miR-33a is up-regulated in chemoresistant osteosarcoma and promotes osteosarcoma cell resistance to cisplatin by down-regulating TWIST

Yong Zhou, Zufa Huang*, Song Wu, Xiaofang Zang, Min Liu and Jian Shi

Abstract

Background: miRNAs are involved in osteosarcoma (OS) chemoresistance, and TWIST reportedly enhances cisplatin-induced OS cell apoptosis by inhibiting multiple signaling pathways. In this study, we profiled miRNAs differentially expressed in chemoresistant OS, with a focus to identify miRNAs that regulate TWIST expression and OS chemoresistance.

Methods: OS patients who showed <90% tumor necrosis after neochemotherapy were defined as poor responders (chemoresistant), and those who showed ≥90% tumor necrosis were defined as good responders (control). miRNA microarray analysis was carried out with a discovery cohort (n = 12) of age-, sex- and tumor stage-matched chemoresistant and control OS patients.

Results: Among the up-regulated miRNAs in chemoresistant OS samples, miR-33a was verified to down-regulate TWIST expression, which was supported by an inverse miRNA-33a/TWIST expression trend in the validation cohort (n = 70), target-sequence-specific inhibition of TWIST-3′ untranslated region-luciferase reporter activity by miR-33a, and alteration of TWIST expression by overexpression or inhibition of miR-33a in human OS cell lines. In Saos-2 cells treated with cisplatin, inhibition of miR-33a by antagomir-33a markedly increased cell apoptosis, which was enhanced by overexpression of TWIST. The apoptosis-inducing effect of TWIST overexpression was reversed by overexpression of miR-33a. In MG-63 cells, overexpression of miR-33a significantly decreased cisplatin-induced cell apoptosis, which was enhanced by knockdown of TWIST. Antagomir-33a significantly increased cisplatin-induced cell apoptosis, which was reversed by knockdown of TWIST.

Conclusions: We have demonstrated in this study that miR-33a is up-regulated in chemoresistant OS and that the miR-33a level is negatively correlated with the TWIST protein level in OS. Our in vitro data indicate that miR-33a promotes OS cell resistance to cisplatin by down-regulating TWIST; on the other hand, inhibition of miR-33a by antagomir-33a enhances cisplatin-induced apoptosis in OS cells by up-regulating TWIST expression. The findings suggest that inhibition of miR-33a/TWIST signaling could be a potential new strategy to enhance neoadjuvant chemotherapy for OS.

Keywords: microRNA, miR-33a, TWIST, Osteosarcoma, Chemoresistance, Apoptosis

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Background

Osteosarcoma (OS) is the most frequent malignant bone tumor in children and adolescents, comprising 2.4% of all malignancies in pediatric patients [1]. The 5-year survival rate of OS patients has significantly improved over the past decades to approximately 60-70% since the introduction of combinatorial chemotherapy [2]. However, a significant proportion of OS patients still respond poorly to chemotherapy, and they have a risk of local relapse or distant metastasis even after curative resection of the primary tumor and intensive chemotherapy. Standard chemotherapy of OS is based on a combination of different drugs: neoadjuvant therapy with methotrexate, cisplatin, and doxorubicin followed by surgery and post-operative chemotherapy (methotrexate, cisplatin, doxorubicin, cyclophosphamide, and vincristine). Despite this, approximately 30% of patients relapse or develop metastasis [3]. The lack of responsiveness to chemotherapy due to intrinsic or acquired chemoresistance is the major reason for poor survival and disease relapse of OS patients [4]. Recently, novel molecular targeted drugs have emerged, but they have not been well established for the treatment of OS [5]. In addition, the molecular mechanisms underlying OS chemoresistance remain largely obscure. Hence, identification of factors that contribute to OS chemoresistance and elucidation of the underlying mechanisms will be pivotal in the development of new therapeutic strategies.

