MicroRNA-376a inhibits cell proliferation and invasion in osteosarcoma via directly targeting SATB1

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Abstract. Aberrantly expressed microRNAs (miRs) are implicated in the regulation of osteosarcoma (OS) onset and development. Therefore, key miRs in OS must be identified to develop promising and effective therapeutic targets for patients with OS. In the present study, reverse transcription-quantitative polymerase chain reaction analysis revealed that miR-376a-3p expression was downregulated in OS tissues and cell lines. Additionally, decreased miR-376a expression was associated with tumor size and lymph node infiltration. Restoration of miR-376a expression reduced cell proliferation and invasion of OS. Furthermore, special AT-rich sequence-binding protein 1 (SATB1) was identified as a direct target gene of miR-376a in OS cells. Furthermore, SATB1 was overexpressed in OS tissues and SATB1 overexpression was inversely correlated with the expression level of miR-376a. In addition, ectopic SATB1 expression counteracted the inhibitory effects of miR-376a overexpression on the proliferation and invasion of OS cells. All these results identified that reduced miR-376a expression may be implicated in the mechanism underlying OS progression, suggesting that the miR-376a/SATB1 axis may be a promising novel target for potential therapeutic methods for the effective treatment of patients with OS.

Introduction

Osteosarcoma (OS) is a common primary malignant tumor of the bone and primarily occurs in adolescents and young adults (1). It accounts for ~2.4% of all tumors in children and adolescents (2). The majority of cases of OS affect the long bones, particularly the metaphysis, proximal tibia, proximal humerus and distal femur (3). Although significant advancements have been achieved in surgical resection combined with radiotherapy and chemotherapy, the prognosis of patients

with OS has only been slightly improved, with the median survival time reaching only 23 months (4,5). The poor treatment outcome of patients with OS is due to local recurrence and metastasis, especially pulmonary metastasis (6). Although numerous molecular targets have been identified to be implicated in OS formation and progression, their detailed mechanisms remain unknown (7). Therefore, the mechanisms underlying OS occurrence and development must be comprehensively understood to identify effective therapeutic targets for treating OS.

MicroRNAs (miRs) are highly conserved, non-coding and small RNAs that comprise 19-23 nucleotides (8). miRs are considered critical gene regulators that directly bind to the 3'-untranslated regions (3'-UTRs) by base pairing and therefore induce mRNA degradation and/or translational suppression (9). miRs regulate ~60% of all human protein coding genes, indicating that miRs serve important roles in physiological and pathological behavior, including cell proliferation, cycle, apoptosis, metastasis, embryogenesis and differentiation (10). The dysregulation of miRs has been recently reported in different types of malignancies, including OS (11), prostate cancer (12), breast cancer (13) and colorectal cancer (14). Abnormally expressed miRs may serve tumor suppressive roles or oncogenic roles in tumorigenesis and tumor development, which depend on the biological functions of their target genes (15). Therefore, the expression and roles of miRs in OS must be further investigated to identify potential therapeutic targets for the therapy of patients with OS.

miR-376a-3p (miR-376a) is dysregulated in several human cancer types and the dysregulation of miR-376a serves an important role in various cell processes, including cell proliferation, migration, invasion, metastasis and apoptosis (16-19). However, to the best of our knowledge, the expression pattern, clinical significance, specific role and detailed mechanisms of miR-376a in OS remains unclear. Therefore, the present study aimed to detect the miR-376a expression in OS and its association with clinicopathological characteristics and to determine the biological roles and regulatory mechanism of miR-376a in OS cells.

Materials and methods

Tissue samples. A total of 49 primary OS tissues and corresponding adjacent normal bone tissues were obtained from patients who underwent surgical resection in the China-Japan Union Hospital of Jilin University (Changchun, China), between

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October 2013 and March 2017. The patient characteristics are presented in Table I. No chemotherapy or radiotherapy was administered prior to surgery. All tissues were snap-frozen in liquid nitrogen (-196°C) at the time of surgery and were stored in liquid nitrogen until further use. The present study was approved by the Ethics Committee of the China-Japan Union Hospital of Jilin University. In addition, written informed consent was provided by all participants.

