

# Upregulation of microRNA-1270 suppressed human glioblastoma cancer cell proliferation migration and tumorigenesis by acting through WT1

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**Background:** Glioblastoma multiforme (GBM) is one of the most aggressive brain tumors among human beings. In this study, we explored the functions of human microRNA-1270 (hsa-miR-1270) on GBM cancer cell proliferation, migration, and tumorigenesis.

**Materials and methods:** In GBM cell lines and clinical tissues, hsa-miR-1270 expression was probed by quantitative real-time PCR (qRT-PCR). In LN-18 and A172 cells, hsa-miR-1270 was upregulated by lentiviral transduction. The effects of hsa-miR-1270 upregulation on GBM in vitro and in vivo functions were probed by proliferation, migration, and xenograft assays, respectively. The correlation between hsa-miR-1270 and Wilms' tumor gene (*WT1*) was probed by dual-luciferase activity assay, qRT-PCR, and Western blot. *WT1* was then secondarily over-expressed in hsa-miR-1270-upregulated LN-18 and A172 cells, to explore its mechanisms in GBM's association with hsa-miR-1270.

**Results:** Hsa-miR-1270 was significantly downregulated in both GBM cell lines and clinical tumors. Upregulating hsa-miR-1270 considerably suppressed GBM cell proliferation and migration in vitro and xenograft in vivo. *WT1* was inversely correlated with hsa-miR-1270 in GBM. *WT1* overexpression in hsa-miR-1270-upregulated GBM cells reversed the anticancer functions of hsa-miR-1270 on cancer proliferation and migration.

**Conclusion:** Hsa-miR-1270 upregulation may have suppressing effects on GBM cancer cells, likely by functionally acting through *WT1*.

**Keywords:** GBM, miRNA, hsa-miR-1270, *WT1*, xenograft

## Introduction

Glioblastoma multiforme (GBM), or grade IV astrocytoma, is a heterogeneous, fast-growing, and aggressive type of brain tumor. Once diagnosed with GBM, cancer patients are often associated with high relapse rate and high mortality rate.<sup>1-3</sup> Although significant progress had been made over the past decades to develop more treatment options for patients with GBM, such as antiangiogenic therapy, molecule-targeted therapies, immunotherapy, or gene therapy, the prognosis remains very poor for those patients with 5-year survival rates to be around 5%.<sup>4-7</sup> Thus, it is very critical to explore the heterogeneous pathology, understand its underlying mechanisms, and seek effective therapeutic targets for GBM.

miRNAs are families of single-stranded, noncoding small (~18–22 nucleotide long) RNAs that attach to the complementary DNA sequences on the 3'-UTR of their downstream targeting genes, inducing gene silencing, thus yielding significantly functional regulations on molecular and cellular developments in both humans and animals.<sup>8-11</sup> In addition, miRNAs were demonstrated to play important roles in

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regulating pathology among many human diseases, such as cancer.<sup>12–15</sup> Specifically, in human brain tumor of glioma, miRNAs were shown to be aberrantly dysregulated in cancer patients' tumor tissues or plasma, and may induce functional regulations on cancer cell development, maturation, metastasis, or drug resistance.<sup>16–18</sup>

Within the families of human cancer-related miRNAs, human mature microRNA 1270, hsa-miR-1270, was discovered to be either upregulated or downregulated in several types of human cancers, for example, upregulated in osteosarcoma or renal cancer,<sup>19,20</sup> but downregulated in endometrial cancer.<sup>21</sup> In addition, hsa-miR-1270 was found to be associated with several cancer-regulating genes, such as glypican 5 or histone deacetylase 4 in human non-small-cell lung cancer,<sup>22,23</sup> and alpha fetoprotein in hepatoma.<sup>24</sup> However, there has been little evidence directly linking hsa-miR-1270 with functional regulations in human cancers.

In a very recent study, Sun et al demonstrated that, among human glioma subtypes, hsa-miR-1270 might be downregulated in grade IV astrocytoma or GBM.<sup>25</sup> To follow up on this discovery, we conducted a systematic approach to explore the expression profile and functional mechanisms of hsa-miR-1270 in human GBM. The results of this study may help to identify the clinical implication of hsa-miR-1270, as to whether it may be used as an epigenetic target for future molecular therapy for patients with GBM.

