MicroRNA-382-5p inhibits osteosarcoma development and progression by negatively regulating VEZF1 expression

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Abstract. Human osteosarcoma is the most frequent malignant primary bone tumor that mainly occurs in young adults and children. MicroRNAs (miRNAs/miRs) are abnormally expressed in human osteosarcoma and contribute to osteosarcoma initiation and development. The present study aimed to investigate the role of miR-382-5p in the nosogenesis of osteosarcoma and to identify a novel target for osteosarcoma treatment. miR-382-5p expression was detected in human osteosarcoma clinical tissues and cell lines, including 143B, U2OS and MG63, via reverse transcription-quantitative PCR analysis. Multiple bioinformatic prediction tools were used to identify the potential target genes of miR-382-5p and vascular endothelial zinc finger 1 (VEZF1), which were validated via the dual-luciferase reporter assay. MG63 and U2OS cells were transfected with miR-382-5p mimics. The Cell Counting Kit-8 assay was performed to assess cell proliferation, while the Transwell assay was performed to assess migration and invasion. Cell colony formation was measured via crystal violet staining, and apoptosis was assessed via Annexin V/propidium iodide staining. The wound healing assay was performed to assess the migratory ability of U2OS and MG63 cells. Antitumor effects of miR-382-5p were evaluated in nude mice xenografts using U2OS cells. The results demonstrated that miR-382-5p expression was markedly downregulated in human osteosarcoma tissues and cell lines compared with adjacent normal tissues. Transfection of miR-382-5p mimics into MG63 and U2OS cells significantly inhibited the malignant behaviors of cells, including decreased proliferation, migration, diminished colony formation and invasion, and promoted osteosarcoma cell apoptosis. Bioinformatics prediction indicated that VEZF1 is a direct target gene of miR-382-5p. Overexpression of VEZF1 restored osteosarcoma tumor development inhibited by miR-382-5p in vivo. In addition, overexpression of miR-382-5p restrained the growth of xenograft osteosarcoma in nude mice following co-transfection, and overexpression of VEZF1 attenuated the inhibitory effect of miR-382-5p in nude mice. miR-382-5p acted as a tumor suppressor gene and inhibited the malignant biological behaviors of human osteosarcoma cells and functions associated with directly targeting VEZF1. Taken together, these results suggest that the miR-382-5p/VEZF1 interaction has an important role in osteosarcoma development and progression, and thus may be used as a diagnostic and therapeutic target for osteosarcoma.

Introduction

Human osteosarcoma is the most frequent malignant bone tumor, and its characteristics of malignant proliferation and invasion have made it the leading cause of cancer-associated mortality among teenagers and children (1-3). The incidence of osteosarcoma is approximately 4-5 cases per million, which accounts for >60% of all bone malignancies and ~20% of all primary osteosarcomas in children >5 years old (4). Surgery and chemotherapy are the most common treatment methods for osteosarcoma (5). With improvements in therapeutic technologies, the 5-year survival rate of newly diagnosed patients with osteosarcoma without metastasis has improved...
to ~60-70% (6). However, the 5-year survival rate of patients with osteosarcoma is only 20% because of the frequency of recurrent or metastatic disease (7). Thus, it is important to understand the molecular mechanisms underlying the initiation, progression, invasion and recurrence of osteosarcoma to identify novel therapeutic targets and develop rational strategies for clinical treatment of osteosarcoma.

MicroRNAs (miRNAs/miRs) are small (17-25 nucleotides) non-coding RNAs (8). These molecules are highly conserved endogenous RNAs that have an important role in post-transcriptional regulation of mRNAs for translation, and influence diverse biological activities (9). Although only the biological functions of a few miRNAs have been elucidated, miRNAs have been associated with osteosarcoma initiation, progression and invasion (10,11). Increasing evidence suggest that miRNAs may perform gene regulation in different types of cancer, such as direct regulation of target genes, including oncogenes or tumor suppressor genes (12). For example, miR-142-3p has been demonstrated to inhibit osteosarcoma cell proliferation by targeting Rac1 (13). In addition, miR-142-3p has been reported to be highly overexpressed in osteosarcoma cells, to have pivotal roles in cellular invasion, proliferation, migration and apoptosis, and to induce E-cadherin expression and decrease expression of matrix metalloproteinase (MMP)2 and MMP9 (13). Thus, miRNAs are considered potential biomarkers and therapeutic targets for osteosarcoma.

Increasing evidence suggest that miR-382-5p is enriched in the serum of patients with ischemic stroke (14). Notably, serum miR-382-5p may be used as a non-invasive biomarker for potential diagnosis of ischemic stroke (15). In addition, aberrant miR-382-5p expression is frequently observed in different types of human cancer, including oral, breast cancer, ovarian, prostate, colorectal, lung and glioblastoma (16-19). However, the underlying molecular mechanisms and functional role of miR-382-5p in osteosarcoma tissues and cell lines remain largely unclear.