TWIST, also known as TWIST1, belongs to the basic helix-loop-helix (bHLH) transcription factor family. During embryonic development, TWIST plays an essential role in specification of the mesoderm and differentiation of the mesoderm-derived tissues [6]. Twist haploinsufficiency was shown to upset bone tissue in both mice and humans [7,8]. In homogeneous cohort of OS patients, the TWIST gene was frequently deleted in the tumors at diagnosis, and its haploinsufficiency was significantly correlated with a poorer patient outcome [6,9]. It has been reported that TWIST decreases OS cell survival against cisplatin by inhibiting β-catenin signaling and endothelin-1/endothelin A receptor signaling pathways [10,11], suggesting that TWIST is an important negative regulator in the development of OS chemoresistance.

MicroRNAs (miRNAs) are noncoding small RNAs, usually 18–25 nucleotides in length, which repress translation and cleave mRNA by base-pairing to the 3′-untranslated region (UTR) of the target genes [12]. Knowledge of individual miRNAs effecting developmental biology, cellular differentiation programs, and oncogenesis continues to grow [13]. Differences in the miRNA expression profiles detected between cancer cells and their normal counterparts have revealed that miRNAs are involved in the pathogenesis of cancer [14]. In addition, miRNAs may play multiple roles as tumor suppressors, oncogenes, or both in some cases [15]. The biological properties of miRNAs may make them useful as diagnostic and prognostic tools as well as therapeutic targets in various cancers, including OS. A number of miRNAs reportedly are involved in OS tumorigenesis and chemoresistance [13].

In the present study, we screened for miRNAs regulating TWIST expression in human OS and explored their functional interaction in modulating human OS chemoresistance.

Methods

Patients

From November 2010 to May 2013, we enrolled two cohorts of OS patients. The discovery cohort consists of six Han Chinese OS patients who showed <90% tumor necrosis (mean 70.8% ± 9.2%) after chemotherapy and were defined as poor responders at the third Xiangya Hospital of Central South University [16]. Another six age-, sex-, and tumor stage-matched Han Chinese OS patients, who showed ≥90% tumor necrosis (mean 94.1% ± 2.8%) as good responders [16], were enrolled as controls. In the validation cohort, 35 Han Chinese poor responders and 35 Han Chinese good responders were enrolled. All patients had OS in the long tubular bones and were treated preoperatively with neoadjuvant chemotherapy as follows: intravenous (i.v.) doxorubicin (3 courses at 25–30 mg/m²/day for 3 days), i.v. methotrexate (4 courses of up to 14 g/m²/day for 1 day) and intraarterial cisplatin (3 courses at 35 mg/m²/day for 3 days). All OS diagnoses were based on biopsy and the response to treatment was determined histologically as the percentage of necrosis after neoadjuvant chemotherapy. Patients with any other malignancies or a family history of OS or any other cancers were excluded. Baseline characteristics of all 82 patients are summarized in Table 1. This study was approved by the Ethics Committee of the Third Xiangya Hospital, Central South University. Written informed consent was obtained from the parent or guardian of minor participants before the start of the study.