## Materials and methods

### GBM cells and clinical tissues

Five human GBM cell lines, LN-18, LN-229, U-138MG, T98G, and A172 cells, were commercially obtained from American Type Culture Collection (Manassas, VA, USA). In addition, a normal human astrocyte (NHA) cell line was commercially obtained from the Chinese Academy of Sciences Cell Bank of Type Culture Collection (Shanghai, China). The cell lines were all cultured in RPMI-1640 medium (Thermo Fisher Scientific, Waltham, MA, USA) supplemented with 10% FBS (Thermo Fisher Scientific), 100 U/mL penicillin (Thermo Fisher Scientific), and 100 mg/mL streptomycin (Thermo Fisher Scientific), in a humidified environment at 37°C with 5% CO<sub>2</sub>.

Clinical tissues were obtained from eleven GBM patients who received maximal surgical resection at the Second Affiliated Hospital of Chongqing Medical University between June 2012 and November 2017. GBM tissues were identified by independent magnetic resonance imaging and histology evaluations according to the WHO classifications.<sup>26</sup> Paired

samples, including GBM tissues and adjacent normal brain tissues, were retrieved, and immediately snap-frozen in liquid nitrogen. Those tissues were then stored in a –70°C biograde freezer until further processing.

### RNA extraction and quantitative real-time PCR (qRT-PCR)

Total RNA was extracted from GBM cell lines or clinical tissues using a PureLink miRNA Isolation Kit (Thermo Fisher Scientific) according to the manufacturer's recommendation. Then, a TaqMan™ MicroRNA Reverse Transcription Kit (Thermo Fisher Scientific) was applied to convert mRNA to cDNA. qRT-PCR was then conducted using an ABI Prism 7900 HT Fast-Real time PCR System (Applied Biosystems, Waltham, MA, USA) according to the manufacturer's recommendation. For quantification on hsa-miR-1270 gene expression, a mirVana™ qRT-PCR miRNA Detection Kit (Invitrogen, Waltham, MA, USA) was used according to the manufacturer's recommendation, and U6 small nuclear RNA (U6 sRNA) was used as a reaction template. Alternatively, for quantification on Wilms' tumor gene (*WT1*), a PowerUp™ SYBR™ Green Master Mix kit (Invitrogen) was used according to the manufacturer's recommendation, and glyceraldehyde 3-phosphate dehydrogenase was used as reaction template. Finally, relative gene expression levels were quantified using the 2<sup>–ΔΔCt</sup> method.

### Hsa-miR-1270 overexpression assay

A lentivirus with the insertion of an hsa-miR-1270 mimics, miR-1270M, was commercially obtained from Shanghai Genepharma Co., Ltd. (Shanghai, China). Correspondingly, another lentivirus with the insertion of a nonspecific human miRNA mimics, miR-MC, was also obtained from Shanghai Genepharma. Among GBM cell lines, LN-18 and A172 cells were transduced with miR-MC or miR-1270M for 48 hours, and then selected by blasticidin (8 μg/mL, Sigma Aldrich, St Louis, MO, USA) for 72 hours. After that, healthy cell colonies were selected and passaged for three times. Then, qRT-PCR was conducted to examine hsa-miR-1270 expression in lentiviral-transduced GBM cells.

### In vitro proliferation assay

LN-18 and A172 cells were seeded in 96-well plates (1.5 × 10<sup>3</sup> cells/well), and allowed to proliferate for five consecutive days. Their in vitro proliferation was examined using a CyQUANT® NF Cell Proliferation Assay Kit (Thermo Fisher Scientific) according to the manufacturer's

recommendation, and quantified as absorbance readings at 570 nm (Multiskan, Thermo Fisher Scientific).

