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Long noncoding RNA DLEU1 promotes cell proliferation and migration of Wilms tumor through the miR-300/HOXC8 axis†

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Wilms tumor (nephroblastoma) is the most common primary renal tumor occurring in children. Long noncoding RNA (lncRNA) deleted in lymphocytic leukemia 1 (DLEU1) is an identified cancer-associated lncRNA that plays an important role in various cancers. However, the role of DLEU1 in Wilms tumor remains unclear. In the present study, we examined the expression and role of DLEU1 in Wilms tumor. We demonstrated that DLEU1 expression was upregulated in Wilms tumor tissues and cell lines. Knockdown of DLEU1 significantly inhibited the proliferation, migration and invasion of GHINK-1 cells. Furthermore, DLEU1 directly sponged miR-300 and regulated the expression level of miR-300 in GHINK-1 cells. Inhibition of miR-300 reversed the inhibitory effects of DLEU1 downregulation on cell proliferation, migration and invasion. Homeobox C8 (HOXC8) was found to be a target gene of miR-300 and mediated the role of miR-300 in GHINK-1 cells. In conclusion, these findings indicated that DLEU1 executed an oncogenic role in Wilms tumor *via* regulating the miR-300/HOXC8 axis, indicating that DLEU1 might be a therapeutic target for the treatment of Wilms tumor.

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1. Introduction

Wilms tumor, also known as nephroblastoma, is the most common primary renal tumor occurring in children, accounting for 5% of pediatric malignant tumors.^{1,2} It is an embryonic tumor composed of stromal, blastemal, and epithelial cells, and is histologically similar to renal embryogenesis.³ According to the diagnosis and staging assessment of Wilms tumor cases, the treatments include surgical resection, radiotherapy, chemotherapy, or multimodal therapies.⁴ Although the overall 5-year survival rate exceeds 90%, a more successful treatment may reduce the late adverse effects, such as secondary malignancies and cardiac dysfunction.⁵ Therefore, better understanding of the molecular pathogenesis of Wilms tumor is conducive to improve the treatment and outcomes of patient with Wilms tumor.

Long non-coding RNAs (lncRNAs) are transcripts longer than 200 nucleotides with no or limited protein-coding ability.⁶ lncRNAs can interact with DNAs, RNAs and proteins to regulate gene expression at epigenetic, transcriptional and post-transcriptional levels.^{7,8} Increasing amounts of evidence have demonstrated that lncRNAs are closely associated with the occurrence and progression of cancers.^{9,10} Recent studies have indicated that lncRNAs may be therapeutic target in the cancer

treatments.¹¹ However, the bio-functions and molecular mechanisms of lncRNAs in Wilms tumor remain limited. Zhu *et al.*¹² reported that LINC00473 functions as an oncogene in Wilms tumor *via* microRNA (miR)-195/IκB kinase (IKK). lncRNA LINP1 enhances cell metastasis and proliferation of Wilms tumor *via* inducing the Wnt/β-catenin signaling pathway.¹³

lncRNA deleted in lymphocytic leukemia 1 (DLEU1) is an identified oncogenic lncRNA in various cancers, such as lung cancer, colorectal cancer, cervical cancer, endometrial cancer, and ovarian carcinoma.^{14–18} DLEU1 expression is upregulated in colorectal cancer tissues, the higher expression of DLEU1 indicates lower survival rate and poorer prognosis.¹⁵ DLEU1 promotes tumorigenesis and progression of non-small cell lung cancer (NSCLC) and might be a promising therapeutic target for NSCLC.¹⁴ These findings suggest that DLEU1 play a crucial role in the tumorigenesis. However, the role of DLEU1 in Wilms tumor remains unclear.

In the present study, we found that DLEU1 expression was increased in Wilms tumor tissues and exerted an oncogenic role. Therefore, we speculated that DLEU1 might be a therapeutic target in the Wilms tumor treatment.

2. Results

2.1 DLEU1 is highly expressed in Wilms tumor tissues and cell lines

To reveal the role of DLEU1 in Wilms tumor, qRT-PCR analysis was performed to determine the expression level of DLEU1 in Wilms tumor samples and adjacent non-tumor tissues. As

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revealed in Fig. 1A, DLEU1 was upregulated in Wilms tumor samples compared with non-tumor tissues ($p < 0.05$). Then we tested the DLEU1 expression in Wilms tumor cell lines including GHINK-1, SK-NEP-1 and G401 cells. The results showed that DLEU1 was highly expressed in Wilms tumor cell lines (Fig. 1B).