Cell culture and transfection. A total of four human OS cell lines (HOS, SAOS-2, MG-63 and U2OS), and a normal human osteoblast hFOB1.19 cell line were acquired from the Shanghai Cell Bank of the Chinese Academy of Sciences (Shanghai, China). All cell lines were cultured at 37°C in a humidified atmosphere containing 5% CO₂ and were maintained in Dulbecco's modified Eagle's medium (DMEM) supplemented with 10% fetal bovine serum (FBS; both from HyClone; GE Healthcare Life Sciences, Logan, UT, USA), 100 U/ml penicillin and 100 U/ml streptomycin (Sigma-Aldrich; Merck KGaA, Darmstadt, Germany).

miR-376a mimic and miR mimic negative controls (miR-NC) were synthesized and purchased from the Shanghai GenePharma Co., Ltd. (Shanghai, China). The miR-376a mimic sequence was 5'-AUCAUAGAGGAAAAUCCACGU-3' and the miR-NC sequence was 5'-UUCUCCGAACGUGUCACGUTT-3'. SATB1 overexpression vector pcDNA3.1-SATB1 and control empty vector pcDNA3.1 were ordered from GeneCopoeia Inc., (Rockville, MD, USA). For transfection, MG-63 and U2OS cells were plated into 6-well plates with a density of 5x10⁵/well and grown in culture medium without antibiotics. MG-63 and U2OS cells were transfected with miR-376a mimic (100 pmol), miR-NC (100 pmol), pcDNA3.1-SATB1 (4 µg) or pcDNA3.1 (4 µg) by using Lipofectamine® 2000 (Invitrogen; Thermo Fisher Scientific, Inc., Waltham, MA, USA), following the manufacturer's protocol. After transfection 48 h, reverse transcription-quantitative polymerase chain reaction (RT-qPCR) was performed to detect miR-376a expression. MTT and invasion assays were carried out at 24 h and 48 h posttransfection. Western blot analysis was performed to detect SATB1 protein expression at 72 h following transfection.

RT-qPCR analysis. The total RNA was isolated from the cells or tissues using the TRIzol® reagent (Invitrogen; Thermo Fisher Scientific, Inc.). miR complementary DNA (cDNA) was produced using a TaqMan MicroRNA Assay kit (Applied Biosystems; Thermo Fisher Scientific, Inc.). The temperature protocol for reverse transcription was as follows: 16°C for 30 min, 42°C for 30 min and 85°C for 5 min. Quantification of miR-376a was performed using a TaqMan MicroRNA assay (Applied Biosystems; Thermo Fisher Scientific, Inc.). The cycling conditions were as follows: 50°C for 2 min, 95°C for 10 min; 40 cycles of denaturation at 95°C for 15 sec; and annealing/extension at 60°C for 60 sec. To quantify the SATB1 mRNA, the total RNA was reverse transcribed into cDNA using a PrimeScripts RT reagent kit and qPCR was conducted using SYBR-Green Premix Ex Taq II (both from Takara Biotechnology Co., Ltd., Dalian, China). The temperature protocol for reverse transcription was as follows: 37°C for 15 min and 85°C for 5 sec. The amplification was performed

Table I. Association between miR-376a and clinicopathological features of osteosarcoma patients.

Clinicopathological features	miR-376a expression		P-value
	Low (n=25)	High (n=24)	
Age (years)			0.567
<20	16	13	
≥20	9	11	
Gender			0.377
Male	14	17	
Female	11	7	
Tumor size (cm)			0.022
<8	7	15	
≥8	18	9	
Location of the primary tumor			0.289
Tibia/femur	22	18	
Elsewhere	3	6	
Lymph node infiltration			0.001
No	9	20	
Yes	16	4	

