, miR-107 has been reported to be involved in the pathogenesis of multiple tumors¹¹. For example, miR-107 inhibits glioma cell proliferation by targeting SALL4¹², and miR-107 overexpression inhibits colon cancer cell proliferation by targeting HIF-1 β ¹³. However, many studies have demonstrated that miR-107 overexpression promotes tumor cell proliferation in cancers such as hepatocellular carcinoma¹⁴ as well as the EMT process and tumor cell metastasis in gastric cancer cells¹⁵. These reports demonstrated the highly complex nature of miR-107 functions.

Tropomyosin 1 (TPM1) has been reported to function as a tumor suppressor gene that is downregulated in several human cancers, including breast cancer, colon cancer,

Address correspondence to Professor Han Wu, Department of Orthopedics, China–Japan Union Hospital of Jilin University, No. 126, Xiantai Street, Changchun, Jilin Province 130033, P.R. China. E-mail: wuhan12017@126.com

and urinary bladder carcinoma^{16–18}. However, the role of TPM1 in the pathogenesis of osteosarcoma remains to be established.

In the present study, we analyzed the effects of miR-107 on human osteosarcoma cells and investigated the mechanism in which TPM1 might be involved.

MATERIALS AND METHODS

Clinical Specimens

Clinical human osteosarcoma tissues and the corresponding normal osteoblasts ($n=25$) were obtained from the China–Japan Union Hospital of Jilin University (Changchun, Jinlin, P.R. China). None of the patients received any therapy before surgery. Informed consent was obtained from all patients, and the present study was approved by the medical ethics committee of the China–Japan Union Hospital of Jilin University.

Cell Culture

The human osteosarcoma cell lines MG63, U2OS, and OS732, and the human osteoblast cell line hFOB1.19 were obtained from Shanghai Institutes for Biological Sciences Cell Resource Center and were cultured in high-glucose Dulbecco's modified Eagle's medium (DMEM) supplemented with 10% fetal bovine serum (Gibco, Carlsbad, CA, USA). All cells were incubated at 37°C in a humidified incubator under 5% CO₂.

miRNA Transfection

U2OS cells were seeded into six-well plates (5×10^4 cells/well) and cultured overnight to approximately 70% confluence in a humidified atmosphere containing 5% CO₂ at 37°C. The cells were then transfected with 100 pmol of miR-107 mimic, miR-107 inhibitor, scramble controls, si-TPM1, or siRNA-negative controls, which were synthesized by GenePharma Co. (Shanghai, P.R. China). Cell transfections were conducted using Lipofectamine 3000 reagent (Invitrogen, Carlsbad, CA, USA) according to the manufacturer's instructions. Forty-eight hours after transfection, the cells were collected for the next analysis.

RNA Extraction and Quantitative Real-Time PCR (qRT-PCR) Analysis

Total RNA was extracted from cells and tissues using TRIzol reagent (Life Technologies, Carlsbad, CA, USA) according to the manufacturer's instructions. The One Step SYBR[®] PrimeScript[®] PLUS RT-RNA PCR Kit (TaKaRa Biotechnology, Dalian, P.R. China) was used for the RT-PCR analysis of TPM1 expression levels. The TaqMan MicroRNA Reverse Transcription Kit and TaqMan Universal Master Mix II were used with the TaqMan MicroRNA Assay (Applied Biosystems, Foster City, CA, USA) for analysis of miR-107 expression levels in cells and tissues.

Cell Viability Assay

Cell viability was assessed using the cell counting kit-8 (CCK-8; Dojindo Molecular Technologies, Gaithersburg, MD, USA). Briefly, after stimulation, cells were seeded into 96-well plates (5×10^3 cells/well), the CCK-8 solution was added to the culture medium, and the cells were incubated for 1 h at 37°C in a humidified incubator under 5% CO₂. Cell viability was determined by measuring the absorbance of the culture supernatants at 450 nm using a microplate reader (Bio-Rad, Hercules, CA, USA).

Cell Migration and Invasion Assays

Cell migration was determined using a modified two-chamber system separated with 8-mm pore size PET membrane of (Millipore, Bedford, MA, USA). Briefly, after treatment as indicated, cells (5×10^4) were suspended in 200 μ l of serum-free medium and seeded into the upper compartment of a 24-well Transwell culture chamber (Corning, Elmira, NY, USA), and 600 μ l of complete medium was added to the lower compartment. After incubation for 24 h at 37°C under 5% CO₂, the cells were fixed with methanol. Nontraversed cells were carefully removed from the upper surface of the filter with a cotton swab. Traversed cells on the lower side of the filter were stained with 0.1% crystal violet and counted using a light microscope (Olympus, Tokyo, Japan).

The invasion behavior of cells was determined using BD BioCoat[™] Matrigel[™] Invasion Chambers with Matrigel (8- μ m pore size polycarbonate filters; BD Biosciences, San Jose, CA, USA). Briefly, after treatment as indicated, 5×10^4 cells resuspended in 200 μ l of serum-free DMEM were plated onto the upper compartment, while 600 μ l of complete medium containing 10% FBS was added to the lower chamber. After incubation for 48 h at 37°C under 5% CO₂, the noninvading cells were removed with a cotton swab. The invading cells were fixed with 100% methanol and stained with 0.1% crystal violet solution, after which they were counted by a light microscope (Olympus). The relative number of cells attached on the bottom surface from five randomly chosen fields was calculated.

Apoptosis Assay

Cell apoptosis analysis was performed using propidium iodide (PI) and fluorescein isothiocyanate (FITC)-conjugated annexin V staining. Briefly, after cells were treated, they were washed in phosphate-buffered saline (PBS) and fixed in 70% ethanol. Fixed cells were then washed twice in PBS and stained using PI/FITC-annexin V with the presence of 50 μ g/ml RNase A (Sigma-Aldrich, St. Louis, MO, USA) and then incubated for 1 h at room temperature in the dark. Flow cytometric analysis was performed using a FACScan (Beckman Coulter, Fullerton, CA, USA), and the data were analyzed using FlowJo software.