### In vitro migration assay

LN-18 and A172 cells were seeded in the upper chambers of a Transwell plate (24-well, 8  $\mu$ m, VWR, USA). The Transwell membranes were precoated with Matrigel (0.1%, Sigma Aldrich). Cells in the upper chambers were maintained in culture medium without FBS ( $1.5 \times 10^4$  cells/well), whereas the lower chambers were filled with serum-supplemented regular culture medium. The Transwell plate was kept in a humidified environment at 37°C with 5% CO<sub>2</sub> for 48 hours. Then, Transwell inserts, floating cells, and culture medium were all removed. LN-18 and A172 cells migrated to the bottoms of Transwell plates were fixed, stained with 0.1% crystal violet (Sigma Aldrich) for 20 minutes at room temperature, and then imaged. In each well, the numbers of migrated GBM cells were measured for five randomly selected areas (200  $\mu$ m  $\times$  200  $\mu$ m). GBM cancer migration was then quantified as the percentages of migrated GBM cells in each well against migrated GBM cells in control wells.

### In vivo xenograft assay

GBM LN-18 cells, after lentiviral transduction, were xenografted into the subcutaneous compartments of male athymic nude mice (5-week old,  $1 \times 10^6$  cells/injection). Twelve mice received xenografts transduced with miR-MC lentivirus. Fourteen mice received xenografts transduced with miR-1270M lentivirus. Every week, length (L, mm) and width (W, mm) of LN-18 xenografts were measured, and their volumes (V, mm<sup>3</sup>) were estimated based on a formula,  $V=L*W*W/2$ . After 5 weeks, mice were killed. LN-18 xenografts were removed from mice bodies and compared under the microscope.

### Dual-luciferase reporter assay

The 3'-UTR of human *WT1* gene was inserted into a pmiR-REPORT luciferase vector to generate a wild-type luciferase plasmid, WT1-WT (Shanghai Genepharma). Alternatively, at position 910–916 and position 2329–2335 of human *WT1* DNA sequence, the sequences were mutated to nullify hsa-miR-1270 binding. The mutant *WT1* 3'-UTR was also inserted into pmiR-REPORT to generate another luciferase plasmid, WT1-MU (Shanghai Genepharma). In a dual-luciferase reporter assay, HEK293T cells were pretransduced with miR-MC or miR-1270M, followed by secondary transfection of WT1-WT or WT1-MU for 24 hours. Relative luciferase activities were measured using a

Multiskan plate-reader (Thermo Fisher Scientific) according to the manufacturer's recommendation.

### Western blotting assay

LN-18 and A172 cells were lysed with a Pierce™ IP Lysis Buffer (Thermo Fisher Scientific). Equal amount of extracted protein samples were then blotted onto a nitrocellulose membrane (Shanghai Genepharma). A *WT1* polyclonal antibody (1:500, Invitrogen) and a polyclonal beta-actin antibody (1:10,000 Invitrogen) were used for primary Western blot reaction, and horseradish peroxidase-conjugated secondary antibodies (Invitrogen) were used for secondary Western blot reaction. Finally, the blots were visualized using a Fusion Solo chemiluminescence system (PEQLAB Biotechnology GmbH, Germany) according to the manufacturer's recommendation.

### WT1 overexpression assay

The whole sequence of human *WT1* gene was inserted into a mammalian overexpression vector, pcDNA/3.1+, to generate a *WT1* overexpression plasmid, pc/WT1 (Shanghai Genepharma). An empty pc/DNA/3.1+ vector was applied as control plasmid, pc/C (Shanghai Genepharma). In LN-18 and A172 cells with hsa-miR-1270 overexpression, they were secondarily transfected with pc/C or pc/WT1 for 72 hours. After that, healthy cell colonies were selected and passaged for three times. Then, qRT-PCR was conducted to examine WT1 expression in GBM cells.

### Statistical analyses

In this study, all statistical analyses were conducted using a SPSS software (SPSS, version 16.0, IBM Corporation, Armonk, NY, USA). Data were averaged from at least three biologic repeats, and demonstrated as means  $\pm$  SEM. A one-way ANOVA, along with Turkey's post hoc test, was conducted across this study. A *P*-value  $<0.05$  was considered as statistically significant.