The present study aimed to investigate the biological effects of miR-382-5p on osteosarcoma cell migration, invasion, proliferation and apoptosis, both in vitro and in vivo. The results demonstrated that miR-382-5p expression was markedly downregulated in human osteosarcoma tumor tissues and cell lines, including MG63 and U2OS cells. In addition, overexpression of miR-382-5p inhibited malignant biological behaviors, including colony formation, proliferation, invasion and migration of osteosarcoma cell lines. Further studies demonstrated that miR-382-5p suppressed osteosarcoma development and progression by targeting VEZF1, and its aberrant expression remarkably reversed the antitumor effects of miR-382-5p overexpression in human osteosarcoma cell lines in vitro. Both the in vivo and in vitro experiments provide novel insights into the molecular functions of miR-382-5p in human osteosarcoma, which may be used as a potential therapeutic target for patients with osteosarcoma.

Materials and methods

Clinical samples and cell culture. A total of 20 paired (13 men and seven women; age range, 22-48 years; mean age, 33.8 years) adjacent normal tissues (within 5 mm from the tumor boundary) and osteosarcoma tumor tissues were collected from patients who underwent surgical resection at Nanchong Central Hospital (Nanchong, China) between January 2018 and December 2019. Resected tissues were immediately frozen in liquid nitrogen and stored at -80°C until subsequent experimentation. The present study was approved by the Institutional Center Ethics Review Committee of Nanchong Central Hospital (Nanchong, China; approval no. 2020-002) and written informed consent was provided by all patients or their guardians prior to the study start.

Human osteosarcoma cell lines, including HFOB, HOBC, 143B and U2OS were purchased from The Cell Bank of Type Culture Collection of the Chinese Academy of Sciences, while MG63 cells were purchased from the American Type Culture Collection (ATCC, cat. no. ATCC® CRL-1690™) and U2 osteosarcoma (ATCC® CRL-2611™) were purchased from the indicated vendors and maintained in Dulbecco's Modified Eagle's Medium (DMEM, Gibco; Thermo Fisher Scientific, Inc.) supplemented with 10% fetal bovine serum (FBS; AccurRef Scientific, https://www.accurerefsci.com), 100 IU/ml penicillin and 100 µg/ml streptomycin (all from Thermo Fisher Scientific, Inc.), at 37°C with 5% CO2.

Transfection of miRNA mimics and plasmids. Human miR-382-5p mimics (50 nM, 5'-GAAGUUGUCCUGGU GGAUUGG-3'), antisense or negative controls (50 nM, NC-mimic; 5'-CTCGCTTCCGACGACA-3') were purchased from Shanghai GenePharma Co., Ltd. The DNA coding sequence for VEZF1 was cloned into the pcDNA™3.1 (+) expression vector (Thermo Fisher Scientific, Inc.,), and the empty vector pcDNA3.1 was used as the NC. Osteosarcoma cell lines (MG63 and U2OS) in logarithmic growth phase were maintained in RPMI-1640 medium (Gibco; Thermo Fisher Scientific, Inc.) supplemented with 10% FBS. Cells were incubated at 37°C with 5% CO2 in a sterility incubator and cultured for 24 h. Cells were transfected with miR-382-5p mimics using the Lipofectamine® 3000 kit (Thermo Fisher Scientific, Inc.), according to the manufacturer's protocol. Cells were collected for subsequent experimentation 48 h post-transfection.

Lentivirus packaging and infection. The 2nd lentiviral system purchased from Shanghai Genechem Co., Ltd., was used for lentivirus generation. Lentiviral particles were produced in Lenti-X™ 293T cells (Takara Bio Inc.; cat. no. 632180) in 10-cm dish by transiently co-transfecting control lentiviral vector (4.5 µg) or miR-382-5p overexpressing lentiviral vector (4.5 µg) together with helper plasmids pHelper 1.0 (Gag and Pol; 2.5 µg) and pHelper 2.0 (VSVG; 2.0 µg) using the Lipofectamine® 3000 kit (Thermo Fisher Scientific, Inc.), according to the manufacturer's protocol. The vector constructions, verification by sequencing, virus packaging and collection of the corresponding viral supernatants were performed by Shanghai Genechem Co., Ltd. The lentivirus in medium of 293T cells were harvested by centrifuging at 75,000 x g for 1.5 h at 4°C 48 h post-transfection (Abcam), according to the manufacturer's protocol. Cells in the logarithmic growth phase were seeded into 96-well microplates (Costar; Corning, Inc.) at a density of 2x104 cells/well. Subsequently, the medium was replaced with DMEM supplemented with 10% FBS after overnight incubation at 37°C. Cells were incubated for an
additional 24, 48, 72, and 96 h, respectively. At each time point, 100 µl CCK-8 reagent was added to each well and cells were incubated for an additional 2 h at 37°C. Against a background control, the sample absorbance was measured at a wavelength of 490, using a microplate reader (Bio-Rad Laboratories, Inc.).