Dual-Luciferase Activity Assay

The wild-type TPM1 3'-UTR (TPM1-wt) sequence carrying the putative miR-107 binding site was generated as shown by TargetScan. To mutate the TPM1 3'-UTR as a negative control, the QuikChange II kit (Stratagene, San Diego, CA, USA) was used according to the manufacturer's instructions. The sequences were cloned into the pGL promoter vector to generate luciferase reporter constructs. Cells were cotransfected with the reporter construct or negative control and miR-107 mimic or scramble using Lipofectamine 3000 (Life Technologies) according to the manufacturer's instructions. After 24 h, luciferase activity was analyzed using the Dual-Luciferase Reporter System (Promega, Madison, WI, USA) following the manufacturer's protocol. Firefly luciferase activity was normalized to that of *Renilla* luciferase. Each transfection was assayed at least three times.

Western Blot Analysis

Total cellular proteins were extracted using RIPA lysis buffer (Beyotime Biotechnology, Shanghai, P.R. China) supplemented with protease inhibitors (Roche, Guangzhou, P.R. China). The proteins were quantified using the BCA™ Protein Assay Kit (Pierce, Appleton, WI, USA). Whole-protein samples were separated using a Bio-Rad Bis-Tris Gel system (Bio-Rad) according to the manufacturer's instructions and transferred to a polyvinylidene fluoride (PVDF) membrane. The membrane was blocked in 5% nonfat milk for 2 h at room temperature and then incubated overnight at 4°C with the following primary antibodies diluted 1:1,000 in blocking buffer: Bcl-2 (ab32124; Abcam, Cambridge, MA, USA), Bax (ab32503; Abcam), caspase 3 (3CSP03; Santa Cruz Biotechnology),

TPM1 (ab55915; Abcam), and anti-GAPDH antibody (ab8245; 1:100 dilution; Abcam). After washing, the membranes were incubated for 1 h at room temperature with a horseradish peroxidase (HRP)-conjugated secondary antibody (1:1,000 dilution; ERP3312, ab197034; Sigma-Aldrich). After washing, the membranes were transferred into the Bio-Rad ChemiDoc™ XRS system, and 200 µl of Immobilon Western Chemiluminescent HRP Substrate (Millipore, Boston, MA, USA) was added to cover the surface. The signals were captured, and the intensity of the bands was quantified using Image Lab™ Software (Bio-Rad, Shanghai, P.R. China).

Statistical Analysis

All statistical analyses were performed using SPSS 19.0 statistical software. Data are presented as the mean ± standard deviation (SD) of three independent experiments analyzed using a one-way analysis of variance (ANOVA). A value $p < 0.05$ was considered to indicate a statistical significance.

RESULTS

miR-107 Was Upregulated in Osteosarcoma Tissues and Cell Lines

miR-107 expression was evaluated in clinical human osteosarcoma tissues and corresponding nontumor osteoblast tissues ($n=25$) by qRT-PCR. miR-107 expression was significantly upregulated in tumor tissues compared to nontumor tissues ($p < 0.01$) (Fig. 1A). Expression of miR-107 levels was also significantly upregulated in the human osteosarcoma cell lines MG63, U2OS, and OS732 compared with that of the normal human osteoblast cell line hFOB1.19 ($p < 0.01$) (Fig. 1B). The highest

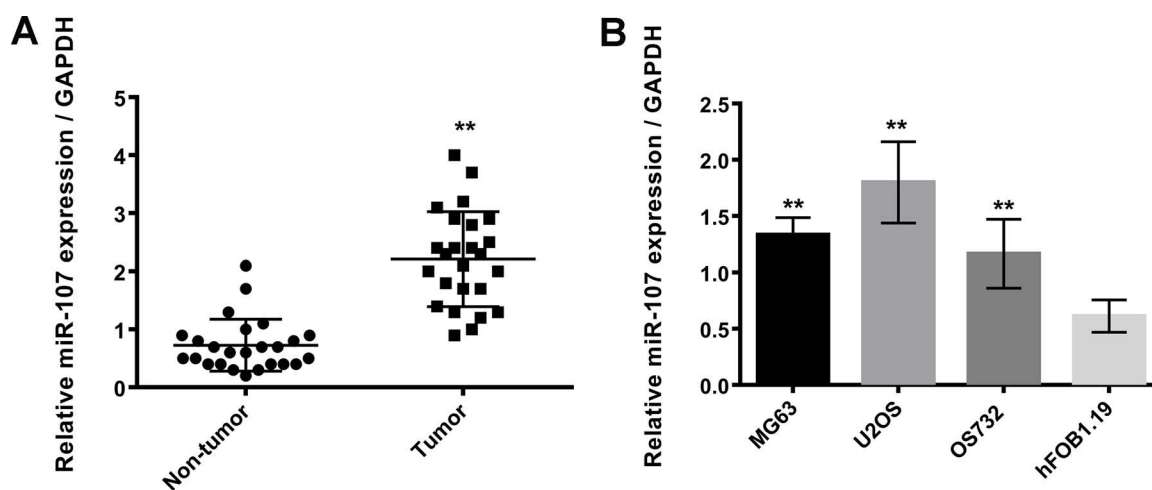


Figure 1. miR-107 was upregulated in osteosarcoma tissues and cell lines. miR-107 expression was evaluated by quantitative real-time (qRT)-PCR in (A) clinical human osteosarcoma tissues and adjacent nontumor osteoblasts ($n=25$) and (B) human osteosarcoma cell lines MG63, U2OS, and OS732 and the normal human osteoblast cell line hFOB1.19. Data represent the mean ± SD of three independent experiments. ** $p < 0.01$.

expression levels of miR-107 were detected in the U2OS cell line, which was selected for further experiments.