Cells lines, reagents and plasmid constructs

Saos-2 and MG-63 human OS cell lines were purchased from the American Type Culture Collection (Manassas, VA, USA). Human Twist cDNA was subcloned into the pcDNA 3.1 expression vector [17]. Twist (sc-38604-V) short hairpin RNA (shRNA) lentiviral particles, control shRNA lentiviral particles-A (sc-108080), and anti-TWIST (sc-81417) antibody were purchased from Santa Cruz Biotechnology (Santa Cruz, CA, USA). The Dead-End™ Fluorometric TUNEL System was purchased from Promega (Madison, WI, USA). Superfect™ transfection reagent was purchased from Qiagen (Valencia, CA, USA). Dual-luciferase reporter assay system was
purchased from Promega (Madison, WI, USA). Puro-
mycin, cisplatin, and all chemicals of reagent grade were
purchased from Sigma (St. Louis, MO, USA). The 3′-
UTR of TWIST was amplified from genomic DNA using
the following primers: 5′-GCGCCTCGAGCAGGCG-
GAGGCCCTCCCCCTCA-3′ (forward) and 5′-GCG
CGCGCGCCGAGAAAAATATACAAAGATATT-3′
(reverse). The TWIST-3′UTR-luciferase reporter was
generated by inserting the TWIST 3′-UTR between
XhoI and NotI restriction sites (underlined in the
above primers) of the psiCheck2 vector (Promega)
downstream of the renilla luciferase gene. PsiCheck2
vector (Promega) was used as a control vector. TWIST-
mut33-luciferase reporter was generated by site-directed
mutagenesis with the following primers: 5′-TTTATT-
GAGGACCCATGGTAACATATGAATAGATCCGGTG
TCTAAATGC-3′ (forward) and 5′-GCATTTAGACAC
CGGATCTATTCATATGTTACCATGGGTCCTCAATA
AA-3′ (reverse). The miRNA-33 anti-seed sequence was
converted to NdeI restriction site (underlined in the
primers). Antagomir-33a (427064-00hsa-miR-33amiR-
CURY LNA™ microRNA Power inhibitor) was purchased
from Exiqon (Woburn, MA, USA). miRNAs potentially
able to suppress TWIST expression were selected by
using TargetScan prediction software (http://www.tar-
getscan.org). The miR-Vecs (miRNA expressing vectors)
and MSCV-hITR (control vector) constructs were made as
previously described [18].

miRNA microarray analysis
Total RNA from OS tissues of the discovery cohort of pa-
tients was isolated using TRIzol reagent. The integrity of
RNA was confirmed by agarose gel electrophoresis and
its concentration determined by spectrophotometry. Taq-
Man Low Density miRNA Arrays (Applied Biosystems,
Carlsbad, CA, USA) was used to assay the expression of
human miRNAs by the manufacturer’s protocol. Manual
inspection of all amplification plots was performed and
miRNAs were excluded from the analysis if CT values
were too high (>35, indicating that a miRNA expression is
too low for accurate detection). Data analysis was per-
fomed using SDS 2.3 software (Applied Biosystems),
which utilizes the delta-delta CT method [19].

Real-time quantitative reverse transcription PCR
Total RNA was prepared from OS tissues or cell lines
using TRIzol reagent followed by purification with TURBO
dNA-free System (Ambion, Austin, TX). The cDNAs were
synthesized using SuperScript II reverse transcriptase
(Invitrogen, Carlsbad, CA, USA). Real-time quantitative
PCR was performed using SYBR Green PCR master
mix (Applied Biosystems) in a 7300 Real-time PCR
System (Applied Biosystems). TaqMan microRNA assays
(Applied Biosystems) that include RT primers and
TaqMan probes were used to quantify the expression
of mature miRNA-33a. The mean Ct was determined
from triplicate PCRs. Gene expression was calculated
relative to GAPDH. For measurement of TWIST
mRNA, the following primers were used: for human TWIST,
5′-ACGAGCTGGACCTCCAAGATG-3′ (forward) and 5′-
CAGGCCCTGTTTCTTCTGAAT-3′ (reverse); for human
GAPDH, 5′-GACTCATGACCACAGTCCATGC-3′ (forward)
and 5′-AGAGGCAGGGATGATGTTCTG-3′ (reverse).
The results were normalized against that of the GAPDH
gene in the same sample. Each experiment was repeated
for two times in triplicates.

Western blot analysis
Briefly, cells were dissolved in 250 μl of 2x SDS loading
buffer (62.5 mM TrisHCl, pH 6.8, 2% SDS, 25% glycerol,
0.01% bromphenol blue, 5% 2-mercaptoethanol), and incubated at 95°C for 10 min. Equal amount of proteins for each sample were separated by 10% SDS-polyacrylamide gel and blotted onto a polyvinylidene difluoride microporous membrane (Millipore, Billerica, MA, USA). Membranes were incubated for 1 h with a 1/1000 dilution of primary antibody, and then washed and revealed using secondary antibodies with horseradish peroxidase conjugate (1/5000, 1 h). Peroxidase was revealed with a GE Healthcare ECL kit (Shanghai, China).