Apolipoprotein assay. MG63 and U2OS cells seeded into 6-well plates were transfected with the indicated miRNA mimics or the indicated plasmids and incubated at 37°C for an additional 24 h. Cells were digested with 0.25% trypsin-EDTA solution (Sigma-Aldrich; Merck KGaA), terminated with cell solution, and harvested via centrifugation at 300 x g for 5 min at 4°C. After washing with PBS, the Annexin V-fluorescein isothiocyanate (FITC)/PI Apoptosis kit (BD Biosciences) was used to detect cell apoptosis according to the manufacturer's protocols. Cells were washed three times with PBS and the binding buffer, and stained with Annexin V-FITC/PI for 15-20 min at room temperature in the dark. Cells were re-washed and the labeled cells were detected using a flow cytometer (CytoFLEX; Beckman Coulter, Inc.). Flow cytometry data were analyzed using Expo32 software (version 1.2; Beckman Coulter, Inc.).

Cell migration and invasion assays. Transwell® Boyden chambers (BD Biosciences) were used for in vitro cell migration assays, and chambers coated with or without Matrigel (BD Biosciences) were used to assess the migratory and invasive abilities of MG63 and U2OS cells in vitro. For pre-coating, Matrigel was maintained at 4°C overnight and then diluted with serum-free DMEM at a dilution of 1:4 on ice. Subsequently, Matrigel was added to each upper chamber for the lower chamber. Following incubation at 37°C for 1 h, the chamber was washed with serum-free DMEM followed by 50 µl of DMEM supplemented with 10 g/l BSA at 37°C. Cells were resuspended in FBS-free medium 48 h post-transfection. Cells were plated at a density of 1x10^5 cells/well into the upper chamber, and 200 µl complete medium was loaded into the lower chamber. Following incubation at 37°C for 24 h, the migratory and invasive cells on the lower surface of the membrane were fixed with 4% formaldehyde at 4°C for 1 h and stained with 0.1% crystal violet (Sigma-Aldrich; Merck KGaA) at room temperature for 15 min, followed by a light microscopy (Nikon E600; Nikon Instrument Inc.) for visualization and photography. Stained cells were counted in five randomly selected fields at low-power (x100) magnification.

Dual-luciferase reporter assay. Given that miRNAs potentially interact with VEZF1, the binding sites between VEZF1 and miR-382-5p were predicted using IncBase version 2.0 (20). Candidate miRNA genes were targeted by miR-382-5p, and the binding sites of miR-382-5p were predicted using the online tools StAR MirDB (http://sfold.wadsworth.org/starmirdb.php) and miRWalk (http://mirwalk.umm.uni-heidelberg.de). The predicted binding sites within the 3'-untranslated region (3'UTR) of VEZF1 mRNA (5'-AGGACAAUA AAGUG ACAACUUU-3') and the corresponding mutated binding region (5'-AGGACAAUA AAGUGUCCACGGU-3') were cloned into a luciferase-expressing vector pCDNA3.1. For the luciferase reporter assay, MG63 cells were transfected with different combinations of miR-382-5p mimics, control mimics, and pCDNA3.1-VEZF1 3'-UTR wild-type or mutant using the Lipofectamine® 3000 kit (Thermo Fisher Scientific, Inc.), according to the manufacturer's protocol. Following transfection for 48 h, cells were collected and luciferase activities were detected using the dual-luciferase reporter assay system (Promega Corporation) according to the manufacturer's instructions. Firefly luciferase activity was normalized to Renilla luciferase activity.