miR-107 Influenced U2OS Cell Viability and Apoptosis

The effect of the aberrant expression of miR-107 was investigated by transfection of U2OS cells with the miR-107 mimic to induce significant overexpression of miR-107, whereas miR-107 suppression was achieved by miR-107 inhibitor-mediated silencing compared with scramble controls, respectively ($p < 0.005$ or $p < 0.01$) (Fig. 2A). Overexpression of miR-107 resulted in a significant increase in cell viability, which was significantly decreased by the miR-107 inhibitor ($p < 0.05$ or $p < 0.05$) (Fig. 2B). Furthermore, miR-107 inhibitor transfection resulted in a significant increase in apoptosis ($p < 0.001$ vs. inhibitor control), whereas transfection with the miR-107 mimic or the scramble control had

no effect on the rate of U2OS cell apoptosis compared with that of untransfected cells (Fig. 2C). Western blot analysis of apoptosis-related protein expressions showed that the miR-107 inhibitor transfected in U2OS cells resulted in decreased expression of the antiapoptotic protein Bcl-2, and that expression of the proapoptotic protein Bax was markedly upregulated (Fig. 2D). In contrast, miR-107 overexpression in U2OS cells resulted in the upregulation of Bcl-2 and the downregulation of Bax. Transfection with scramble controls had no significant effect on the expression of these apoptosis-related proteins. These observations suggested that miR-107 inhibited cell apoptosis of U2OS cells.

Caspase 3 is activated by proteolytic cleavage to generate the active form of the enzyme, which plays a central role in apoptosis. Transfection of the miR-107 inhibitor

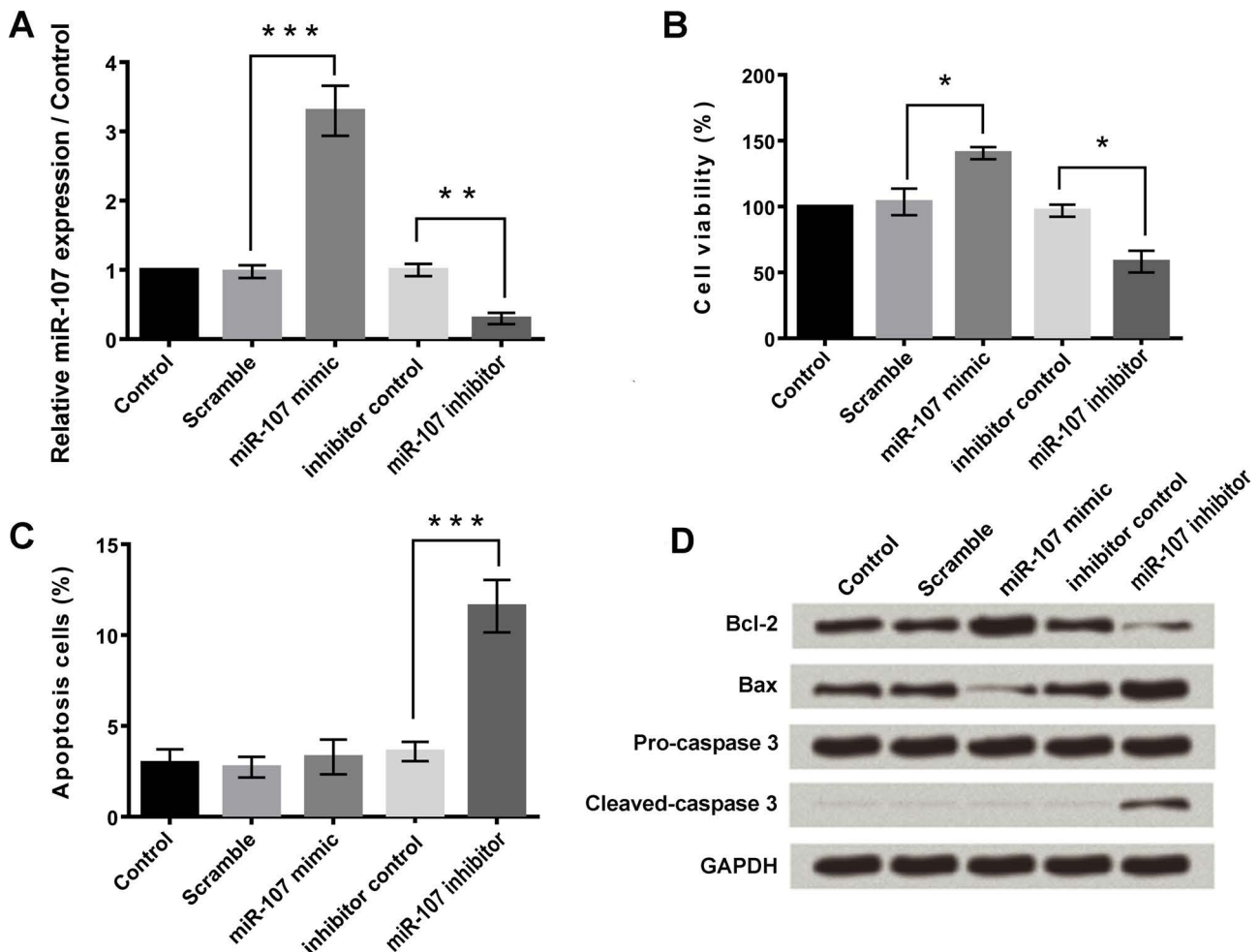


Figure 2. miR-107 influenced U2OS cell viability and apoptosis. U2OS cells were transfected with miR-107 mimic, miR-107 inhibitor, or scramble as the respective controls. (A) qRT-PCR analysis of miR-107 mRNA expression. (B) Cell counting kit-8 (CCK-8) assay of cell viability. (C) Annexin V/fluorescein isothiocyanate (FITC) analysis of apoptosis. Data represent the mean \pm SD of three independent experiments. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$. (D) Western blot analysis of apoptosis-related protein expressions. Numbers represent the expression relative to GAPDH as the internal control.

in U2OS cells had no effect on caspase 3 expression and resulted in increased levels of cleaved caspase 3. In contrast, miR-107 overexpression had no effect on the expression levels of either pro-caspase 3 or cleaved caspase 3. These data indicated that inhibition of miR-107 promoted cell apoptosis in U2OS cells.

miR-107 Influences U2OS Cell Migration and Invasion

The effects of miR-107 on the migration and invasion capacities of U2OS cells were investigated by Transwell assays with or without Matrigel, respectively. Cell migration was significantly promoted by overexpression of miR-107 and inhibited by its inhibitor ($p < 0.05$) (Fig. 3A). Similarly, the invasive capacity of U2OS cells was promoted by the overexpression of miR-107 and inhibited by its inhibitor ($p < 0.05$) (Fig. 3B). Transfection with scramble controls had no effect on either cell migration or invasion compared with the controls. Results suggested that miR-107 promoted U2OS cell migration and invasion.