Transfection and lentiviral transduction
Plasmid constructs were transfected into cells using Superfect™ transfection reagent (Qiagen) according to the manufacturer's instructions. Pools of stable transfec-
tants of TWIST were generated via selection with G418 (800 μg/ml) by the manufacturer's protocol. Lentiviral transduction of TWIST-shRNA was performed and pools of stable transductants were generated via selection with puromycin (5 μg/ml).

Luciferase assay
MG-63 cells were transfected with luciferase reporter constructs using Superfect™ transfection reagent (Qiagen). Luciferase activity was measured 72 hours after transfection using the Dual-luciferase reporter assay system (Promega) following the manufacturer's instructions. Experiments were conducted in triplicates and results were expressed as ratios between renilla and firefly luciferase counts.

Measurement of apoptosis by TUNEL (terminal deoxynucleotidyl transferase mediated nick-end labeling) assay
The TUNEL assay was performed using the DeadEnd™ Fluorometric TUNEL System by the manufacturer's protocol (Promega). Cells were treated with cisplatin (15 nM) for 8 hours. Apoptotic cells exhibit a strong nuclear green fluorescence that could be detected using a standard fluorescein filter. All cells stained with DAPI exhibit a strong blue nuclear fluorescence. The slides were observed under fluorescence microscopy with relative apoptotic cells determined by counting TUNEL-positive cells in five random fields (magnification, ×100) for each sample.

Statistical analysis
Statistical analyses were performed with SPSS for Windows 10.0. All continuous variable values were expressed as Mean±SD. Comparison of means between two groups was performed with student t tests. Comparisons of means among multiple groups were performed with one-way ANOVA followed by post hoc pairwise comparisons using Tukey's tests. A two-
tailed p < 0.05 was considered statistically significant in this study.

Results
miRNA expression profiling in chemoresistant and control OS
In the discovery cohort, patients were matched by age, sex and tumor stage. Thus, there was no significant difference in age, sex and distribution of tumor stages between poor responders (chemoresistant OS patients, n = 6, tumor necrosis 70.8% ± 9.2%) and good responders (control OS patients, n = 6, tumor necrosis 94.1% ± 2.8%) to neoadjuvant chemotherapy (Table 1). All patients had OS in the long tubular bones. As the inclusion rate for adult OS patients was low, we only performed this study with pediatric OS patients. miRNA microarray analyses

| Table 2 Up-regulated miRNAs in chemoresistant vs control osteosarcomas |
|-----------------------------|-----------------------------|-----------------------------|
| miRNA  | Ct (chemoresistant) | Ct (control) | ΔΔct |
| hsa-miR-132 | 22.81 | 29.40 | −6.57 |
| hsa-miR-21 | 25.16 | 32.13 | −6.97 |
| hsa-miR-33a | 21.57 | 28.70 | −7.13 |
| hsa-miR-215 | 26.97 | 33.13 | −6.16 |
| hsa-miR-221 | 26.15 | 32.07 | −5.92 |
| hsa-miR-140 | 27.62 | 32.78 | −5.16 |
| hsa-miR-375 | 22.46 | 26.92 | −4.46 |
| hsa-miR-181a | 23.22 | 28.09 | −4.87 |
| hsa-miR-30a | 26.62 | 25.75 | −0.87 |
| hsa-miR-25 | 31.54 | 33.87 | −2.33 |
| hsa-miR-200c | 31.96 | 32.86 | −0.90 |
| hsa-miR-26b | 26.70 | 29.17 | −2.47 |
| hsa-miR-23a | 30.05 | 33.08 | −3.03 |
| hsa-miR-625 | 29.36 | 31.12 | −1.76 |
| hsa-miR-363 | 25.73 | 28.24 | −2.51 |
| hsa-miR-17-5p | 31.02 | 32.25 | −1.23 |