### Ethical statement

The approval to conduct this study was granted by the Human Research and Ethical Committees at all participating institutes, including The Second Affiliated Hospital of Chongqing Medical University and Chongqing Medical University, both in Chongqing City, China. Consent forms were signed by all participating patients. In addition, all protocols were conducted in accordance with the principles of the Declaration of Helsinki. All animal experiments were approved by the Ethical Committees prior to the commencement of

the study, and conducted according to the institutional and national guidelines and regulations.

## Results

### Hsa-miR-1270 is downregulated in both GBM cell lines and clinical tumors

We examined hsa-miR-1270 gene expression in GBM cell lines. Through qRT-PCR analysis, we found that hsa-miR-1270 was significantly downregulated in LN-18, N-229, U-138MG, T98G, and A172 cells, compared with a NHA cell line (Figure 1A,  $*P < 0.05$ ). In addition, in eleven patients diagnosed with GBM, qRT-PCR showed that hsa-miR-1270 was significantly downregulated in GBM tissues, compared with paired normal tissues, in nine patients (Figure 1B,  $*P < 0.05$ ).

### Hsa-miR-1270 upregulation suppressed GBM in vitro functions of proliferation and migration

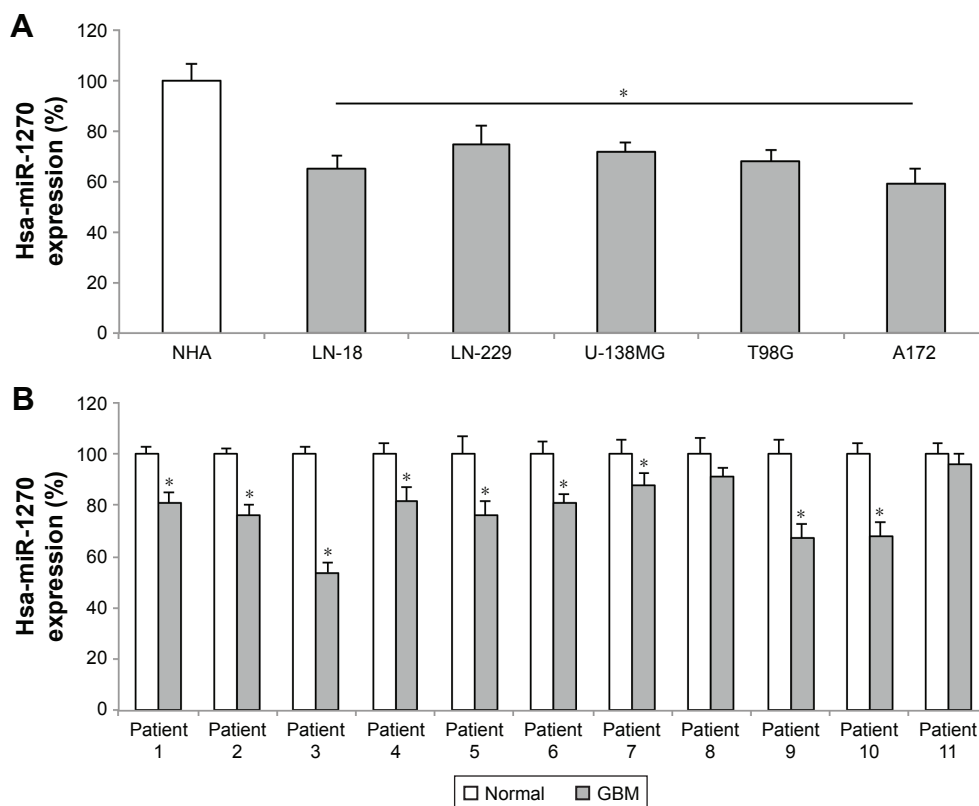
To understand the biologic functions of hsa-miR-1270 in GBM, we transduced LN-18 and A172 cells with lentivirus of miR-1270M. Analysis of qRT-PCR confirmed that,

in miR-1270M-transduced GBM cells, their endogenous hsa-miR-1270 expressions were significantly upregulated, compared with miR-MC-transduced GBM cells (Figure 2A,  $*P < 0.05$ ).