TPM1 Was a Target of miR-107

We next investigated the effects of miR-107 on the expression of the tumor suppressor TPM1 in U2OS cells. TPM1 expression was significantly downregulated at both the mRNA and protein levels after transfection of the miR-107 mimic ($p < 0.05$) (Fig. 4A and B, respectively). In contrast, TPM1 expression was significantly upregulated at both the mRNA and protein levels following miR-107 inhibitor transfection ($p < 0.01$) (Fig. 4A and B, respectively). Transfection with scramble controls had no effect on TPM1

expression compared with that observed in the control cells without transfection.

The binding sites of miR-107 in TPM1 3'-UTR suggested that TPM1 was a potential target of miR-107, as per the prediction (Fig. 4C). The target effect of miR-107 was investigated by dual-luciferase reporter assays. We generated the U2OS cell line stably expressing the pMiR-reporter vector containing the wild-type 3'-UTR of the TPM1 gene carrying a putative miR-107 binding site (TPM1-wt) or a mutated version (TPM1-mt). Comparing the luciferase activity, miR-107 overexpression resulted in a significant reduction in luciferase activity of TPM1-wt cotransfected cells but not in TPM1-mt cells ($p < 0.05$) (Fig. 4C). These data indicated miR-107 downregulated expression of TPM1 and could target TPM1.

miR-107 Inhibitor Inhibited U2OS Cell Viability and Apoptosis via TPM1

To further elucidate the involvement of TPM1 in the effect of miR-107 on U2OS cells, we investigated the effects of the miR-107 inhibitor and/or TPM1 on cell viability and apoptosis in U2OS cells (Fig. 5). miR-107 inhibitor transfection resulted in a significant increase in TPM1 expression at both mRNA and protein levels ($p < 0.01$). This effect was abolished by concurrent transfection of both the miR-107 inhibitor and TPM1 siRNA ($p < 0.01$) (Fig. 5A and B, respectively). Furthermore, the decrease in U2OS cell viability mediated by the miR-107 inhibitor alone was significantly inhibited by concurrent transfection of the miR-107 inhibitor and TPM1 siRNA

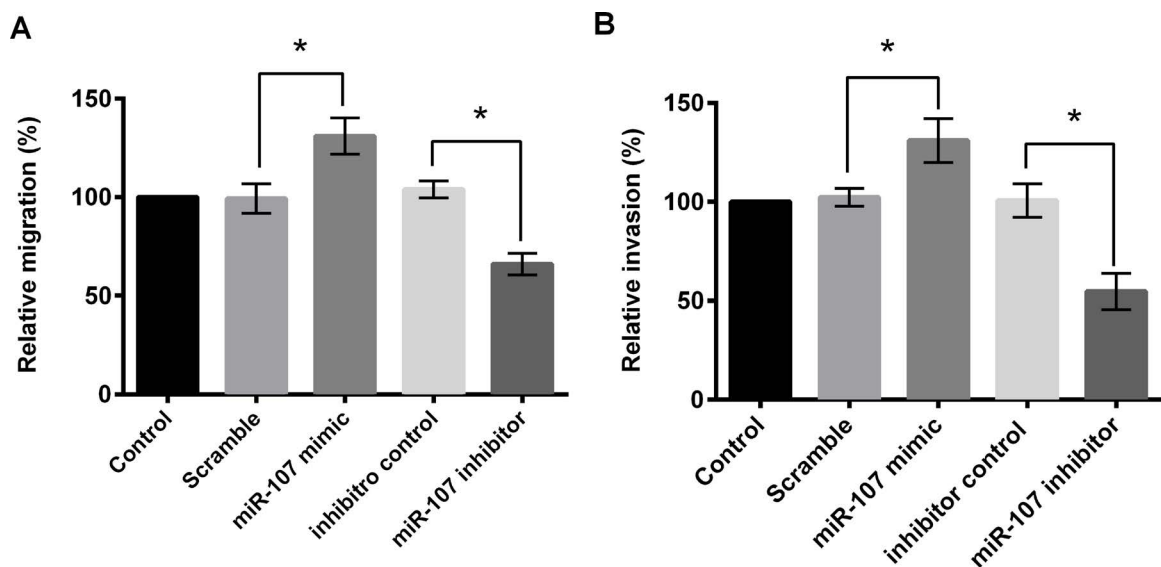


Figure 3. miR-107 influenced U2OS cell migration and invasion. U2OS cells were transfected with miR-107 mimic, miR-107 inhibitor, or scramble miRNA as the respective controls. (A) Transwell assay of migration. (B) Matrigel assays of invasion. Data represent the mean \pm SD of three independent experiments. * $p < 0.05$.