| Table 3 Down-regulated miRNAs in chemoresistant vs control osteosarcomas |
|-----------------------------|-----------------------------|-----------------------------|
| miRNA  | Ct (chemoresistant) | Ct (control) | ΔΔct |
| hsa-miR-451 | 33.46 | 29.79 | 3.67 |
| hsa-miR-92a | 33.29 | 29.14 | 4.15 |
| hsa-miR-200a | 32.85 | 28.31 | 4.54 |
| hsa-miR-15b | 32.08 | 27.15 | 4.93 |
| hsa-miR-143 | 31.15 | 24.85 | 6.30 |
| hsa-miR-145 | 30.37 | 23.66 | 6.71 |
| hsa-miR-422a | 30.15 | 22.74 | 7.41 |
| hsa-miR-611 | 29.72 | 22.96 | 6.76 |
| hsa-miR-34c | 28.64 | 21.29 | 7.35 |
Figure 1 Western blot analysis of TWIST expression in chemoresistant and control osteosarcoma (OS) tissues. (A) OS tissue lysates from the chemoresistant OS and the non-chemoresistant control groups (n = 6 each) were subject to western blot analysis for TWIST expression. Glyceraldehyde-3-phosphate dehydrogenase (GAPDH) blotting was used as a loading control. Lanes 1, 3, 5, 7, 9, 11 were samples from the chemoresistant OS group. Lanes 2, 4, 6, 8, 10, 12 were samples from the control group. (B) Density of the TWIST blots was normalized against that of GAPDH to obtain a relative blot density. The relative TWIST blot density of the non-chemoresistant control group was expressed as fold changes to that of the chemoresistant OS group (designated as 1). *p < 0.05 vs chemoresistant OS.

Figure 2 Effects of selected miRNAs on TWIST 3′-untranslated region (UTR). Twelve miRNAs potentially able to regulate TWIST 3′-UTR were selected based on TargetScan prediction software and individually co-transfected with a TWIST-3′UTR-luciferase reporter in MG-63 cells. Among the selected miRNAs, miR-33a, miR-25 and miR-363 were differentially expressed between chemoresistant and control osteosarcoma tissues based on results of the microarray analysis. miRNAs that significantly reduced renilla luciferase activity compared to the control miR-Vec (hTR) (designated as 1, cut off value: 0.8) were selected for validation (light bars). *p < 0.05 vs 0.8 (cut off value).
showed that 25 miRNAs were differentially expressed in OS tissues from the chemoresistant OS patients compared with those from the control OS patients, 16 being up-regulated (Table 2) and 9 down-regulated (Table 3).

**Screening of miRNAs able to regulate TWIST**
The expression of TWIST in the chemoresistant OS patients was significantly lower than that in the control OS patients, either individually or by group (Figure 1), suggesting that TWIST is an important negative regulator in the development of OS chemoresistance. We screened for miRNAs able to regulate TWIST in OS. To this purpose, the 3′-UTR of the TWIST gene was inserted downstream of the renilla luciferase gene in the psi-Check2 vector to generate a TWIST-3′-UTR-luciferase reporter. By combining the microarray expression data with results of the prediction software TargetScan, three up-regulated miRNAs (miR-33a, miR-25 and miR-363) potentially able to regulate TWIST 3′-UTR were selected and individually tested for their ability to affect luciferase expression in MG-63 human OS cells co-transfected with the TWIST-3′-UTR-luciferase reporter.