Lentiviral-transduced LN-18 and A172 cells were examined by an in vitro proliferation assay for five consecutive days. It demonstrated that in vitro cancer cell growth was significantly suppressed in miR-1270M-transduced cells compared with miR-MC-transduced cells (Figure 2B,  $*P < 0.05$ ).

Then, lentiviral-transduced LN-18 and A172 cells were examined by an in vitro migration assay for 24 hours. GBM cells migrated into the lower chambers of Transwell assay were stained by crystal violet. The imaging results showed that significantly less miR-1270M-transduced cells were observed than miR-MC-transduced cells (Figure 2C). In addition, quantification on cancer migration demonstrated that hsa-miR-1270 upregulation markedly inhibited GBM migrating capability in vitro (Figure 2D,  $*P < 0.05$ ).

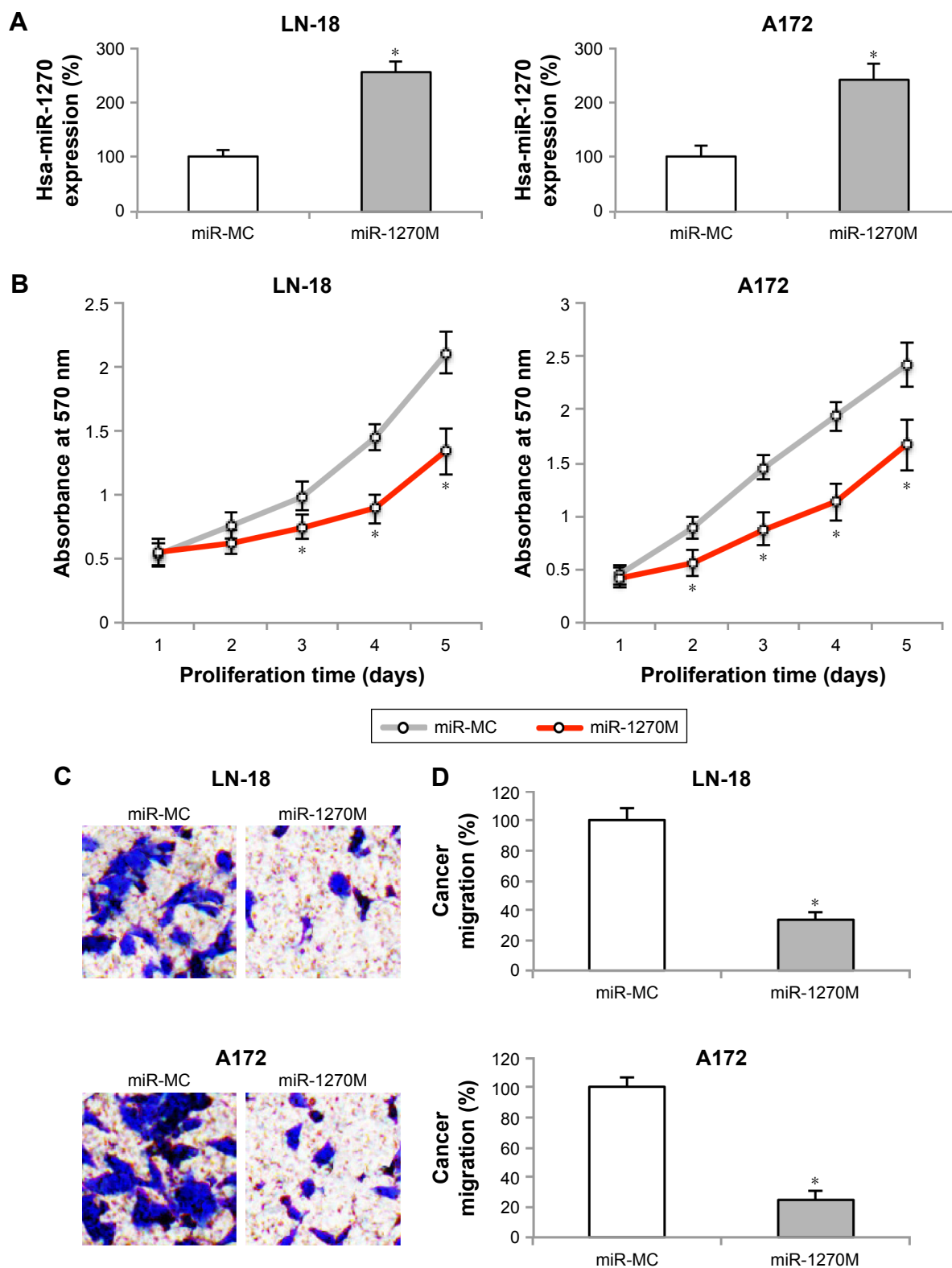
Thus, our data clearly demonstrated that hsa-miR-1270 upregulation suppressed GBM in vitro functions of proliferation and migration.