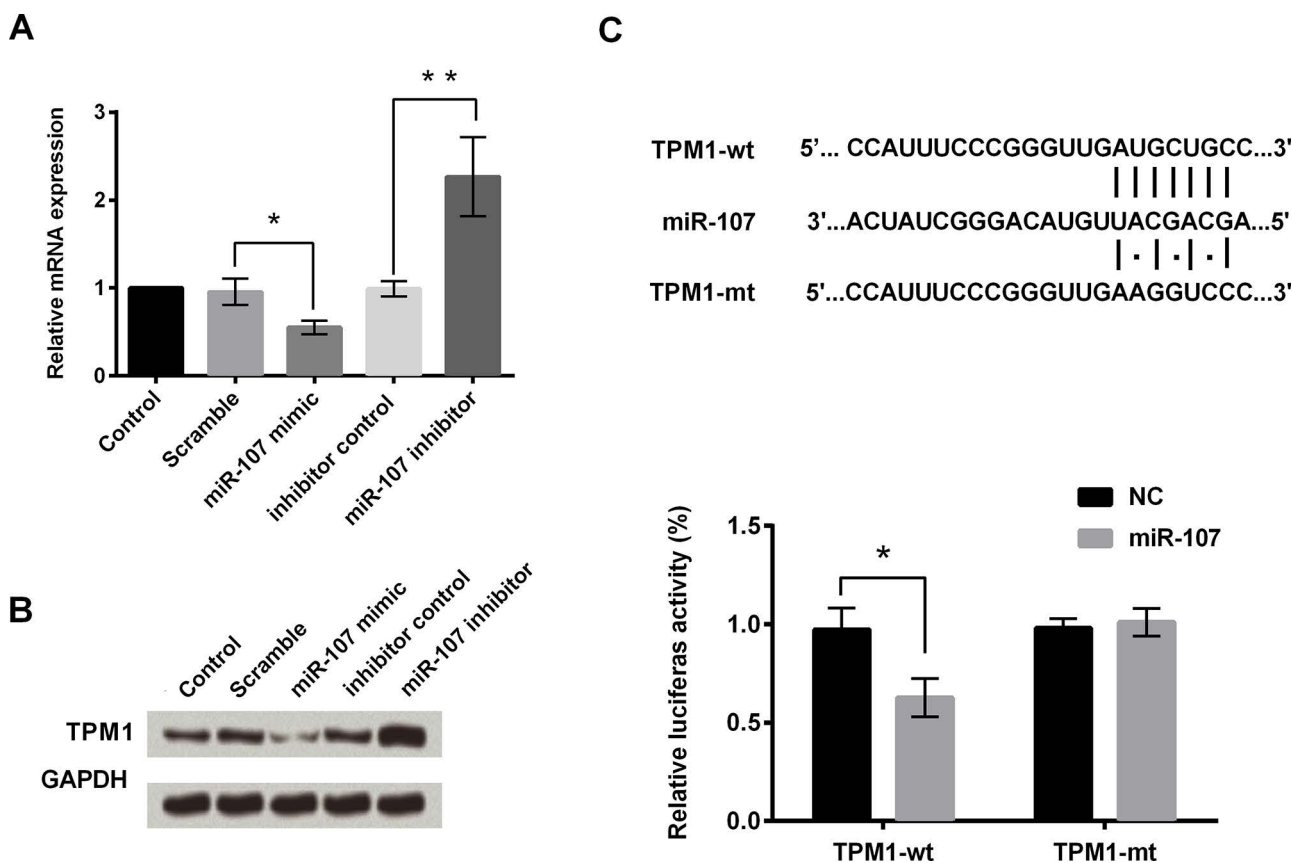


Figure 4. Tropomyosin 1 (TPM1) was a target of miR-107. U2OS cells were transfected with miR-107 mimic, miR-107 inhibitor, or scramble miRNA as the respective controls. The effect of miR-107 on the expression of the tumor suppressor TPM1 was analyzed by (A) qRT-PCR and (B) Western blotting. (C) The binding effect of miR-107 on TPM1 in U2OS cell stably expressing the pMiR-reporter vector containing the wild-type 3'-UTR of the TPM1 gene sequence carrying the putative miR-107 binding site (TPM1-wt) or a mutated version of TPM1 3'-UTR (TPM1-mt) was investigated by dual-luciferase reporter assays. Binding sequences were shown as TargetScan prediction. Data represent the mean \pm SD of three independent experiments. * p <0.05, ** p <0.01.

(p <0.01) (Fig. 5C). In contrast, the increased proportion of apoptotic cells observed following miR-107 inhibitor transfection alone was significantly inhibited by concurrent transfection of the miR-107 inhibitor and TPM1 siRNA (p <0.01) (Fig. 5D). Western blot analysis of apoptosis-related protein expressions showed that transfection of the miR-107 inhibitor alone in U2OS cells resulted in a decreased expression of the antiapoptotic protein Bcl-2, whereas expression of the proapoptotic protein Bax was markedly upregulated (Fig. 5E). In contrast, concurrent transfection of both the miR-107 inhibitor and TPM1 siRNA resulted in the upregulation of Bcl-2 and a downregulation of Bax. Transfection with the scramble control or siNC had no effect on the expression of these apoptosis-related proteins.

Compared with the controls, transfection of the miR-107 inhibitor alone in U2OS cells resulted in the increased expression level of cleaved caspase 3; this effect was weakened by cotransfection with both the

miR-107 inhibitor and TPM1 siRNA. In contrast, the miR-107 inhibitor and/or TPM1 siRNA in U2OS cells had no effect on cleaved caspase 3 expression. These results indicated that miR-107 inhibition decreased cell viability and promoted cell apoptosis in U2OS cells possibly through regulating TPM1.

miR-107 Inhibitor Influenced U2OS Cell Migration and Invasion via TPM1

In further experiments, to determine the role of TPM1 in the functions of miR-107, we investigated the effects of miR-107 inhibitor and/or TPM1 siRNA on the migration and invasion ability of U2OS cells (Fig. 6). The decreases in U2OS cell migration and invasion mediated by miR-107 inhibitor alone were significantly inhibited by concurrent transfection with both miR-107 inhibitor and TPM1 siRNA (p <0.01) (Fig. 6A and B). These results indicated that miR-107 inhibition decreased cell migration and invasion in U2OS cells possibly through regulating TPM1.