Reverse transcription-quantitative (RT-q)PCR. According to the manufacturer’s protocols, total RNA and miRNA were extracted from osteosarcoma tissues and cell lines using TRIzol® reagent (AccuRef Scientific) and the miRNeasy mini kit (Qiagen GmbH), respectively. The PrimeScript® RT Master Mix Perfect Real-Time Reagent kit (Takara Bio, Inc.) was used to synthesize cDNA from total RNA, according to the manufacturer’s instructions. For miRNA RT, the corresponding cDNA was synthesized using a universal tag (miScript II RT kit; Qiagen GmbH). qPCR for miRNA and mRNA was subsequently performed using a standard protocol from the SYBR Green PCR kit (AccuRef Scientific) on an AB7500 RT-PCR instrument (Applied Biosystems; Thermo Fisher Scientific, Inc.) The following thermocycling conditions were used for qPCR: 95°C for 10 min, and 40 cycles of 95°C for 10 sec, 60°C for 30 sec and 72°C for 10 sec. Relative quantification was determined by normalization to U6 or GAPDH. The following primer sequences were used for qPCR: VEZF1 forward, 5'-GCC TCA ACT CGAC AGGAGGA AG-3' and reverse, 5'-TCC AAA GTACT CTT GGT GCT CC-3'; AKIRINI forward, 5'-GCAC CT AGCT CTCC AGA AACA-3' and reverse, 5'-CTTGG CCA TATT CCA GC-3'; SLAINI forward, 5'-TTT TCCC CAGTGT TTG TTCGAG-3' and reverse, 5'-CCTT GGAGTA GTG AAC GCT T-3' and reverse, 5'-TTCAT CCGT CTCA A AACTGT-3'; LMTK3 forward, 5'-GTGTA AAT GTCG TAAC CGC-3' and reverse, 5'-CCC CCA GTT GAC AGA ACT CC-3'; GEN2X forward, 5'-GGGCGTGA AAT CTCG TGGT AAGG-3' and reverse, 5'-TCCTAT GTGGT GTT CCA AACTGT-3'; TOP1 forward, 5'-GT TCA AGC CAG CAC GAG AAG-3' and reverse, 5'-GCGTT GGT AAG C CGT TAC GAC AAG-3'; AMOTL2 forward, 5'-GGGTGATT CTC G GTT GCT-3' and reverse, 5'-GTA GCCG TGG TTA GCTG CA-3'; FOXN2 forward, 5'-GGGCGTGA AAT CTCG TGGT AAGG-3' and reverse, 5'-TCCTAT GTGGT GTT CCA AACTGT-3'; LMTK3 forward, 5'-GTGTA AAT GTCG TAAC CGC-3' and reverse, 5'-CCC CCA GTT GAC AGA ACT CC-3'; GAPDH forward, 5'-GTC TCC TCT GAC TTC CCA A-3'; miR-382-5p forward, 5'-ATC CGT GAA GTT GTT CTT GGT GAC GTT CT-3' and reverse, 5'-ATCCGTGAA GTT GAC GTT CT-3'; U6 forward, 5'-CGC TTC ACG AAT TTG CGT-3'; and U6 forward, 5'-CGC TTC ACG AAT TTG CGT-3'.

Relative expression levels were calculated using the 2^(-ΔΔCq) method (21).

Western blotting. Protein samples from osteosarcoma tumor tissues and cell lines were isolated using RIPA lysis buffer (Sigma-Aldrich; Merck KGaA) and subsequently centrifuged at 12,000 x g for 30 min at 4°C. Total protein was quantified using the BCA method (Thermo Fisher Scientific, Inc.). Equal amounts of protein samples (20 µg/lane) were separated via 10% SDS-PAGE, transferred onto PVDF membranes and
blocked with 5% skimmed milk at room temperature for 1 h. The membranes were incubated with primary antibodies against VEZF1 (1:1,000; cat. no. SAB2102675; Sigma-Aldrich; Merck KGaA) and β-actin (1:10,000; cat. no. MFCD00164531; Sigma-Aldrich; Merck KGaA) at 4°C overnight. All primary antibodies were diluted in a Primary Dilution Solution (AccuRef Scientific) prior to incubation. Following the primary incubation, membranes were incubated with secondary HRP-conjugated polyclonal antibody anti-rabbit IgG (1:10,000; cat. no. MFCD00163923; Sigma-Aldrich; Merck KGaA) at room temperature for 1 h. Protein bands were visualized using the Enhanced Chemiluminescence kit (Millipore, Sigma) and qualified using ImageJ software (version 1.49, National Institutes of Health).

Colony formation assay. Following trypsinization, MG63 and U2OS cells were suspended in medium that included 0.3% agar (low melt, Bio-Rad Laboratories, Inc.) and 10% FBS (AccuRef Scientific), seeded into 12-well plates and subsequently transfected with the indicated miRNA mimics or the indicated plasmids followed by culturing for an additional 24 h. Subsequently, 500 cells of each group were seeded into 35-mm dishes, and the medium was replaced with fresh complete DMEM every 2 days. Following culturing for 14 days at 37°C in the incubator and washing three times with PBS, the cells were fixed with 4% paraformaldehyde for 10 min at room temperature. Cells were subsequently stained with 0.1% crystal violet (Sigma-Aldrich; Merck KGaA) for 20 min at room temperature and washed with tap water. Cells were observed under a light microscope (Nikon E600; Nikon Instrument Inc.) with x200 magnification. All experiments were performed in triplicate.

Human osteosarcoma xenograft model. A total of 15 female BALB/c nude mice (4-6 weeks old and weighed 18-20 g) were obtained from Taconic Biosciences Company and acclimated in a specific-pathogen-free facility at the Experimental Animal Center of Nanchong Central Hospital, with a 12/12 h light/dark cycle, 40-60% humidity, 24-26°C temperature condition and free access to food and water for ≥1 week. Stable NC lentivirus and stable miR-382-5p-expressing lentivirus-infected MG63 cells were resuspended in normal saline and mixed with an isopycnic of Matrigel® (BD Biosciences) at a final concentration of 5x10⁶ cells/ml, respectively. Cells were injected into the left flank of nude mice subcutaneously at a density of 1x10⁶ cells/mouse. Tumor size was measured every 7 days using vernier calipers. Following tumor cell inoculation, the mice were euthanatized at day 21 using CO₂ inhalation with a flow rate of 3 l/min and euthanasia was confirmed via cardiac and respiratory arrest. The tumors were subsequently isolated, weighed, imaged and subjected to subsequent analyses. All animal experiments were performed in accordance with the Guide for the Care and Use of Laboratory Animals (National Academies Press, 2011; Washington, USA). The animal experimental protocol was approved by the Ethical Review Committee for Animal Experiments of Nanchong Central Hospital (Nanchong, China; approval no. NSMC-2020-42), and all experiments were performed according to the AVMA guidelines.