![Figure 3 Regulation of TWIST 3′-untranslated region (UTR) by miR-33a.](image)

(A) MG-63 cells were co-transfected with a TWIST-3′UTR-luciferase reporter and miR-33a or control miR-Vec (hTR). The relative luciferase activity in cells co-transfected with hTR was designated as 1. (B) Schematic presentation of generation of a TWIST-mut33-luciferase reporter (mut33) by site-directed mutagenesis of the predicted binding sequence of miR-33a in TWIST 3′-UTR. (C) MG-63 cells were co-transfected with the TWIST-3′UTR-luciferase reporter or TWIST-mut33-luciferase reporter and miR-33a or control miR-Vec (hTR). The relative luciferase activity in cells co-transfected with the TWIST-3′UTR-luciferase reporter (wild type) and hTR was designated as 1. *p < 0.05 vs hTR.
In addition, a group of nine miRNAs (miR-33b, miR-32, miR-92a, miR-92b, miR-92c, miR-367, miR-137, miR-137a, and miR-137b) predicted by TargetScan but not differentially expressed in our microarray analysis were also assayed (Figure 2). miR-Vec control (hTR) was used as a negative control. We used MG-63 cells because, according to our microarray data, the predicted miRNAs were absent or expressed at very low levels in the cell line (data not shown). miR-33a, miR-33b and miR-367 significantly reduced renilla luciferase activity compared to the control miR-Vec (cut off value: 0.8) (Figure 2), and miR-33a was found the best candidate to significantly and consistently reduce renilla luciferase activity compared to the control (Figure 3A).

To demonstrate a direct interaction between miR-33a and TWIST, the potential binding sequence for the miRNA within the 3′-UTR of TWIST, as predicted by TargetScan, was mutated to generate a TWIST-mut33-luciferase reporter (Figure 3B). MG-63 cells were co-transfected with miR-33a or miR-Vec control together with either TWIST-3′-UTR-luciferase reporter or TWIST-mut33-luciferase reporter. The reduction of renilla luciferase activity caused by miRNA-33a was specifically abolished by the mutation of the corresponding anti-seed sequence (Figure 3C), suggesting that miR-33a could suppress TWIST expression by acting on its predicted sequence in the 3′-UTR.

To confirm the findings, we determined miRNA-33a and TWIST protein levels in chemoresistant OS patients (n = 35) and control patients (n = 35) in the validation cohort. As shown in Figure 4A, the chemoresistant OS group presented a significantly higher range of miR-33a levels than the control group (0.32 ± 0.08 vs. 0.13 ± 0.05; \( p < 0.001 \)). On the other hand, the chemoresistant OS group presented a significantly lower range of TWIST protein levels than the control group (0.20 ± 0.08 vs. 0.67 ± 0.19; \( p < 0.001 \)) (Figure 4B). Correlation analyses in the entire validation cohort (n = 70) showed that the miR-33a level was negatively correlated with the TWIST protein level in the OS tissue (\( r = -0.627, \ p < 0.001 \)). The miR-33a was negatively correlated with the tumor necrosis rate (\( r = -0.352, \ p < 0.001 \)), while the TWIST protein level was positively correlated with the tumor necrosis rate (\( r = 0.562, \ p < 0.001 \)).

**Effect of overexpression and inhibition of miR-33a on TWIST expression in OS cells**

We next examined the effects of miRNA-33a on TWIST expression in human OS cells. As shown in Figure 5, miR-33a was highly expressed in Saos-2 cells, which had a low constitutive expression of TWIST at both the mRNA and the protein levels. In contrast, MG-63 cells had a constitutive low expression of miR-33a, and a high expression of TWIST at both the mRNA and the protein levels (Figure 5). Thus, overexpression and knockdown of TWIST were respectively performed in the two cell lines to approach the study objectives. As shown in Figure 6A, inhibition of miR-33a by antagonimir-33a increased TWIST expression by over 1.5 fold in Saos-2 cells. On the other hand, overexpression of miR-33a decreased TWIST expression by about 30%. Overexpression of TWIST led to an approximately two-fold increase of TWIST expression in Saos-2 cells, which was largely reversed by overexpression of miR-33a and doubled by antagonimir-33a. As shown in Figure 6B, overexpression of miR-33a decreased TWIST expression by nearly 70% in MG-63 cells, while antagonimir-33a increased TWIST expression by 0.4 fold. Knockdown of
TWIST by shRNA resulted in an approximately 80% decrease of endogenous TWIST expression in MG-63 cells, which was partially reversed by antagomir-33a.