with cycling conditions as follows: 5 min at 95°C, followed by 40 cycles of 95°C for 30 sec and 65°C for 45 sec. The primers were designed as follows: miR-376a, 5'-GTGCAGGGTCCGAGGT-3' (forward) and 5'-ATCATAGAGGAAAATCCACG-3' (reverse); U6, 5'-CTCGCTTCGGCAGCACA-3' (forward) and 5'-AACGCTTCACGAATTTGCGT-3' (reverse); SATB1, 5'-AGAGGAAGGCTTGGGAGTA-3' (forward) and 5'-GGG CAGCAGAGCTATGTG-3' (reverse); and GAPDH, 5'-CGG AGTCAACGGATTTGGTTCGTAT-3' (forward) and 5'-AGC CTTCTCCATGGTGGTGAAGAC-3' (reverse). U6 snRNA and GAPDH were used as the internal control for miR-376a and SATB1 mRNA, respectively. Data were analyzed by 2^{-ΔΔCq} method (20).

MTT assay. Following transfection for 24 h, the cells were re-suspended and inoculated into 96-well plates at a density of 3x10³ cells per well. The cells were incubated at 37°C under a humidified atmosphere containing 5% CO₂ for 0-72 h. An MTT assay was performed every 24 h to detect cell proliferation. A total of 10 µl MTT solution (Sigma-Aldrich; Merck KGaA) was added to each well and the cells were incubated for 4 h at 37°C. After the supernatant was discarded, 100 µl dimethyl sulfoxide (Sigma-Aldrich; Merck KGaA), which was used to resolve the crystalline formazan, was added to each well. The absorbance at a wavelength of 490 nm was monitored with a microplate reader (Bio-Rad Laboratories, Inc., Hercules, CA, USA).

Invasion assay. The transfected cells were digested at 48 h post-transfection, washed with PBS and suspended in FBS-free DMEM medium. A total of 1x10⁵ cells in FBS-free DMEM were seeded into the upper chamber of Transwell

Figure 1. miR-376a expression is downregulated in OS. (A) Reverse transcription-quantitative polymerase chain analysis was utilized to detect miR-376a expression in 49 pairs of OS tissues and adjacent normal bone tissues. * $P < 0.05$ vs. the normal bone tissues. (B) The expression level of miR-376a was determined in four human OS cell lines, namely, HOS, SAOS-2, MG-63 and U2OS, as well as the normal human osteoblast hFOB1.19 cell line. * $P < 0.05$ vs. hFOB1.19. miR, microRNA; OS, osteosarcoma.

plates (Corning Inc., Corning, NY, USA), which were pre-coated with Matrigel (BD Biosciences, Franklin Lakes, NJ, USA). Subsequently, 500 μ l DMEM supplemented with 10% FBS was added to the lower chamber of the Transwell plates. Following 24 h of culture at 37°C, the cells remaining on the top of the chamber were removed using cotton swabs. The invasive cells were fixed with 4% paraformaldehyde at room temperature for 20 min and stained with 0.05% crystal violet at room temperature for 20 min. A total of five fields per chamber were randomly selected and the number of invasive cells was counted under an inverted light microscope (x200; Olympus Corporation, Tokyo, Japan).

Target genes of miR-376a. TargetScan Human 7.0 (<http://www.targetscan.org/>) and miRBase (<http://www.mirbase.org/>) were employed to predict the putative targets of miR-376a.

Luciferase reporter assay. A wild-type (Wt) fragment of the 3'-UTR of SATB1 containing the predicted miR-376a binding sequences and its mutated (Mut) sequence was chemically produced by Shanghai GenePharma Co., Ltd., and cloned into pMIR-GLO™ Luciferase vector (Promega Corporation, Madison, WI, USA). The Wt and Mut luciferase plasmids were defined as pMIR-SATB1-3'-UTR Wt and pMIR-SATB1-3'-UTR Mut, respectively. Cells were seeded into 24-well plates with a density of 1.5×10^5 /well and cotransfected with miR-376a mimic or miR-NC, and pMIR-SATB1-3'-UTR Wt or pMIR-SATB1-3'-UTR Mut, using Lipofectamine 2000 according to the manufacturer's protocol. Following transfection for 2 days, the dual-Luciferase Reporter Assay System (Promega Corporation) was applied to measure the luciferase activities, according to the manufacturer's protocol. Firefly luciferase activity was normalized to *Renilla* luciferase activity.