Immunohistochemistry. While the tumors were pictured and weighed, partial tumor tissues were collected and cut into 5 mm³ size. Subsequently, tissues were fixed with 4% formaldehyde at 4°C more than 12 h. Then, tissues were paraffin-embedded at 60°C overnight and cut into 5-μm thick slices. Tumor tissues from xenograft mice were deparaffinized and rehydrated in an ethanol series. Antigen retrieval was performed according to previous experimental protocols (22). After washing with PBS and subsequent blocking with 3% hydrogen peroxide at room temperature for 10 min, the sections were incubated with rabbit-anti-Ki-67 polyclonal antibody (1:400; cat. no. PA5-19462; Thermo Fisher Scientific, Inc.) overnight at 4°C. Sections were washed three times with PBS, and the samples were incubated with HRP-conjugated secondary donkey-anti-rabbit monoclonal antibody (1:200; cat. no. ab6802; Abcam) at room temperature for 1 h. Samples were re-washed five times with PBS and subsequently stained with 0.5% dianaminobenzidine and counterstained with Mayer’s hematoxylin at room temperature for 2-3 min. The mounted slices were observed under a light microscope (Nikon E600; Nikon Instrument Inc., magnification, x200).

Wound healing assay. U2OS and MG63 cells were seeded into 6-well plates at a density of 2x10⁶ cells/well and allowed to adhere for 24 h. Mimics were introduced into the cells and monolayer cells were scratched using a 200 μl pipette tip after 24 h. The medium was replaced with fresh serum-free RPMI-1640 medium. Wounded areas were observed at 0 and 48 h after culturing under a light microscope (Nikon E600; Nikon Instrument Inc., magnification, x100). The wound-healing rate (%) was calculated using the following equation: Wound-healing rate (%) = (width at 0 h - width at 48 h) / width at 0 h) x100. All experiments were performed in triplicate.

Statistical analysis. All quantitative data are presented as the mean ± standard deviation. Normal distribution of all data were confirmed using the Kolmogorov-Smirnov D test, and differences between unpaired two groups were compared using unpaired two-tailed Student’s t-test. Paired two-tailed Student’s t-test was used to compare differences between adjacent tissues and tumor tissues. Comparisons among multiple groups were performed using ANOVA analysis. All experiments were performed in triplicate. SPSS v.19.0 software (IBM Corp.) was used to assess the correlation between miR-382-5p and VEZF1 mRNA expression, using Pearson’s correlation analysis. P<0.05 was considered to indicate a statistically significant difference.

Results

miR-382-5p expression is downregulated in osteosarcoma tumor tissues and cell lines. To investigate the role of miR-382-5p in osteosarcoma, RT-qPCR analysis was performed to detect miR-382-5p expression in 20 paired human osteosarcoma tissues and adjacent normal tissues. The results demonstrated that miR-382-5p expression was significantly downregulated in osteosarcoma tumor tissues compared with adjacent normal tissues (P<0.01; Fig. 1A). In addition, miR-382-5p expression was detected in human osteosarcoma cell lines. The results demonstrated that miR-382-5p expression was significantly downregulated in human osteosarcoma cell lines (143B, MG63 and U2OS) compared with the normal
HFOB and HOBC cell lines (P<0.01; Fig. 1B). Notably, MG63 and U2OS cells exhibited ~80% reduction of miR-382-5p expression (Fig. 1B). Thus, these cell lines were selected for subsequent experimentation.

**Overexpression of miR-382-5p inhibits proliferation, invasion and migration of osteosarcoma cells, and promotes apoptosis.** According to the identifications in Fig. 1, downregulated miR-382-5p expression in human osteosarcoma tissues and osteosarcoma cell lines suggests that miR-382-5p may have important antitumor functions in the initiation and progression of osteosarcoma. Thus, the effects of miR-382-5p overexpression on the malignant behaviors of human osteosarcoma cells were investigated, including colony formation, proliferation, invasion and migration. MG63 and U2OS cells were transfected with miR-382-5p mimic or a NC miRNA mimic, and a 40-fold increase in miR-382-5p expression was observed at 48 h in miR-82-5p mimic-transfected MG63 and U2OS cells (P<0.01; Fig. 2A). We then compared the proliferation of these cells by performing CCK-8 assays. In addition, overexpression of miR-382-5p significantly impeded the proliferation of both MG63 and U2OS cells (both P<0.01; Fig. 2B). A nearly 50% decrease in OD_{scan} absorbance values was observed in miR-382-5p mimic-transfected MG63 and U2OS cells at 72 h after culturing. Furthermore, similar reductions in colony numbers were also observed in MG63 (P<0.01; Fig. 2C) and U2OS (P<0.01) cells with ectopic miR-382-5p expression, which suggests that miR-382-5p notably decreases the colony formation ability of human osteosarcoma cells. Notable reductions in the migratory (P<0.01; Fig. 2D) and invasive (P<0.01; Fig. 2E) abilities of both MG63 and U2OS cells were also observed following transfection with miR-382-5p mimics. In addition, overexpression of miR-382-5p significantly elevated the apoptotic rates of MG63 and U2OS cells (both P<0.01; Fig. 2F). Taken together, these results suggest that miR-382-5p exhibits tumor-suppressive functions in human osteosarcoma cells.