2.2 Knockdown of DLEU1 inhibits the proliferation, migration and invasion of Wilms tumor cells

To further investigate the roles of DLEU1 on Wilms tumor, GHINK-1 cells were transfected with si-DLEU1 or si-NC. The knockdown of DLEU1 was confirmed by qRT-PCR analysis (Fig. 2A). The results from MTT assay indicated that GHINK-1 cells with lower DLEU1 exhibited reduced proliferation as compared to the GHINK-1 cells transfected with si-NC (Fig. 2B). Additionally, cell migration and invasion were respectively decreased by si-DLEU1 (Fig. 2C and D). Furthermore, we found that DLEU1 had no significant effect on cell proliferation and migration of normal kidney HK2 cells (ESI Fig. 1†).

2.3 DLEU1 targets miR-300 in Wilms tumor cells

We next analyzed the molecular mechanism of DLEU1 using the online software to predict the target miRNA of DLEU1. As shown in Fig. 3A, putative complementary sequences between DLEU1 and miR-300 were found. Luciferase reporter assay proved that co-transfection with psiCHECK2-DLEU1 and miR-300 mimics significantly reduced the luciferase activity (Fig. 3B). In addition, miR-300 expression was significantly increased by si-DLEU1 in GHINK-1 cells (Fig. 3C). Furthermore, correlation analysis showed that there was a negative correlation between DLEU1 expression and miR-300 expression in Wilms tumor tissues (Fig. 3D).

2.4 MiR-300 regulates the effects of DLEU1 on cell proliferation, migration and invasion

To better understand the role of miR-300 in Wilms tumor, the expression of miR-300 was examined. As illustrated in Fig. 4A and B, the expressions of miR-300 were markedly decreased in

Wilms tumor tissues and cell lines. Furthermore, we observed that the decreased cell proliferation caused by si-DLEU1 was mitigated by transfection with miR-300 inhibitor (Fig. 4C). Moreover, miR-300 inhibitor also attenuated si-DLEU1-caused decrease in cell migration and invasion (Fig. 4D and E). Furthermore, we found that miR-300 mimics reserved the promotion effects of over-DLEU1 on GHINK-1 cell proliferation, migration and invasion (ESI Fig. 2A–C†).

2.5 MiR-300 targets HOXC8 in Wilms tumor cells

It has been well-known that miRNAs usually bind to the 3'UTR of target mRNA and thereby regulating gene expression.^{19,20} Then we used bioinformatic prediction software to explore the potential target gene of miR-300. We identified that miR-300 was able to bind to the 3'UTR of HOXC8 (Fig. 5A). To confirm the interaction between miR-300 and HOXC8, a luciferase assay was performed and demonstrated that miR-300 dramatically reduced the luciferase activity of the psiCHECK2-HOXC8 (Fig. 5B). Moreover, miR-300 mimics significantly suppressed the HOXC8 expression in GHINK-1 cells (Fig. 5C).

2.6 HOXC8 overexpression reverses the effects of miR-300 mimics on cell proliferation, migration and invasion

To validate whether HOXC8 mediated the role of miR-300, HOXC8-overexpressing plasmid pcDNA3.1-HOXC8 was transfected into GHINK-1 cells. The results from qRT-PCR and western blot showed that HOXC8 expression was markedly elevated after transfection with pcDNA3.1-HOXC8, compared with transfection with pcDNA3.1 vector (Fig. 6A and B). Moreover, the inhibitory effects of miR-300 mimics on cell proliferation, migration and invasion were mitigated by HOXC8 overexpression (Fig. 6C–E).

3. Discussion

Increasing evidences have indicated that many lncRNAs are involved in the tumorigenesis of various types of cancers. However, the role of lncRNAs in Wilms tumor remains largely unknown. In the present study, we first evaluated the function

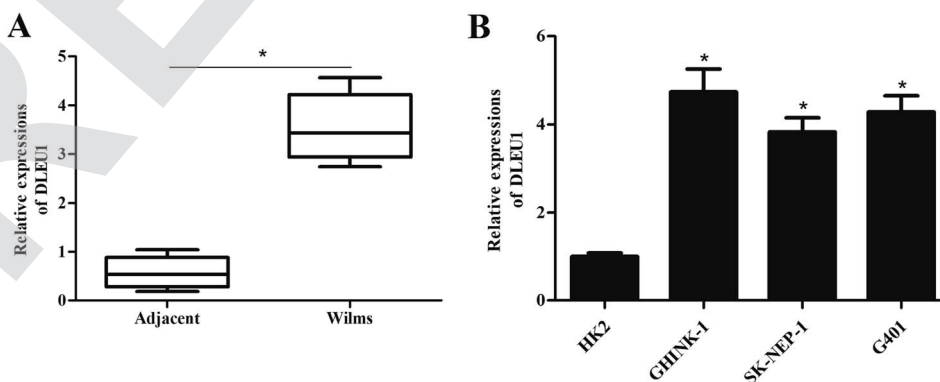


Fig. 1 DLEU1 expression levels in tissue specimens and cultured cells. To reveal the role of DLEU1 in Wilms tumor, qRT-PCR analysis was performed to determine the expression levels of DLEU1 in Wilms tumor samples and cell lines. (A) Expression levels of DLEU1 in Wilms tumor samples and adjacent non-tumor tissues. (B) Expression levels of DLEU1 in Wilms tumor cell lines. * $p < 0.05$.