Western blot analysis. Tissue samples or cells were solubilized in cold radioimmunoprecipitation assay lysis buffer (Beyotime Institute of Biotechnology, Shanghai, China). The total protein concentration was determined using a Bicinchoninic Acid Protein Assay kit (Pierce Biotechnology, Inc., Rockford, IL, USA). Equal amounts of proteins (20 μ g) were separated using 10% SDS-PAGE, electrotransferred to polyvinylidene

fluoride membranes (EMD Millipore, Billerica, MA, USA) and blocked at room temperature for 2 h in Tris-buffered saline with 0.5% Tween (TBST) buffer containing 5% dried skimmed milk. Following being washed three times with TBST, the membranes were incubated with rabbit anti-human monoclonal SATB1 antibody (cat. no. ab109122; 1:1,000; Abcam, Cambridge, MA, USA) or rabbit anti-human monoclonal GAPDH antibody (cat. no. ab181603; 1:1,000; Abcam) at 4°C overnight. Afterwards, the membranes were washed three times with TBST and further incubated with goat anti-rabbit horseradish peroxidase-conjugated secondary antibody (cat. no. ab6721; 1:5,000; Abcam) at room temperature for 1 h. Finally, an ECL Protein Detection kit (Pierce Biotechnology, Inc., Rockford, IL, USA) was used to visualize the protein bands. GAPDH was used as the loading control. Protein expression was quantified using Quantity One software version 4.62 (Bio-Rad Laboratories, Inc.).

Statistical analysis. Data were presented as the mean $\pm$ standard deviation of at least 3 independent experiments and analyzed with SPSS software, version 21.0 (IBM SPSS, Armonk, NY, USA). The association between miR-376a expression and clinicopathological characteristics of OS was evaluated by a χ^2 test. The two-tailed Student's t-test and one-way analysis of variance combined with Student-Newman-Keuls were used to compare differences between groups. Spearman's correlation analysis was carried out to reveal the correlation between expression levels of miR-376a and SATB1 mRNA in OS tissues. $P < 0.05$ was considered to indicate a statistically significant difference.

Results

miR-376a expression is decreased in OS tissues and cell lines. To understand the expression pattern of miR-376a in OS, miR-376a expression was initially measured in 49 pairs of OS tissues and adjacent normal bone tissues. RT-qPCR analysis revealed that miR-376a was significantly downregulated in OS tissues compared with the adjacent normal bone tissues ($P < 0.05$; Fig. 1A). Subsequently, the clinical significance of miR-376a expression in OS was investigated.

Figure 2. miR-376a overexpression inhibits proliferation and invasion of MG-63 and U2OS cells. (A) MG-63 and U2OS cells were transfected with miR-376a mimic or miR-NC. The transfection efficiency was quantified using reverse transcription-quantitative polymerase chain reaction. * $P < 0.05$ vs. miR-NC. (B) MTT assay was used to detect the proliferation of MG-63 and U2OS cells transfected with miR-376a mimic or miR-NC. * $P < 0.05$ vs. miR-NC. (C) The effect of miR-376a overexpression on the invasion abilities of MG-63 and U2OS cells was evaluated using an invasion assay. * $P < 0.05$ vs. miR-NC. miR, microRNA; miR-NC, microRNA negative control.

Figure 3. SATB1 is a direct target gene of miR-376a in OS cells. (A) miR-376a putative Wt binding sequences and corresponding Mut sequences in the 3'-UTR of SATB1. (B) Luciferase activity was quantified in MG-63 and U2OS cells co-transfected with miR-376a mimic or miR-NC, and pMIR-SATB1-3'-UTR Wt or pMIR-SATB1-3'-UTR Mut. * $P < 0.05$ vs. miR-NC. The expression levels of SATB1 were detected in MG-63 and U2OS cells transfected with miR-376a mimic or miR-NC by (C) reverse transcription-quantitative polymerase chain reaction and (D) western blot analysis. * $P < 0.05$ vs. miR-NC. miR, microRNA; miR-NC, microRNA negative control; Mut, mutant; Wt, wild-type; OS, osteosarcoma cells; UTR, untranslated region; SATB1, special AT-rich sequence-binding protein 1.