miR-382-5p directly targets VEZF1 in osteosarcoma cells. To further investigate the molecular mechanisms underlying the tumor-suppressive function of miR-382-5p in osteosarcoma, bioinformatics analysis was performed, using STarMiRDB and miRWalk (23,24), to predict the target genes based on the consensus binding sites of miR-382-5p. In the present study, the potential targets of miR-385-5p were predicted using ENCORI and TargetScan databases and the overlapped targets were screened using a Venn analysis (Fig. 3A). A total of nine overlapped targets of miR-382-5p were identified, and overexpression of miR-382-5p significantly inhibited the expression levels of VEZF1, FOXN2, and AMOTL2, particularly VEZF1 (P<0.01; Fig. 3B). The dual-luciferase reporter assay was performed to determine whether VEZF1 is a direct target of miR-382-5p in osteosarcoma. The results demonstrated that luciferase activity significantly decreased following co-transfection of wild-type VEZF1 3'-UTR and miR-382-5p mimics but remained unchanged following co-transfection with mutated VEZF1 3'-UTR and miR-382-5p mimics or NC in MG63 cells (P<0.01; Fig. 3C). These results suggest direct binding of miR-382-5p to the predicted 3'-UTR of VEZF1.

RT-qPCR and western blot analyses were performed to further verify the association between miR-382-5p and VEZF1 expression in osteosarcoma tissues and adjacent normal tissues, and the respective cell lines. The results demonstrated that the mRNA and protein expression levels of VEZF1 were significantly lower in adjacent normal tissues compared with osteosarcoma tissues (P<0.01; Fig. 3D). Similarly, the mRNA and protein expression levels of VEZF1 were significantly lower in normal cells compared with osteosarcoma cells (P<0.01; Fig. 3E). Furthermore, mRNA VEZF1 expression was inversely correlated with miR-382-5p expression in osteosarcoma tissues (r=-0.2182; P<0.01; Fig. 3F).

VEZF1 mRNA expression significantly decreased in MG63 cells transfected with miR-382-5p mimic compared with the control group (P<0.05; Fig. 3G). Furthermore, western blot analysis demonstrated that VEZF1 protein expression significantly decreased in MG63 cells transfected with miR-382-5p mimic compared with the control group (Fig. 3H). Collectively, these results suggested that VEZF1 is a direct downstream target of miR-382-5p in human osteosarcoma.

**Overexpression of VEZF1 enhances cell proliferation, invasion and migration, while inhibiting apoptosis in osteosarcoma.**
To determine whether VEZF1 promotes the malignant behaviors of osteosarcoma cell, including invasion, proliferation and migration in MG63 or U2OS cells, MG63 and U2OS cells were transfected with VEZF1-expressing plasmid to elevate VEZF1 protein expression (P<0.01; Fig. 4A). The results of the CCK-8 assay demonstrated that overexpression of VEZF1 significantly enhanced the proliferation of MG63 and U2OS cells (P<0.05; Fig. 4B). Furthermore, overexpression of VEZF1 significantly increased the colony formation ability of MG63 and U2OS cells (P<0.01; Fig. 4C). The results of the wound healing assay demonstrated that overexpression of VEZF1 significantly increased the migratory ability of MG63 and U2OS cells transfected with NC mimic or miR-382-5p mimic. Scale bar, 200 µm. (E) The Transwell assay was performed to assess invasive ability of MG63 and U2OS cells transfected with NC mimic or miR-382-5p mimic. Scale bar, 200 µm. (F) Flow cytometric analysis was performed to detect apoptosis in MG63 and U2OS cells transfected with NC mimic or miR-382-5p mimic. *P<0.01, miR, microRNA; NC, negative control; OD, optical density.