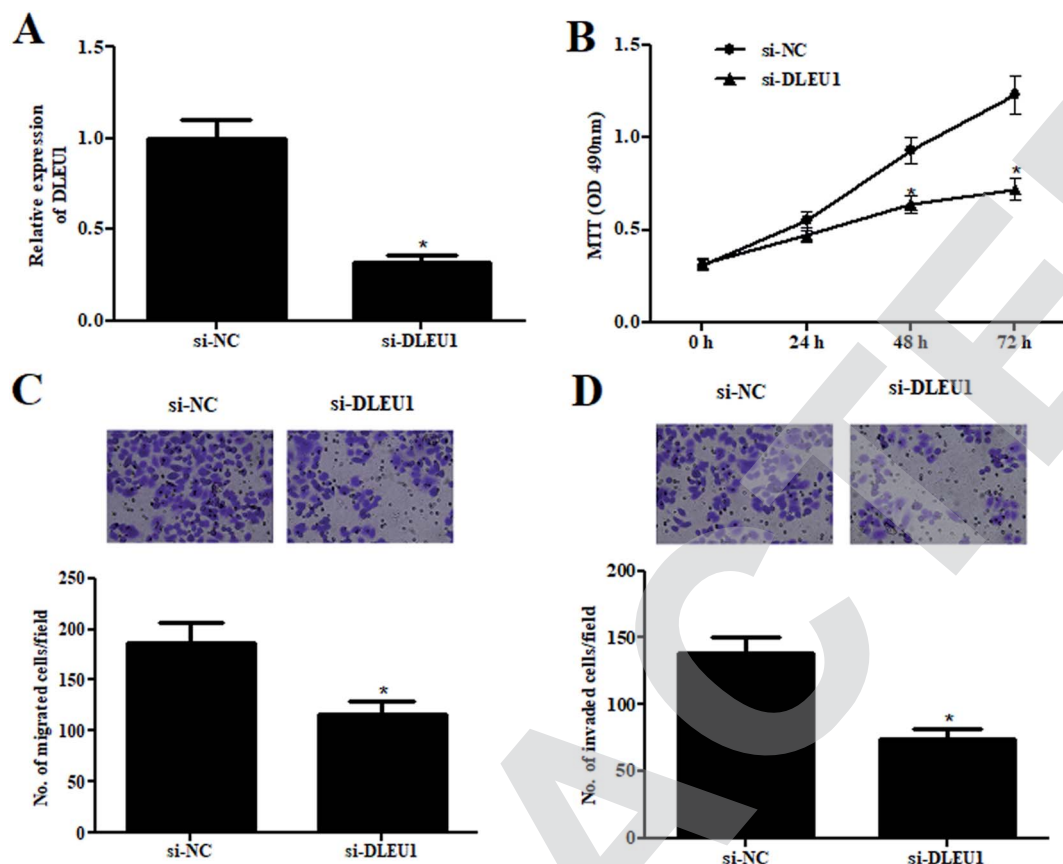


Fig. 2 Wilms tumor cells proliferation, migration and invasion were decreased in si-DLEU1-transfected cells. (A) The knockdown of DLEU1 was confirmed by qRT-PCR analysis after transfection with si-DLEU1 or si-control. (B) Cell proliferation was detected by MTT assay. (C and D) Cell migration and invasion were determined using transwell assay. * $p < 0.05$.