**Figure 1** Gene expression of hsa-miR-1270 in GBM.

**Notes:** (A) In five human GBM cell lines, LN-18, N-229, U-138MG, T98G, and A172 cells, their endogenous hsa-miR-1270 expressions were measured by qRT-PCR, and compared to endogenous hsa-miR-1270 expression in a normal human astrocyte cell line ( $*P < 0.05$ ). (B) In situ GBM tissues, and paired normal brain tissues, were collected from eleven patients. Their endogenous hsa-miR-1270 expressions were compared by qRT-PCR ( $*P < 0.05$ ).

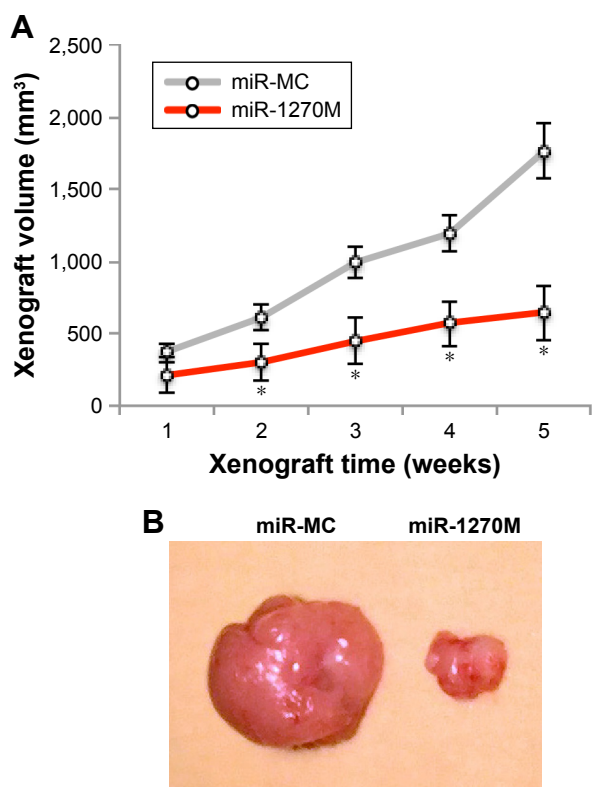
**Abbreviations:** GBM, glioblastoma multiforme; qRT-PCR, quantitative real-time PCR.



**Figure 2** Effects of hsa-miR-1270 upregulation on GBM proliferation and migration in vitro.

**Notes:** (A) LN-18 and A172 cells were transduced with a lentivirus of hsa-miR-1270 mimics, miR-1270M, or a lentivirus of nonspecific human miRNA mimics, miR-MC. After transduction was stabilized, we applied quantitative real-time PCR to examine endogenous hsa-miR-1270 expressions in lentiviral-transduced LN-18 and A172 cells (\* $P < 0.05$ ). (B) In an in vitro proliferation assay, lentiviral-transduced LN-18 and A172 cells were allowed to grow for five consecutive days. Their proliferation rates were measured and compared at absorbance of 570 nm (\* $P < 0.05$ ). (C) In an in vitro migration assay, lentiviral-transduced LN-18 and A172 cells were given 24 hours to migrate in a Transwell plate. GBM cells attached to the bottom of Transwell plate were stained by 0.1% crystal violet and imaged. (D) For imaging result in (C), cancer migration was compared between miR-MC- and miR-1270M-transduced GBM cells (\* $P < 0.05$ ).