Overexpression of VEZF1 abrogates the effect of miR-382-5p on cell proliferation, invasion, migration and cell apoptosis. To further investigate the association between VEZF1 and miR-382-5p in human osteosarcoma cells, the effects of overexpressing VEZF1 on malignant behaviors of MG63 and U2OS cells were assessed. Western blot analysis demonstrated that overexpression of VEZF1 significantly decreased cell apoptosis compared with the control group (P<0.01; Fig. 4F). Taken together, these results suggest that VEZF1 promotes malignant behaviors of MG63 and U2OS cells.
Figure 3. VEZF1 is a direct target of miR-382-5p in osteosarcoma. (A) Bioinformatics analysis was performed to identify the potential targets of miR-382-5p, among which nine mRNAs were selected for subsequent experimentations. (B) Relative expression of potential targets of miR-382-5p in MG63 cells following transfection with miR-382-5p mimic. (C) The mature miR-382-5p sequence, the putative binding sites of miR-382-5p and the corresponding wild-type and mutant sites of VEZF1 mRNA 3'‑UTR (left) and the luciferase activity of the reporter vectors demonstrated that VEZF1 mRNA expression significantly decreased in MG63 cells following transfection with miR-382-5p mimics (right). (D) RT‑qPCR analysis was performed to detect VEZF1 mRNA expression in osteosarcoma tissues and adjacent normal tissues. *P<0.05, **P<0.01. (E) RT‑qPCR analysis was performed to detect VEZF1 mRNA expression in osteosarcoma cell lines (143B, MG63 and U2OS) and normal cell lines (HFOB and HOBC). **P<0.01 vs. HFOB cells; ##P<0.01 vs. HOBC cells. (F) Pearson's correlation analysis was performed to assess the correlation between miR-382-5p and VEZF1 expression in osteosarcoma. (G) RT‑qPCR analysis was performed to detect VEZF1 mRNA expression in MG63 cells transfected with miR-382-5p mimic or NC mimic. *P<0.05. (H) Western blot analysis was performed to detect VEZF1 protein expression in cells transfected with miR-382-5p mimic or NC mimic. **P<0.01. VEZF1, vascular endothelial zinc finger 1; miR, microRNA; UTR, untranslated; RT‑qPCR, reverse transcription‑quantitative PCR; NC, negative control; WT, wild-type; MUT, mutant.
miR-382-5p targets VEZF1 in osteosarcoma.

Furthermore, simultaneous overexpression of VEZF1 in miR-382-5p-transfected MG63 and U2OS cells significantly increased the number of cell colonies (P<0.01; Fig. 5C), and significantly induced the cell migratory (P<0.01; Fig. 5D) and invasive (P<0.01; Fig. 5E) abilities, while decreasing apoptosis (P<0.01; Fig. 5F). Taken together, these results suggested that simultaneous overexpression of VEZF1 reverses the effects of miR-382-5p overexpression on malignant behaviors of human osteosarcoma cells.

Overexpression of VEZF1 attenuates the inhibitory effect of miR-382-5p in an animal model. To further investigate the tumor-inhibiting function of miR-382-5p via VEZF1 in human osteosarcoma in vivo, empty lentivirus-infected control stable U2OS cells (LV), miR-382-5p-expressing lentivirus-infected stable U2OS cells (LV-miR-382-5p) and VEZF1 overexpression vectors (VEZF1) were inoculated and co-transfected into LV-miR-382-5p cells on the left flank of model mice via subcutaneous injection. Some of the mice were sacrificed at day 18 following tumor inoculation. The
results demonstrated that 21 days after xenografting, the tumor volume significantly decreased in U2OS cells stably transfected with LV-miR-382-5p significantly. In addition, the weight of the mice decreased by 70% compared with the miR-382-5p mimics and VEZF1 co-transfection or miR-NC groups (Fig. 6A and B). RT-qPCR analysis demonstrated a notable increase in miR-382-5p expression following transduction with miR-382-5p lentivirus but had no obvious change after overexpressing VEZF1, whereas VEZF1 mRNA expression significantly decreased following transfection with miR-382-5p mimics but significantly increased after overexpressing VEZF1 (P<0.01; Fig. 6C), which confirms the association between miR-382-5p and VEZF1 in human osteosarcoma.

Immunohistochemical analysis of Ki-67 in tumor tissues indicated pleomorphic poorly differentiated neoplastic cells, confirming cancerous tissue growth (Fig. 6D). Subsequent experiments focused on miR-382-5p-mediated regulation of B-cell lymphoma 2 (Bcl2) and Bax, which are downstream of the vascular endothelial growth factor (VEGF) signaling pathway (25). Western blot analysis demonstrated that overexpression of miR-382-5p promoted Bax expression and suppressed Bcl2 expression, while simultaneously promoting cleaved caspase-3 expression (Pc0.01; Fig. 6E). In addition, overexpression of VEZF1 attenuated the suppression of miR-382-5p in osteosarcoma cells. Taken together, these results suggest that the size and weight of xenograft tumors in nude mice significantly decrease following transfection with miR-372-3p mimics.
Discussion

The first miRNAs, which are small non-coding RNA segments (~22 nucleotides in length) were identified in 1993 (in Caenorhabditis elegans) (26). These miRNAs have diverse roles in cellular development, differentiation, apoptosis and metabolism, including osteoma initiation and normal osteogenesis (9). These molecules have important roles in various
functions, such as those of tumor-suppressor genes or oncogenes, so when they are disordered, they contribute to initiation and progression of cancer (27). Recent studies have reported that aberrant expression of miRNAs has been demonstrated in a range of human diseases, including various malignancies (28-30). miR-382-5p has been repeatedly observed in various human malignancies, including lung, oral, ovarian, breast and prostate cancers (16-19). Given that miR-382-5p has been reported to be frequently and significantly reduced in tumor tissues and tumor-associated cell lines, it was speculated that miR-382-5p functions as a tumor-suppressor gene or oncogene in osteosarcoma. However, the highly complex molecular mechanisms underlying miR-382-5p inhibition of osteosarcoma cells remain unclear.