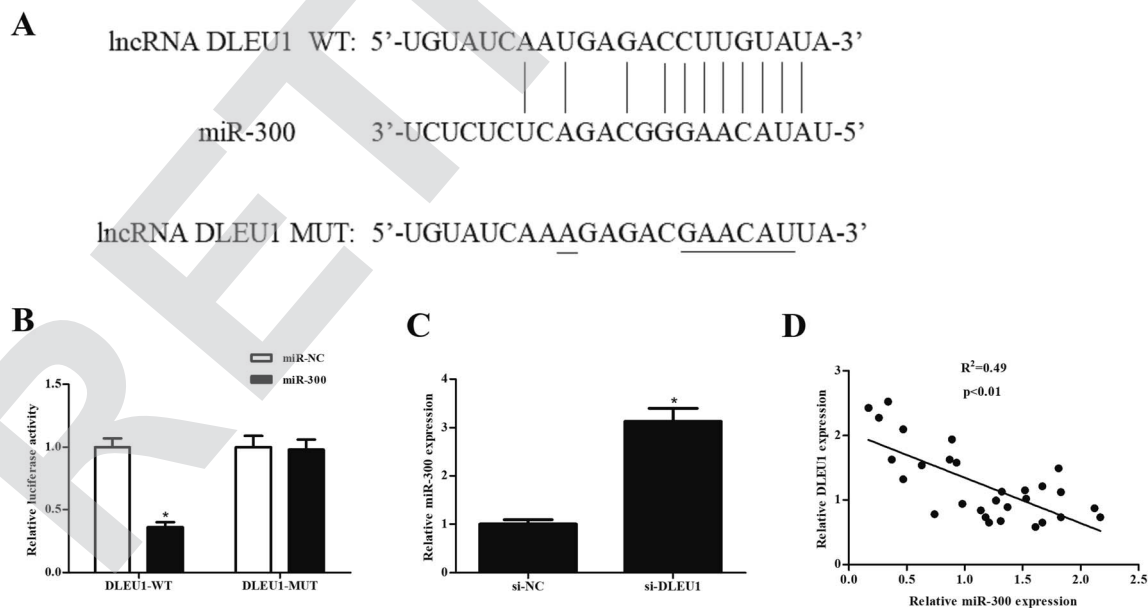


Fig. 3 MiR-300 might be a target mRNA of DLEU1 in GHINK-1 cells. (A) Putative complementary sequences between DLEU1 and miR-300. (B) Luciferase reporter assay was performed to confirm the interaction of DLEU1 and miR-300. * $p < 0.05$ vs. psiCHECK-DLEU1-MUT and miR-300 mimics co-transfected cells. (C) Knockdown of DLEU1 elevated the expression level of miR-300 in GHINK-1 cells. (D) Correlation analysis showed that there was a negative correlation between DLEU1 expression and miR-300 expression in Wilms tumor tissues. * $p < 0.05$.



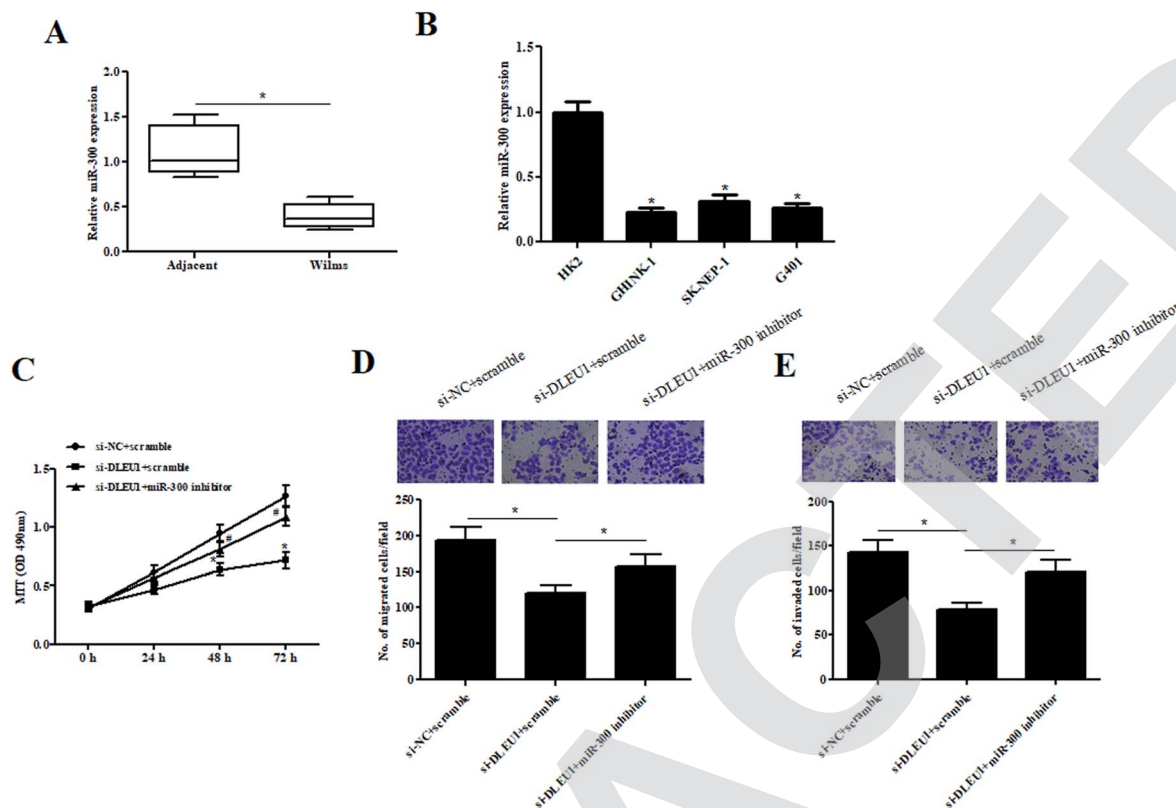


Fig. 4 The effects of si-DLEU1 on GHINK-1 cells were reversed by miR-300 inhibitor. (A and B) Expression of miR-300 was detected by qRT-PCR analysis in Wilms tumor samples and cell lines. (C) Cell proliferation was detected by MTT assay. * $p < 0.05$ vs. si-NC + scramble group; # $p < 0.05$ vs. si-DLEU1 + scramble group. (D and E) Cell migration and invasion were determined using transwell assay. * $p < 0.05$.