Figure 4. miR-376a expression is negatively correlated with SATB1 mRNA level in OS tissues. (A) The mRNA and (B) protein levels of SATB1 in OS tissues and adjacent normal bone tissues were quantified using reverse transcription-quantitative polymerase chain reaction and western blot analysis, respectively (C) Densitometry data from the western blotting. * $P < 0.05$ vs. the normal bone tissues. (D) Spearman's correlation analysis was performed to investigate the association between miR-376a and SATB1 mRNA expression in OS tissues. $r = -0.5288$, $P < 0.0001$. OS, osteosarcoma; miR, microRNA; SATB1, special AT-rich sequence-binding protein 1.

The median value of miR-376a expression was defined as the cutoff point and this cutoff point was used to divide all OS patients into a low miR-376a expression group ($n = 25$) or a high miR-376a expression group ($n = 24$). A low miR-376a level was significantly correlated with tumor size ($P = 0.022$) and lymph node infiltration ($P = 0.001$) in OS but was not associated with age, gender, and location of the primary tumor (all $P > 0.05$; Table I). In addition, the expression of miR-376a in four human OS cell lines, namely, HOS, SAOS-2, MG-63 and U2OS, as well as a normal human osteoblast hFOB1.19 cell line, was determined. As indicated in Fig. 1B, the expression level of miR-376a in all OS cell lines was significantly decreased compared with in hFOB1.19 ($P < 0.05$). These results suggested that reduced expression of miR-376a may be associated with OS progression.

miR-376a upregulation prohibits the proliferation and invasion of OS cells. To determine whether miR-376a can affect the malignant phenotypes of OS cells, MG-63 and U2OS cells, which exhibited relatively lower miR-376a expression among the four OS cell lines, were transfected with an miR-376a mimic or miR-NC. Transfection with the miR-376a mimic significantly increased the miR-376a levels in MG-63 and U2OS cells when compared with the cells transfected with miR-NC ($P < 0.05$; Fig. 2A). MTT and invasion assays were used to examine the effects of miR-376a overexpression on OS cell proliferation and invasion, respectively. The results revealed that ectopic expression of miR-376a significantly inhibited the proliferation ($P < 0.05$; Fig. 2B) and invasion ($P < 0.05$; Fig. 2C) of MG-63 and U2OS cells compared with

the miR-NC groups. These results suggested that miR-376a may serve a tumor suppressive role in OS progression.

miR-376a directly targets SATB1 and inhibits its expression in OS cells. To illustrate the molecular mechanism by which miR-376a inhibits OS cell proliferation and invasion, bioinformatics analysis was conducted to predict the potential targets of miR-376a. SATB1 was predicted as a primary target of miR-376a (Fig. 3A) and was chosen for additional confirmation analysis due to its regulatory roles in OS pathogenesis and development (21,22). To confirm this prediction, luciferase reporter assays were conducted in MG-63 and U2OS cells, which were transfected with pMIR-SATB1-3'-UTR Wt or pMIR-SATB1-3'-UTR Mut, together with the miR-376a mimic or miR-NC. Upregulation of miR-376a reduced the luciferase activities of the plasmid carrying the Wt 3'-UTR of SATB1 in MG-63 and U2OS cells ($P < 0.05$) but had no effect on that of the Mut 3'-UTR of SATB1 (Fig. 3B). Furthermore, the effects of miR-376a overexpression on endogenous SATB1 expression were determined using RT-qPCR and western blot analysis, respectively. As expected, SATB1 mRNA ($P < 0.05$; Fig. 3C) and protein ($P < 0.05$; Fig. 3D) levels were significantly reduced in MG-63 and U2OS cells transfected with miR-376a mimic. Therefore, SATB1 is a direct target gene of miR-376a in OS.