**Abbreviation:** GBM, glioblastoma multiforme.



**Figure 3** Effect of hsa-miR-1270 upregulation on GBM xenograft in vivo. **Notes:** (A) Lentiviral-transduced LN-18 cells were subcutaneously xenografted into athymic nude mice for 5 weeks. Twelve mice received xenografts transduced with miR-MC lentivirus. Fourteen mice received xenografts transduced with miR-1270M lentivirus. The volumes of LN-18 xenografts were compared, weekly, between miR-MC- and miR-1270M-transduced LN-18 xenografts ( $*P<0.05$ ). (B) After in vivo xenograft assay, subcutaneous LN-18 tumors were removed and imaged. **Abbreviation:** GBM, glioblastoma multiforme.

## Hsa-miR-1270 upregulation suppressed GBM in vivo xenograft

GBM LN-18 cells, after lentiviral transduction, were xenografted into the subcutaneous compartment of male athymic nude mice. The xenografts were allowed to grow for 5 weeks in vivo. Their subcutaneous tumor volumes were estimated on weekly basis. It showed that LN-18 xenografts transduced with miR-1270M grew much more slowly than those transduced with miR-MC (Figure 3A,  $*P<0.05$ ). After 5 weeks, subcutaneous GBM xenografts were removed from mice, and imaged. It demonstrated that hsa-miR-1270 upregulation significantly suppressed GBM in vivo xenograft (Figure 3B).

## WT1 was regulated by hsa-miR-1270 in GBM

Next, we asked whether other signaling pathways were correlated with hsa-miR-1270 to regulate GBM functions. By studying several online miRNA targeting algorithms (TargetScan,

[www.targetscan.org](http://www.targetscan.org) and miRDB, [www.mirdb.org](http://www.mirdb.org)), we found that human *WT1* was a possible downstream molecular target of hsa-miR-1270 with two potential DNA binding sites on its 3'-UTR (Figure 4A). Then, through the examination of a dual-luciferase reporter assay, we confirmed that hsa-miR-1270 did physically bind human *WT1* gene (Figure 4B,  $*P<0.05$ ). In addition, analysis of qRT-PCR showed that, in lentiviral-transduced LN-18 and A172 cells, *WT1* gene expressions were significantly downregulated by forced hsa-miR-1270 upregulation (Figure 4C,  $*P<0.05$ ). Moreover, Western blot analysis showed that, in lentiviral-transduced LN-18 and A172 cells, WT1 protein expressions were also significantly reduced by forced hsa-miR-1270 upregulation (Figure 4D).

Thus, our data clearly showed that *WT1* was regulated by hsa-miR-1270 in GBM.