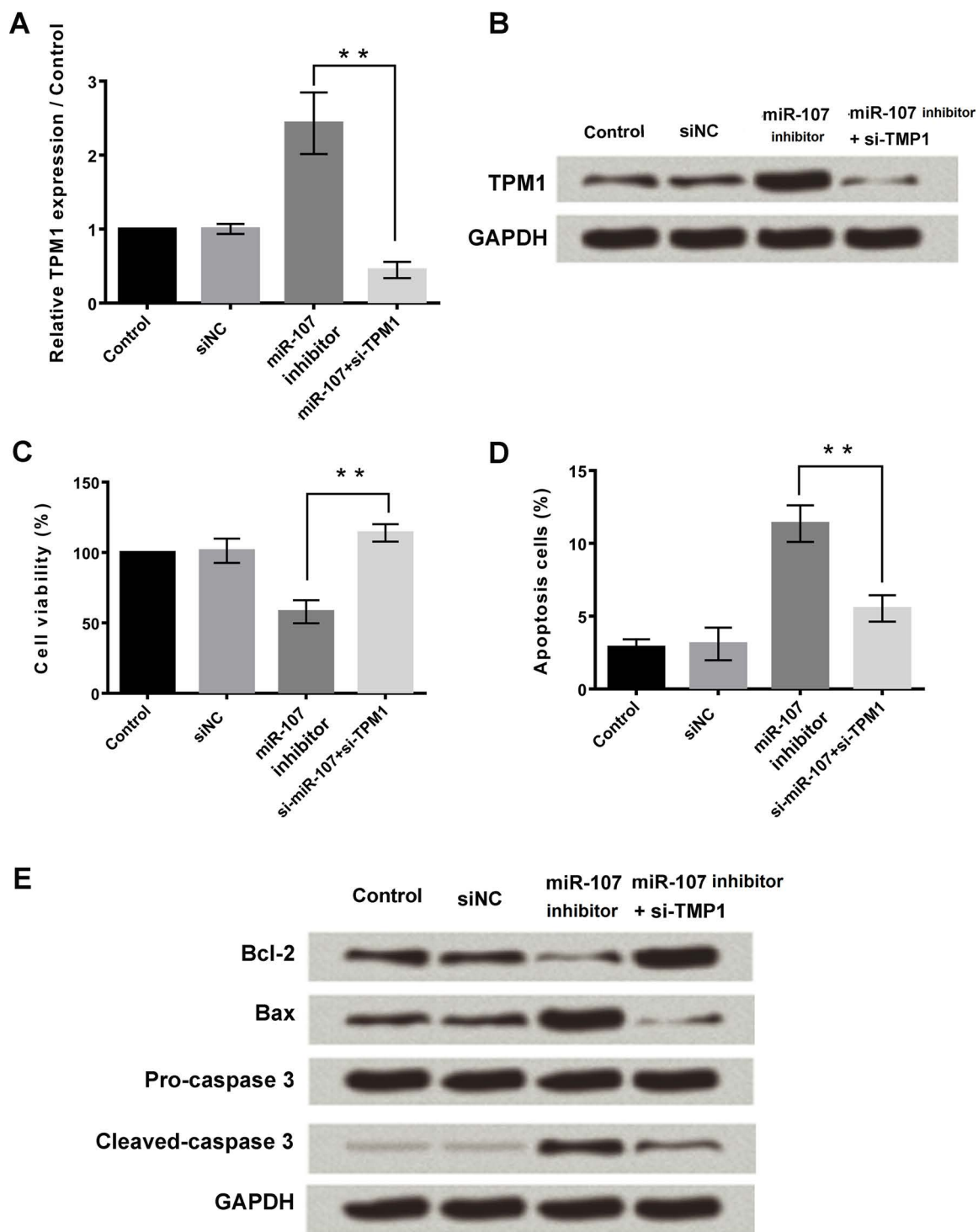


Figure 5. miR-107 suppression inhibited U2OS cell viability and apoptosis by inhibiting TPM1. U2OS cells were transfected with the miR-107 inhibitor and/or TPM1 siRNA (si-TPM1), or respective negative controls (siNC). Cells without transfection were used as controls. (A) qRT-PCR analysis of TPM1 mRNA expression. (B) Western blot analysis of TPM1 protein expression. Numbers represent the expression relative to GAPDH as internal control. (C) CCK-8 assay of cell viability. (D) Annexin V/FITC analysis of apoptosis. (E) Western blot analysis of apoptosis-related protein expressions. Data represent the mean \pm SD of three independent experiments. $**p < 0.01$.

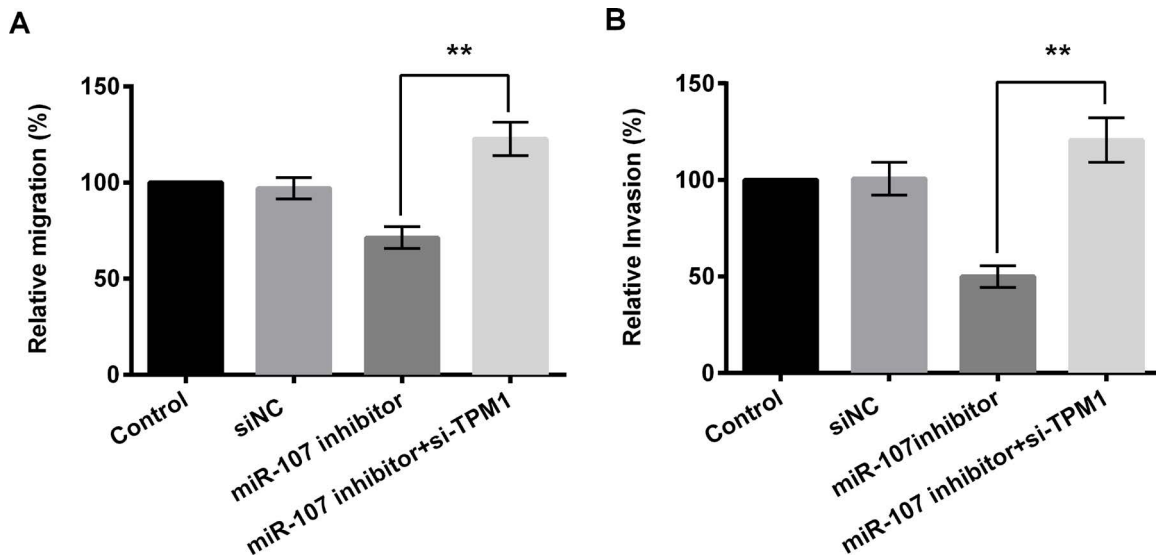


Figure 6. miR-107 suppression influenced U2OS cell migration and invasion via TPM1. U2OS cells were transfected with the miR-107 inhibitor and/or TPM1 siRNA (si-TPM1); respective negative controls (siNC) or cells without transfected were used as controls. (A) Transwell assay of migration. (B) Relative invasion of cells. Data represent the mean \pm SD of three independent experiments. ** $p < 0.01$.