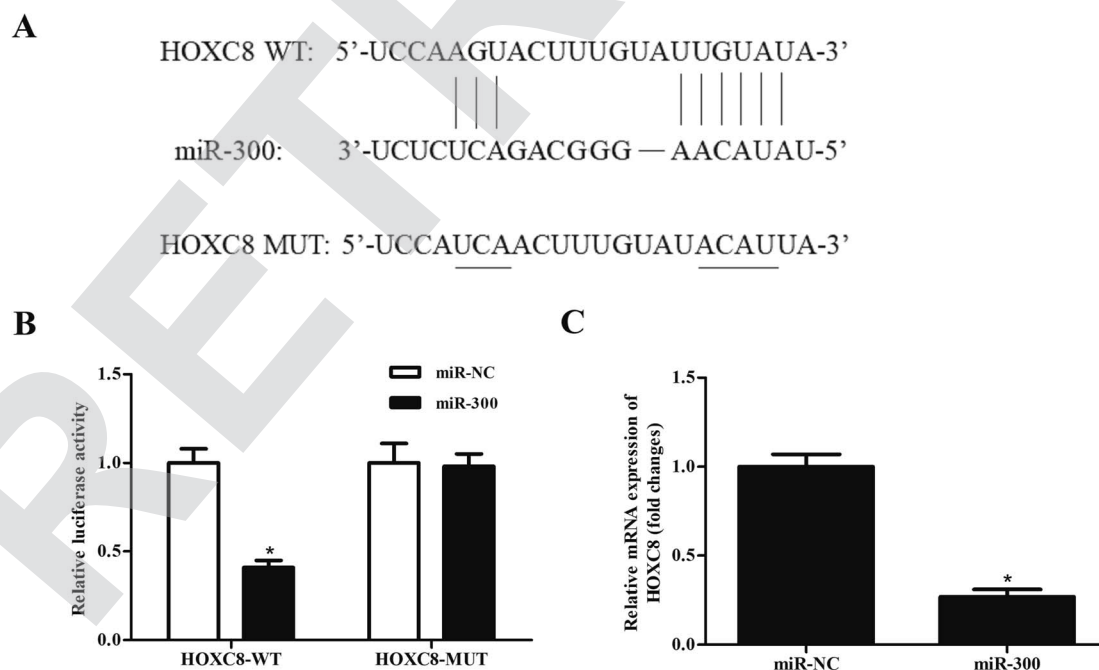


Fig. 5 HOXC8 might be a target gene of miR-300 in GHINK-1 cells. (A) Putative complementary sequences between miR-300 and HOXC8 predicted by bioinformatic software. (B) Luciferase reporter assay was carried out to confirm the interaction of miR-300 and HOXC8. * $p < 0.05$ vs. psiCHECK-HOXC8-MUT and miR-300 mimics co-transfected cells. (C) Overexpression of miR-300 suppressed the expression level of HOXC8 in GHINK-1 cells. * $p < 0.05$.