SATB1 expression is upregulated in OS and inversely correlated with miR-376a expression. SATB1 was validated as a direct target gene of miR-376a in OS. Therefore, the SATB1 expression was detected in 49 pairs of OS tissues and adjacent normal bone tissues using RT-qPCR. The expression level of

Figure 5. Recovered SATB1 expression rescues OS cells from the inhibitory effects of miR-376a overexpression. (A) MG-63 and U2OS cells were transfected with miR-NC, miR-376a mimic+pcDNA3.1 or miR-376a mimic+pcDNA3.1-SATB1. Following transfection for 72 h, SATB1 protein expression was detected using western blot analysis. * $P < 0.05$ vs. miR-NC. # $P < 0.05$ vs. miR-376a mimic+pcDNA3.1-SATB1. The (B) proliferation and (C) invasion of indicated cells was examined using MTT and invasion assays, respectively. * $P < 0.05$ vs. miR-NC. # $P < 0.05$ vs. miR-376a mimic+pcDNA3.1-SATB1. miR, microRNA; miR-NC, microRNA negative control; SATB1, special AT-rich sequence-binding protein 1; OS, osteosarcoma.

SATB1 was significantly upregulated in OS tissues compared with the adjacent normal bone tissues ($P < 0.05$; Fig. 4A). In addition, the SATB1 protein expression was also determined in several pairs of OS tissues and adjacent normal bone tissues chosen at random using western blot analysis. As demonstrated in Fig. 4B, the SATB1 protein expression in OS tissues was significantly increased compared with the adjacent normal bone tissues in the 4 cases tested ($P < 0.05$; Fig. 4B and C). Furthermore, miR-376a expression was inversely correlated with SATB1 mRNA levels in OS tissues ($P < 0.0001$; $r = -0.5288$; Fig. 4D). These results further suggested that SATB1 is a direct target gene of miR-376a in OS.

SATB1 overexpression attenuates the suppressive effects of miR-376a on OS cell proliferation and invasion. Rescue experiments were performed to assess whether SATB1 mediated the inhibitory roles of miR-376a in OS. miR-376a-overexpressing MG-63 and U2OS cells were co-transfected with a SATB1 overexpression vector pcDNA3.1-SATB1 or control empty vector pcDNA3.1. The decreased expression level of SATB1 induced by miR-376a overexpression was recovered in MG-63 and U2OS cells co-transfected with pcDNA3.1-SATB1 ($P < 0.05$; Fig. 5A). MTT and invasion assays revealed that restored SATB1 expression eliminated the inhibitory effects of miR-376a overexpression on the proliferation ($P < 0.05$; Fig. 5B) and invasion ($P < 0.05$; Fig. 5C) of MG-63 and U2OS cells. These results suggested that miR-376a serves a tumor suppressive role in OS partly by inhibiting SATB1 expression.

Discussion

Aberrantly expressed miRs are implicated in the regulation of OS onset and development (23-25). Therefore, key miRs in OS must be identified to develop promising and effective

therapeutic targets for patients with OS. In the present study, the expression level and clinical significance of miR-376a in OS was measured and the roles, and associated regulatory mechanism of miR-376a were determined in OS. The results demonstrated that miR-376a expression was downregulated in OS tissues and cell lines. Decreased expression of miR-376a was associated with tumor size and lymph node infiltration of OS patients. In addition, miR-376a overexpression restricted cell proliferation and invasion in OS. Furthermore, SATB1 was confirmed as a direct target gene of miR-376a in OS. Additionally, SATB1 expression was significantly overexpressed in OS tissues and this overexpression was inversely correlated with miR-376a levels in OS tissues. Reintroduction of SATB1 expression rescued OS cells from the tumor suppressive roles of miR-376a. These results revealed that miR-376a may serve as a tumor suppressor in OS by directly targeting SATB1.