miR-107 Overexpression Downregulated TPM1 and Activated MEK/ERK and NF- κ B Signaling Pathways

To clarify the signaling pathways involved in the effects on U2OS cells mediated by the interaction between miR-107 and TPM1, we investigated the expression of proteins related with the MEK/ERK and NF- κ B pathways by Western blot analysis (Fig. 7). miR-107 overexpression resulted in downregulated TPM1 expression, accompanied by the upregulated expression of both p-MEK and p-ERK (Fig. 7A). In contrast, miR-107 overexpression had no effect on the expression levels of inactive MEK and ERK. Transfection with the miR-107 inhibitor alone resulted in the upregulated expression of TPM1, and the downregulated expression of both p-MEK and p-ERK. These effects were abolished by concurrent transfection of both the miR-107 inhibitor and TPM1 siRNA. miR-107 overexpression resulted in downregulated TPM1 expression, followed by the upregulated expression of both p-I κ B α and p-p65 (Fig. 7B). In contrast, miR-107 overexpression had no effect on the expression level of inactive I κ B α and p65. Transfection of the miR-107 inhibitor alone resulted in the upregulated expression of TPM1, whereas expression of both p-I κ B α and p-p65 was downregulated. These effects were abolished by concurrent transfection of the miR-107 inhibitor and TPM1 siRNA. These observations showed miR-107 activated MEK/ERK and NF- κ B signaling pathways possibly through downregulating TPM1.

DISCUSSION

In this study, we investigated the expression of miR-107 in human osteosarcoma tissues and cells and the

effects of miR-107 on human osteosarcoma cells in vitro as well as its underlying mechanisms. We found that miR-107 expression was upregulated in both osteosarcoma tissues and osteosarcoma cell lines, suggesting that miR-107 might be involved in the pathogenesis of osteosarcoma. We demonstrated that miR-107 overexpression promoted cell viability of human osteosarcoma U2OS cells as well as migration and invasion abilities, whereas it inhibited apoptosis of cells. These effects were reversed by miR-107 inhibitor transfection. Further results showed that miR-107 overexpression promoted U2OS cell viability, migration, and invasion by activating the MEK/ERK and NF- κ B signaling pathways via downregulation of TPM1.

The role of miR-107 in human cancer appears to be context dependent. It has been proven to be a tumor suppressor in lung cancer and glioma^{19,20}, whereas oncogenic activity has been reported in breast cancer and gastric carcinoma^{15,21}. In this study, we found that miR-107 expression was significantly upregulated in human osteosarcoma tissues compared with the adjacent nontumor tissues. Furthermore, miR-107 was found to be upregulated in human osteosarcoma cell lines compared with human osteoblast cell line hFOB1.19. These results suggested that upregulated miR-107 expression might be a biomarker in the pathogenesis of osteosarcoma. This result was consistent with the upregulated miR-107 expression reported in other cancers, such as gastric carcinoma, pancreatic carcinoma, and breast cancer^{22,23}.

We further investigated the effect of miR-107 on U2OS cells by transfection with the miR-107 mimic or inhibitor. The efficiency of miR-107 overexpression or suppression was confirmed by qRT-PCR. Transfection of the miR-107