Functional role of miR-33a in TWIST-inhibited OS cell survival against cisplatin

TWIST reportedly decreases OS cell survival against cisplatin, an apoptosis-inducing chemotherapeutic agent commonly used to treat OS [10,11,16]. To explore the effect of interaction between miR-33a and TWIST on OS chemoresistance, we examined cell apoptosis rate in both cell lines treated with cisplatin (15 nM) using TUNEL (terminal deoxynucleotidyl transferase mediated nick-end labeling) assays. Overexpression or knockdown/inhibition of TWIST and/or miR-33a did not significantly alter cell apoptosis in both Saos-2 and MG-63 cells under normal culture conditions (Figure 7A). In Saos-2 cells treated with cisplatin, inhibition of miR-33a by antagomir-33a markedly increased cell apoptosis, which was enhanced by overexpression of TWIST (Figure 7B). The apoptosis-inducing effect of TWIST overexpression was reversed by overexpression of miR-33a (Figure 7B). In MG-63 cells, overexpression of miR-33a significantly decreased cisplatin-induced cell apoptosis, which was enhanced by knockdown of TWIST (Figure 7C). Antagomir-33a significantly increased cisplatin-induced cell apoptosis, which was reversed by knockdown of TWIST (Figure 7C).

Discussion

Chemoresistance is the major reason for poor survival of OS patients. Previous studies reported that TWIST could decrease OS cell survival against cisplatin by inhibiting multiple signaling pathways [10,11], suggesting that TWIST is a pivotal negative regulator of OS chemoresistance. miRNAs reportedly are involved in the pathogenesis and chemoresistance of various cancers, including OS. In the present study, we profiled miRNAs differentially expressed in chemoresistant OS by microarray analysis, with a focus to identify miRNAs that regulate TWIST expression and OS chemoresistance. We provide the first evidence suggesting that miR-33a promotes OS chemoresistance by down-regulating TWIST.

OS is the most common pediatric bone malignancy in the world [1]. As the inclusion rate for adult OS patients was low, we performed this study only in pediatric OS patients. Patients (n = 12) in the discovery cohort were...
Figure 6 (See legend on next page.)
matched on age, sex and tumor stages to reduce the effects of confounders on miRNA profiling between chemoresistant and control OS samples. Patients (n = 70) in the validation cohort were not matched in order to verify the profiling findings in a more generalizable setting. Among the up-regulated miRNAs identified in chemoresistant OS samples in this study, miR-140, miR-215 and miR-221 have been reported to induce human OS chemoresistance [20-22]. Among the down-regulated miRNAs identified in chemoresistant OS samples, miR-451 and miR-15b have been reported to increase chemosensitivity of OS [13]. Thus, our findings were in agreement with previous studies, indicating good reliability of the data.

High expression of TWIST has been detected in several cancers and has been associated with the initial phase of metastatic progression [23]. One recent study reported that TWIST overexpression correlated with disease progression and a poor clinical outcome in OS patients [23]. On the other hand, it has been reported that in homogeneous cohort of OS patients, the TWIST gene was frequently deleted in the tumors at diagnosis, and its haploinsufficiency was significantly correlated with a poorer patient outcome [6,9]. In addition, two recent studies reported that TWIST could decrease OS cell survival against cisplatin by inhibiting β-catenin signaling and endothelin-1/endothelin A receptor signaling pathways [10,11], suggesting that TWIST is an important negative regulator in the development of OS chemoresistance. In this study, our in vitro results showed that overexpression and knockdown of TWIST increased and decreased cisplatin-induced OS cell apoptosis, respectively. This was corroborated by our findings that the expression of TWIST in the chemoresistant OS group was significantly lower than that in the control OS group in both the discovery and validation cohorts, which provides further evidence supporting a critical counteracting role of TWIST in the development of OS chemoresistance.