Significant alterations in miR-376a expression have been observed in multiple human cancer types. For instance, miR-376a is under-expressed in glioma tissues, cell lines and serum (16,26). Low miR-376a expression is correlated with the World Health Organization grade and KPS score of glioma patients. In addition, miR-376a is validated as an independent factor that predicts the poor prognosis of glioma patients (26). In hepatocellular carcinoma, miR-376a is downregulated in tumor tissues and this downregulation is associated with α -fetoprotein levels (17). In colorectal cancer, the expression level of miR-376a in tumor tissues is decreased compared with the adjacent normal mucosa. Low miR-376a expression is associated with lymph node metastasis of colorectal cancer patients. Colorectal cancer patients with low miR-376a expression present lower survival rates compared with patients with high miR-376a levels (27). Low expression of miR-376a has also been observed in prostate cancer (18) and non-small-cell

lung cancer (19). Conversely, miR-376a is overexpressed in ovarian cancer. High miR-376a expression is significantly associated with the clinical stage and the International Federation of Gynaecological Oncologists stage of ovarian cancer (28,29). These conflicting results indicate that the expression patterns of miR-376a exhibit tissue specificity and suggest that miR-376a may be developed as a meaningful prognostic biomarker for predicting the prognosis of patients with these human cancer types.

Differently expressed miRs have been associated with the carcinogenesis and cancer progression of several human malignancies. For instance, miR-376a suppresses the proliferation and invasion of glioblastoma multiform cells (16). Zheng *et al* (17) reported that miR-376a overexpression inhibits cell proliferation and promotes apoptosis in hepatocellular carcinoma. Formosa *et al* (18) demonstrated that the resumption of miR-376a expression attenuates the growth of prostate cancer cells while inducing apoptosis. Wang *et al* (19) indicated that miR-376a re-expression restricts cell proliferation and invasion while promoting apoptosis in non-small-cell lung cancer. Conversely, miR-376a serves oncogenic roles in ovarian cancer by promoting cell proliferation and motility, inhibiting cell apoptosis *in vitro* and increasing tumor growth *in vivo* (29). These studies suggested that miR-376a may be a potential therapeutic target for patients with these human malignancies.

Previous studies have identified several direct targets of miR-376a, including SP1 (16) in glioblastoma multiform, p85 α (17) and H3K18 (30) in hepatocellular carcinoma, frizzled-4 in prostate cancer (18), c-Myc (19) in non-small-cell lung cancer, Krueppel-like factor 15 (29) and Caspase-8 (29) in ovarian cancer. SATB1, which is a cell type-specific nuclear matrix attachment region binding protein, was a direct target gene of miR-376a in OS. SATB1 is reportedly upregulated in several types of cancer, including hepatocellular carcinoma (31), ovarian cancer (32), gastric cancer (33), prostate cancer (34) and colorectal cancer (35). The expression of SATB1 was also upregulated in OS tissues and cell lines. The overexpression of SATB1 contributes to OS progression by promoting cell proliferation, migration and invasion, inhibiting apoptosis and reducing the chemosensitivity of cells to arsenic trioxide (21,22). These results suggest that knockdown of SATB1 may be a valuable therapeutic strategy for the treatment of patients with OS.

In conclusion, miR-376a expression was downregulated in OS tissues and cell lines, and the low miR-376a expression in OS was significantly associated with the tumor size and lymph node infiltration. In addition, forced expression of miR-376a repressed OS cell proliferation and invasion by directly targeting SATB1. Therefore, the results of the present study suggest that miR-376a may be a promising therapeutic target for treating patients with OS.

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Availability of data and materials

The datasets used and/or analyzed during the present study are available from the corresponding author on reasonable request.

Authors' contributions

HJ designed the present study. GZ and LM performed the functional experiments. LM analyzed the data of this study. All authors have read and approved the final draft.

Ethics approval and consent to participate

The present study was approved by the Research Ethics Committee of China-Japan Union Hospital of Jilin University (Jilin, China) and was performed in accordance with the Declaration of Helsinki and the guidelines of the Ethics Committee of China-Japan Union Hospital of Jilin University. Written informed consent was obtained from all patients for the use of their clinical tissues.

Patient consent for publication

Written informed consent was obtained from all patients for the use of their clinical tissues.

Competing interests

The authors declare that they have no competing interests.

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