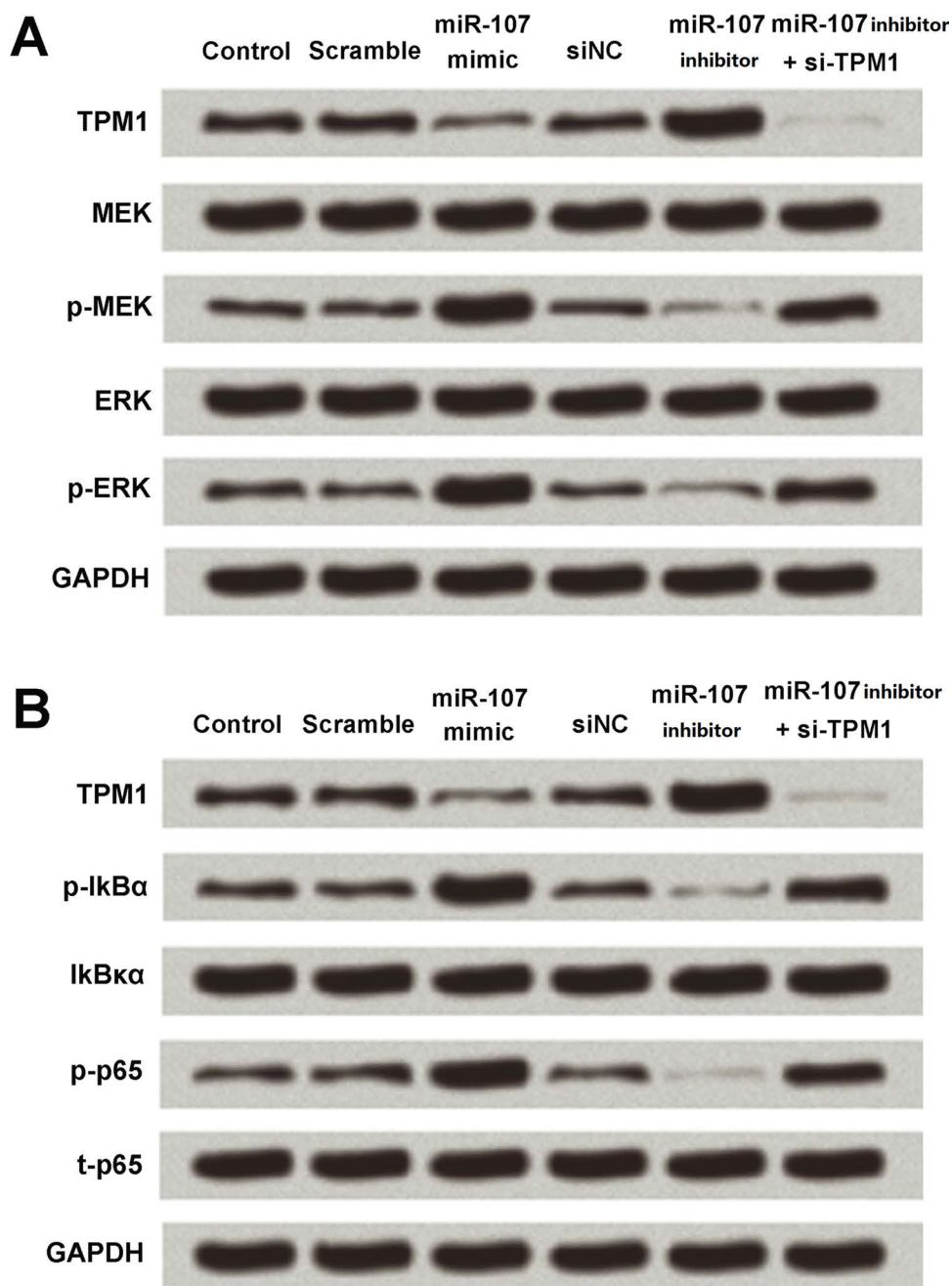


Figure 7. miR-107 overexpression downregulated TPM1 and activated MEK/ERK and NF- κ B signaling. U2OS cells were transfected with miR-107 mimic, miR-107 inhibitor, and/or TPM1 siRNA (si-TPM1); scramble miRNAs and siRNA negative control (siNC) or untransfected cells were used as controls. (A) Western blot analysis of expression of proteins related with the MEK/ERK pathway. (B) Western blot analysis of expression of proteins related with the NF- κ B signaling pathway. GAPDH acted as the internal control.

mimic increased U2OS cell viability, and this effect was abolished by the miR-107 inhibitor. These findings are consistent with the role of miR-107 in the development and progression of cancer. Furthermore, transfection of the miR-107 inhibitor was associated with significantly increased apoptosis, which was accompanied by increased expression of Bax and decreased expression of Bcl-2. In addition,

the miR-107 inhibitor had no effect on pro-caspase 3 expression, whereas a significant increase in cleaved caspase 3 was detected. Combined with the fact that Bax is a proapoptotic protein and Bcl-2 is an antiapoptotic protein, these results suggested that miR-107 overexpression might enhance cell viability and proliferation by influencing the balance of expression of apoptotic-related proteins.

We then conducted similar investigations to explore the effects of miR-107 on the migration and invasion abilities of U2OS cells. Overexpression of miR-107 resulted in the increased migration and invasion of U2OS cells, while these effects were inhibited by miR-107 knock-down. Thus, these observations indicate that miR-107 promotes the migration and invasion abilities of osteosarcoma cells. Similarly, several reports have demonstrated that miR-107 upregulation has a similar effect in gastric cancer cells^{15,24–26}.

miRNAs have the function of regulating gene expression by interacting with specific sequences in the 3'-UTR of their target genes. For instance, miR-107 has been shown to modulate the expression of the large tumor suppressor 2 (LATS2) gene and the neurofibromin 1 (NF1) gene in gastric cancer^{27,28}. These reports indicated that miR-107 regulated the development of gastric cancer by regulating the expressions of multiple target genes. TPM proteins are a vast family of actin-binding proteins encoded by four different genes, and all four genes generate multiple isoforms. The function of TPM isoforms is poorly understood. TPM1 has been reported to function as a tumor suppressor^{16–18}. In this study, we found that TPM1 expression was significantly decreased at both the mRNA and protein levels after miR-107 mimic transfection, whereas the opposite effects were observed after transfection of the miR-107 inhibitor. The dual-luciferase reporter assay results showed that miR-107 targets the 3'-UTR of the TPM1. To further clarify this interaction, we investigated the effects of miR-107 and/or TPM1 knockdown on U2OS cells. Although miR-107 inhibitor transfection alone significantly decreased U2OS cell viability and increased apoptosis, these effects were inhibited by cotransfecting with TPM1 siRNA. Furthermore, transfection of miR-107 inhibitor alone decreased Bcl-2 expression and expressions of Bax and cleaved caspase 3, and the opposite trends of expression change were observed following concurrent TPM1 knockdown. Similarly, the decreases in U2OS cell migration and invasion mediated by transfection of the miR-107 inhibitor alone were significantly inhibited by concurrent siRNA-mediated knockdown of TPM1. Based on these results, it can be speculated that miR-107 promoted cell viability, migration, and invasion, and inhibited cell apoptosis in osteosarcoma cells by targeting TPM1. Further analysis of the signaling pathways involved in the effects of miR-107 and TPM1 on U2OS cells suggested that the inhibitory effect of miR-107 on TPM1 expression was associated with activation of the MEK/ERK and NF- κ B signaling pathways.

The limitations of this study should be noted. First, we evaluated the expression of miR-107 in only 25 clinical specimens; therefore, our results showing that miR-107 was upregulated in human osteosarcoma sarcoma tissues should be confirmed by evaluation of a greater

number of clinical samples. Second, in vivo studies are required to confirm the role of miR-107 in the pathogenesis of osteosarcoma.

In conclusion, the results of this study demonstrate that miR-107 is upregulated in human osteosarcoma tissues and cell lines. Furthermore, we showed that miR-107 promotes cell viability and enhances the migration and invasion abilities of U2OS cells, which might be related to activating the MEK/ERK and NF- κ B signaling pathways via targeted tumor suppressor TPM1. These findings provide an improved understanding of the mechanism by which miR-107 is involved in the pathogenesis of osteosarcoma, and it is critical for the development of novel therapies for osteosarcoma.

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