With an aim to identify miRNAs regulating TWIST expression in OS, we found that miR-33a could significantly down-regulate TWIST expression, which was supported by an inverse miRNA-33a/TWIST expression trend in the validation cohort, target-sequence-specific inhibition of TWIST-3′ UTR-luciferase reporter activity by miR-33a, and alteration of TWIST expression by overexpression or inhibition of miR-33a in human OS cell lines. Saos-2 and MG-63 cells were employed as OS cell models in this study. Saos-2 cells have a constitutive high expression of miR-33a and low expression of TWIST, while MG-63 cells have a constitutive low expression of miR-33a and high expression of TWIST. This explains why inhibition of miR-33a by antagoni-33a had more pronounced effects on TWIST expression than overexpressing miR-33a in Saos-2 cells. Likewise, overexpressing miR-33a had more pronounced effects on TWIST expression than antagoni-33a treatment in MG-63 cells. The effects of overexpression and inhibition of miR-33a on TWIST expression significantly altered OS cell resistance to cisplatin, a chemotherapeutic agent routinely used in neoadjuvant chemotherapy for OS [16]. In the presence of cisplatin, antagoni-33a significantly enhanced cisplatin-induced apoptosis in both Saos-2 and MG-63 cells, suggesting that inhibition of miR-33a could be a potential new strategy to enhance neoadjuvant chemotherapy for OS. The effects of antagoni-33a was reversed and enhanced by knockdown and overexpression of TWIST, respectively, indicating that miR-33a promotes OS cell resistance to cisplatin by down-regulating TWIST, or antagoni-33a enhances cisplatin-induced cell apoptosis by up-regulating TWIST. miR-33a has been shown to regulate genes involved in fatty acid metabolism and insulin signaling [24]. A recent study indicated that miR-33a targets the proto-oncogene Pim-1 and suggested overexpression of miR-33a as an anticancer treatment [25]. However, another study described the down-regulation of tumor suppressor p53 by miR-33 [26], suggesting a complex and possible context-dependent response to miR-33 manipulations. As p53 is often mutated in OS [27], it is unlikely that miR-33a promotes OS chemoresistance through down-regulating p53-induced apoptosis. Thus, the enhancing effect of miR-33a on OS chemoresistance via down-regulating TWIST expression is a new function of this miR, and the miR-33a/TWIST signaling could be a novel mechanism involved in development of OS chemoresistance.
There are some limitations of this study: [1] This study was only performed in pediatric OS patients. Despite that adult OS patients only occupy a small portion of total OS patients, it would still be interesting to verify the findings in adult patients in future studies. [2] Cisplatin elicits DNA repair mechanisms by crosslinking DNA, which in turn activates apoptosis when repair proves impossible [28]. In this study, we only examined the effect of miR-33a/TWIST signaling on OS cell resistance to cisplatin. It is unclear whether miR-33a/TWIST would impact OS cell resistance to other types of chemotherapy agents. Further studies with more types of chemotherapy agents and OS cell lines would elaborate this issue.

In conclusion, we demonstrate that miR-33a is up-regulated in chemoresistant OS and that the miR-33a level is negatively correlated with the TWIST protein level and the tumor necrosis rate in OS. Our in vitro data indicate that miR-33a promotes OS cell resistance to cisplatin by down-regulating TWIST; on the other hand, inhibition of miR-33a by antagonim-33a enhances cisplatin-induced apoptosis in OS cells by up-regulating TWIST expression. The findings suggest that inhibition of miR-33a/TWIST signaling could be a potential new strategy to enhance neoadjuvant chemotherapy for OS.

Competing interests
The authors declare that they have no competing interests.

Authors’ contributions
YZ participated in study design, collected data, carried out data analysis, and drafted the manuscript. SW and XZ participated in study design, carried out data analysis, and performed data check and proofreading. ML and JS participated in data collection, carried out data analysis, and performed data check and proofreading. ZH participated in study design and data analysis, drafted the manuscript, and performed data check and proofreading. All authors have read and approved the final manuscript.
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