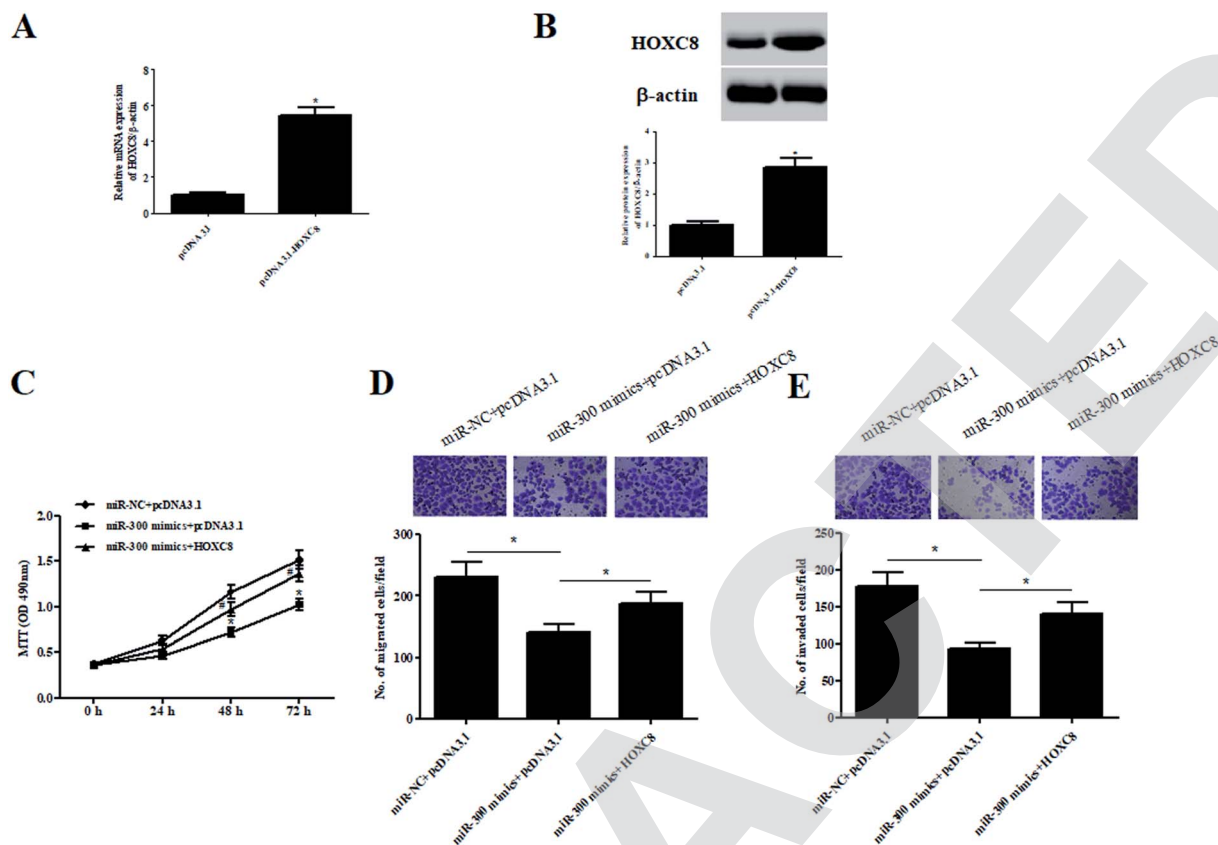


Fig. 6 The inhibitory effects of miR-300 mimics on cell proliferation, migration and invasion were mitigated by HOXC8 overexpression. (A and B) Expression of HOXC8 was detected by qRT-PCR and western blot analysis after transfection with pcDNA3.1-HOXC8 or pcDNA3.1. (C) Cell proliferation was detected by MTT assay. * $p < 0.05$ vs. miR-NC + pcDNA3.1 group; # $p < 0.05$ vs. miR-300 mimics + pcDNA3.1 group. (D and E) Cell migration and invasion were determined using transwell assay. * $p < 0.05$.

of DLEU1 in Wilms tumor. The results showed that DLEU1 expression was upregulated in Wilms tumor tissues and acted as an oncogene. Further investigations proved that DLEU1 sponged miR-300 in GHINK-1 cells and regulated the expression of HOXC8, indicating that DLEU1 exerted its oncogenic role *via* regulating miR-300/HOXC8 axis.

Previous studies have demonstrated that DLEU1 promotes the progression of many cancers. DLEU1 is highly expressed in endometrial carcinoma and contributes to tumorigenesis and development of endometrial carcinoma both *in vitro* and *in vivo*.²¹ DLEU1 expression is closely associated with advanced tumor-node-metastasis stage, vascular metastasis and poor overall survival in hepatocellular carcinoma patients, indicating that DLEU1 promotes hepatocellular carcinoma progression.²² DLEU1 is overexpressed in osteosarcoma tissue specimens, moreover, downregulation of DLEU1 in osteosarcoma cells inhibits the cell proliferation, migration and invasion, indicating that DLEU1 aggravates osteosarcoma carcinogenesis.²³ Our results proved that DLEU1 expression was significantly upregulated in Wilms tumor tissues. *In vitro* investigations revealed that downregulation of DLEU1 inhibited the proliferation, migration and invasion of Wilms tumor cells. However, according to a previous study, DLEU1 functions as a tumor suppressor gene and confer chemoimmunotherapy resistance

in children and adolescents with Burkitt lymphoma (BL).²⁴ These findings suggest that the role of DLEU1 in cancers may depend on the cancer type.

It has been suggested that DLEU1 acts as endogenous miRNA sponges by binding to target miRNAs. Chen *et al.*²³ reported that DLEU1 acts as an oncogene in osteosarcoma by directly sponging miR-671-5p. DLEU1 promotes cervical cancer cell proliferation and invasion by interacting with miR-381.¹⁶ Therefore, we used bioinformatic software to predict the target miRNA of DLEU1, and we found putative complementary sequences between DLEU1 and miR-300. DLEU1 directly bound to miR-300 and regulated the miR-300 expression in GHINK-1 cells. Recent studies have denoted that miR-300 is associated with tumor development. For instance, Liu *et al.*²⁵ showed that miR-300 levels in the serum of osteosarcoma patients are higher than healthy controls and correlates with clinical stage, distant metastasis and poor survival, indicating that serum miR-300 may be an independent prognostic marker for osteosarcoma. MiR-300 promotes proliferation and epithelial-mesenchymal transition (EMT)-mediated migration and invasion in colorectal cancer cells.²⁶ Additionally, miR-300 expression is down-regulated in laryngeal squamous cell carcinoma (LSCC) tissues. Overexpression of miR-300 inhibits cell proliferation and invasion in Hep-2 cells.²⁷ Our results proved that miR-300