The results of the present study demonstrated that miR-382-5p expression was markedly downregulated in clinical osteosarcoma tumor tissues and osteosarcoma-associated cell lines compared with normal tissues and cell lines. In addition, miR-382-5p exerted antitumor functions in human osteosarcoma cells, including restraining cell colony formation, proliferation, migration and invasion, as well as enhancing apoptosis in vitro and suppressing tumor growth in vivo. These results are in accordance with previous findings, demonstrating that miR-382-5p expression is notably downregulated in cancer cells (16,31,32). Previous study had demonstrated that miR-382-5p acts as tumor suppressor to inhibit the metastasis and relapse of osteosarcoma by targeting YB-1 (33), while circ_0001658 can significantly promote the proliferation and metastasis of osteosarcoma cells. Taken together, these results suggest that overexpression of miR-382-5p exerts a tumor-suppressive effect in osteosarcoma.

A previous study reported that miRNAs recognize and combine specific target mRNAs depending on complete or partial base-pairing, mostly at the 3'-UTR of the target genes, to regulate gene expression following transcription, which includes both tumor suppressors and oncogenes (41). Bioinformatic analysis was performed in the present study to predict gene targets of miR-382-5p. The results revealed that VEZF1 has a highly conserved miR-382-5p binding sequence in the mRNA 3'-UTR. The results of the dual-luciferase reporter assay confirmed the direct association between miR-382-5p and VEZF1 in human osteosarcoma cells.

VEZF1 is a Krüppel-like zinc finger protein that plays a vital role in vascular development, and is a member of the VEGF signaling pathway (42). Abnormal expression of VEZF1 contributes to the pathogenesis of different disorders, including cancer and cardiovascular diseases (43). Recent study has reported that a small molecule inhibitor of VEZF1, Vec6, prevents wound healing and angiogenesis (44). Increasing evidences suggest that VEZF1 plays a major role in the development and progression of malignancy (43,45,46). The results of the present study demonstrated that overexpression of VEZF1 in vitro and in a nude mouse model increased proliferation, invasion and migration of osteosarcoma cells, and decreased apoptosis. Furthermore, overexpression of VEZF1 notably attenuated the tumor-suppressive effects of miR-382-5p mimics in human osteosarcoma cells and the animal model, which suggests that VEZF1 may act as an oncogene in human osteosarcoma cells. However, further studies are required to screen potential targets of miR-382-5p other than VEZF1 to determine the tumor-suppressive functions of miR-382-5p in human osteosarcoma.

In conclusion, the results of the present study demonstrated that miR-382-5p acts as a tumor-suppressor gene, whose ectopic expression inhibits the malignant biological behaviors of human osteosarcoma cells in vitro and in a nude mice model. VEZF1 was identified as a direct target gene of miR-382-5p, significantly downregulated in clinical osteosarcoma tissues and osteosarcoma-associated cell lines. Despite the limitation in serum samples, the results of the present study and previous findings suggest that miR-382-5p may serve as a novel biomarker for the prognosis of patients with osteosarcoma. Considering the prognostic significance, serum miR-382-5p expression and its prognostic value in osteosarcoma will be investigated in prospective studies.
and its overexpression remarkably attenuated the inhibitory effect of miR-382-5p mimics in human osteosarcoma cells. The miR-382-5p/VEZF1 axis provides novel insights into the pathogenesis of osteosarcoma, and miR-382-5p expression may have effects on the malignancy of osteosarcoma cells. Thus, miR-382-5p expression may potentially act as a prognostic biomarker or a therapeutic target in osteosarcoma.

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Availability of data and materials

The datasets used and/or analyzed in this study are available from the corresponding author upon reasonable request.

Authors' contributions

KL and GS conceived and designed the experiments. HW, YXL, MLZ and JB performed the experiments. WH, MY, RP, NXH and GF analyzed and interpreted the data. WH and HW drafted the initial manuscript. KL and GS revised the manuscript for important intellectual content. KL and GS confirm 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 the Institutional Center Ethics Review Committee of Nanchong Central Hospital (Nanchong, China; approval no. 2020-002) and written informed consent was provided by all patients or their guardians prior to the study start. All animal experiments were performed according to the Guide for the Care and Use of Laboratory Animals (National Academies Press, 2011; Nanchong, China; approval no. NSMC-2020-42) and all experiments were performed according to the AVMA guidelines.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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