overexpression reduced cell proliferation, migration and invasion of GHINK-1 cells. Inhibition of miR-300 reversed the effects of DLEU1 knockdown on cell proliferation, migration and invasion. These findings suggested that miR-300 exhibited tumor suppressive activity, which contributed to the oncogenic activity of DLEU1.

To analyze the molecular mechanism of miR-300, we found that HOXC8 might be a target gene of miR-300. MiR-300 significantly suppressed the expression of HOXC8 in GHINK-1 cells. Interestingly, HOXC8, a member of HOX family, has been proven to play a crucial role in cancer development. HOXC8 represents an interesting tumor suppressor candidate in breast cancer since it can regulate self-renewal, differentiation and transformation of breast cancer stem cells.²⁸ HOXC8 is involved in the tumorigenesis of osteosarcoma with predictive potential and oncogenic effects.²⁹ Significantly increased expression of HOXC8 has been observed in cervical cancer tissues and cell lines. Downregulation of HOXC8 inhibits the proliferation rate of cervical cancer cells, which suggest that HOXC8 may serve as a therapeutic target for cervical cancer.³⁰ Liu *et al.* reported that HOXC8 was upregulated in NSCLC specimens, and overexpression of HOXC8 significantly promoted the proliferation, anchorage-independent growth and migration of NSCLC cells.³¹ These results indicated that targeting HOXC8 may be new strategy for the therapeutic intervention. Our study proved that HOXC8 was a target gene of miR-300 and mediated the effects of miR-300 on GHINK-1 cells, suggesting an oncogenic role of HOXC8 in Wilms tumor.

In summary, we demonstrated that DLEU1 expression was upregulated in Wilms tumor tissues and cell lines. Knockdown of DLEU1 inhibited the proliferation, migration and invasion of Wilms tumor cells. Furthermore, DLEU1 executed its oncogenic role in Wilms tumor *via* regulating miR-300/HOXC8 axis.

4. Materials and methods

4.1 Clinical tissues and cell culture

Thirty pairs of Wilms tumor tissues and adjacent normal tissues were collected from patients who has Wilms tumor underwent surgery at Xi'an Children's Hospital (Xi'an, China) from July 2016 to June 2017. All samples were removed and immediately frozen in liquid nitrogen and stored at $-80\text{ }^{\circ}\text{C}$ until use. All experiments were performed in compliance with the Ethics Committee of Xi'an Children's Hospital and approved by the Ethics Committee of Xi'an Children's Hospital. Informed consent was obtained for any experimentation with human subjects.

A human immortalized normal kidney cell line (HK2; Shanghai Model Cell Bank, Shanghai, China) and Wilms' Tumor cell lines including GHINK-1 (RIKEN, Wako, Japan), SK-NEP-1 and G401 (Shanghai Model Cell Bank) were cultured in Dulbecco's Modified Eagle Medium (DMEM; Invitrogen, Carlsbad, CA, USA) supplemented with 10% fetal bovine serum (FBS; Invitrogen), 100 U ml^{-1} penicillin and $100\text{ }\mu\text{g ml}^{-1}$ streptomycin (Invitrogen). All cells were maintained in an incubator with 5% CO_2 at $37\text{ }^{\circ}\text{C}$.

4.2 Cell transfection

Small interfering RNA (siRNA) targeting DLEU1 (si-DLEU1) and nonspecific negative control siRNA (si-NC) were synthesized (Biotend, Shanghai, China). MiR-300 mimics, miR-300 inhibitor and non-targeting scramble controls were purchased from Genechem (Shanghai, China). Homeobox C8 (HOXC8) coding sequence was inserted into pcDNA3.1 vector to construct HOXC8-overexpressing plasmid pcDNA3.1-HOXC8. Cell transfection was performed using Lipofectamine 3000 Reagent (Invitrogen) according to the manufacturer's protocols.

4.3 Cell proliferation assay

Cell proliferation was evaluated by MTT assay. The cells (5×10^3 cell per well) with different treatments were cultured for 0, 24, 48, or 72 h, $20\text{ }\mu\text{l}$ MTT were added into each well and continued to culture for another 4 h. After that, $150\text{ }\mu\text{l}$ DMSO was added to the cells, and then the plates were placed on the oscillator for 10 min to dissolve the crystal. The optical density (OD) values were detected and recorded with a microplate reader (Bio-Tek, Winooski, VT, USA) at 490 nm wavelength.

4.4 Cell migration and invasion assays

GHINK-1 cells (5×10^4 cells) in serum-free culture medium were placed in the upper chamber of Transwell chambers (Corning Inc., Corning, NY, USA) for the transwell assay. For the determination of cell invasive ability, the chambers were coated with Matrigel (Sigma). The lower chamber was added with normal medium containing 10% FBS. After 24 h incubation, the cells remained on the upper side of the filters were removed with a cotton swab. Cells moved to the lower side of the filters were fixed using methanol and stained with 0.05% crystal violet. Five random pictures were taken, and the cell numbers were counted under microscope.

4.5 qRT-PCR

TRIzol reagent (Invitrogen) was used for the extraction of the total RNA from tissue samples or culture cells. Then the total RNA (500 ng) was transcribed to cDNA using the PrimeScriptRT Kit (TaKaRa, Dalian, China). The relative expression levels of DLEU1 and HOXC8 were evaluated by qRT-PCR using STBR Premix Ex Taq (TakaRa) according to the instructions. The relative expression levels of miR-300 were measured using All-in-One miRNA qRT-PCR Detection Kit (GeneCopoeia, Rockville, MD, USA). The primer sequences were listed as following: DLEU1 F 5'-TCA GAA TGC CGA CTC TAT GCT-3'; R 5'-GGT GAG GAC AGA GTT AAA CGC-3'; HOXC8 F 5'-GGC CAT AGA GAT TAG GGG TTC-3'; R 5'-GGG GCT CAT GAC CTA AGC TA-3'; GAPDH F 5'-AAT GGG CAG CCG TTA GGA AA-3'; R 5'-TGA AGG GGT CAT TGA TGG CA-3'; MiR-300 F 5'-TAT ACA AGG GCA GAC TCT CTC T-3'; R 5'-GTG CAG GTT CCG AGG T-3'; U6 F 5'-CTC GCT TCG GCA GCA CAT ATA CT-3'; R 5'-ACG CTT CAC GAA TTT GCG TGT C-3'.



4.6 Western blot

GHINK-1 cells were harvested and lysed with RIPA lysis buffer containing 1% v/v PMSF (Beyotime, Haimen, China). A total of 40 µg proteins were separated by 10% sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE), and then the separated proteins were transferred onto polyvinylidene difluoride (PVDF) membranes (Thermo Fisher Scientific, Waltham, MA, USA). The membranes were then blocked in 5% w/v non-fat milk solution at room temperature for 1 h, followed by an incubation with a specific primary antibody against HOXC8 (PA5-27979; 1 : 3000; Invitrogen) or β-actin (PA5-85490; 1 : 5000; Invitrogen) overnight at 4 °C. Then the membranes were incubated with HRP-conjugated secondary antibody (A32740; 1 : 2500; Invitrogen) at room temperature for 1 h. Immune complexes were visualized using the enhanced chemiluminescence (ECL) system (Thermo).

4.7 Target prediction

LncRNATargets (<http://starbase.sysu.edu.cn>) was used to predict target miRNAs of DLEU1. TargetScan (<http://www.targetscan.org>) was applied for the prediction of the target genes of miR-300.

4.8 Luciferase reporter assay

According to the predicted results from bioinformatics software, the wide-type sequences of DLEU1 (DLEU1-WT) and the HOXC8 3'UTRs containing the predicted miR-300 binding sites were respectively inserted into the psiCHECK vector (Promega Corporation, Madison, WI, USA) to generate psiCHECK-DLEU1-WT and psiCHECK-HOXC8-WT plasmids. Meanwhile, the mutant type of DLEU1 (DLEU1-MUT) or HOXC8 (HOXC8-MUT) was inserted into psiCHECK vector to construct psiCHECK-DLEU1-MUT and psiCHECK-HOXC8-MUT plasmids. GHINK-1 cells (2.5×10^5 cells per well) were cultured in a 24-well plate 24 h prior to transfection. The psiCHECK-DLEU1-WT/psiCHECK-DLEU1-MUT or psiCHECK-HOXC8-WT/psiCHECK-HOXC8-MUT and miR-300 mimics/control mimics were used for co-transfection of GHINK-1 cells using Lipofectamine 2000 (Invitrogen). After 48 h, luciferase activities were determined with a dual-luciferase assay kit (Promega) following the manufacturer's protocols.

4.9 Statistical analysis

All experiments were repeated for three times, and the data were presented as mean ± standard deviation. The analyses of the data were performed by SPSS 19.0 software (IBM, Chicago, IL, USA) using unpaired two-tailed Student's *t*-test or one-way analysis of variance (ANOVA). A value of $p < 0.05$ was considered as statistically significant.

Conflicts of interest

There are no conflicts to declare.

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