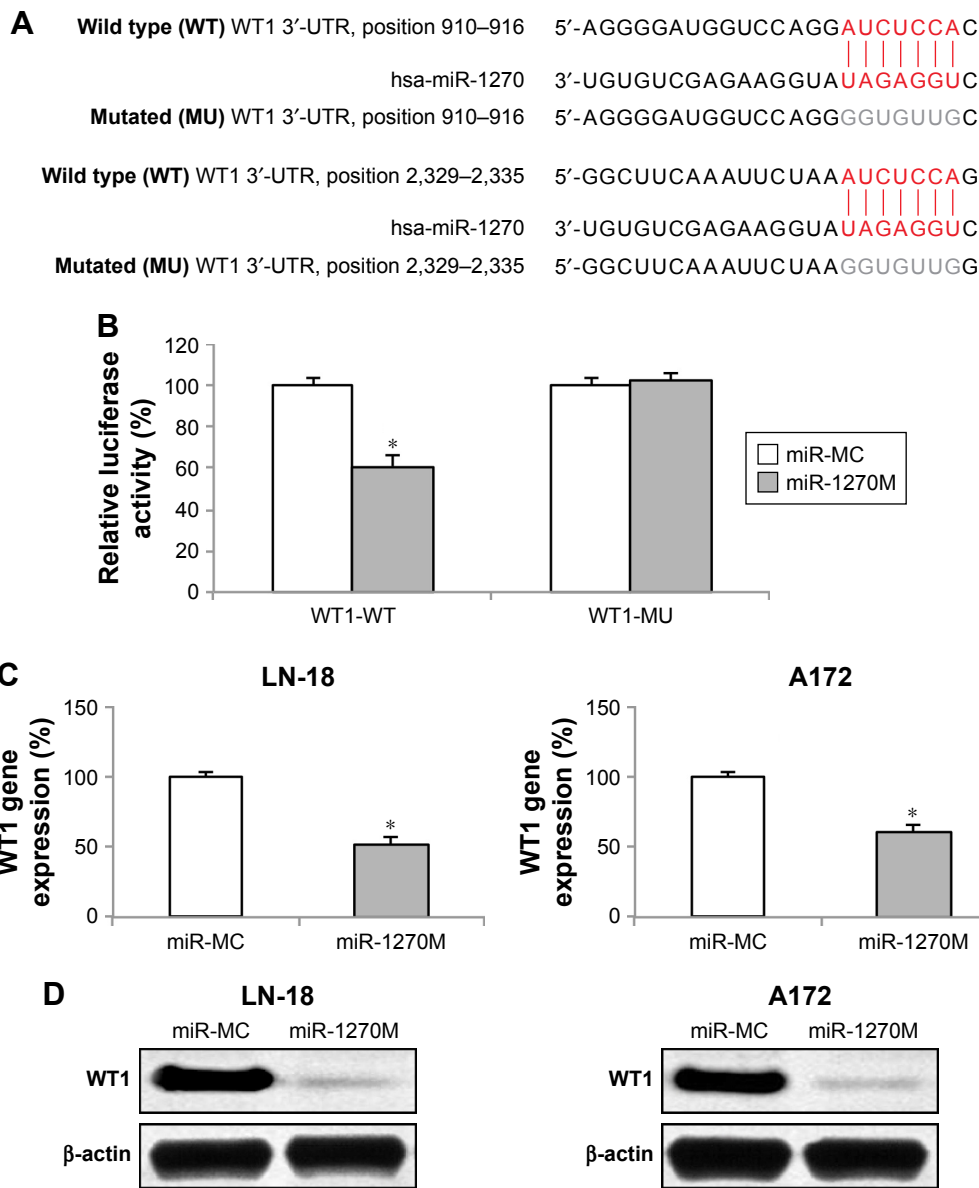
## *WT1* upregulation reversed hsa-miR-1270-induced suppression on GBM in vitro functions

Finally, in hsa-miR-1270-upregulated LN-18 and A172 cells, we transduced them with a mammalian overexpression plasmid, pc/WT1. Analysis of qRT-PCR showed that, in cells transfected with pc/WT1, their endogenous *WT1* expressions were strongly upregulated, as compared to cells transfected with a control overexpression plasmid, pc/C (Figure 5A,  $*P<0.05$ ).

The mechanistic implications of *WT1* upregulation on GBM in vitro functions were explored. First, secondary-transfected LN-18 and A172 cells were examined by the in vitro proliferation assay for five consecutive days. It showed that, in hsa-miR-1270-upregulated GBM cells, in vitro cancer cell growth was significantly augmented in pc/WT1-transfected cells compared with pc/C-transfected cells (Figure 5B,  $*P<0.05$ ).

Then, secondary-transfected LN-18 and A172 cells were examined by an in vitro migration assay. Twenty four hours later, the imaging results showed that, in hsa-miR-1270-upregulated GBM cells, significantly more pc/WT1-transfected cells were observed than pc/C-transfected cells (Figure 5C). In addition, quantification on cancer migration demonstrated that, in hsa-miR-1270-upregulated GBM cells, *WT1* upregulation significantly increased GBM migrating capability in vitro (Figure 5D,  $*P<0.05$ ).

Thus, our data clearly showed that *WT1* upregulation reversed hsa-miR-1270-induced suppression on GBM in vitro functions.



**Figure 4** Inverse correlation of hsa-miR-1270 with human *WT1* gene.

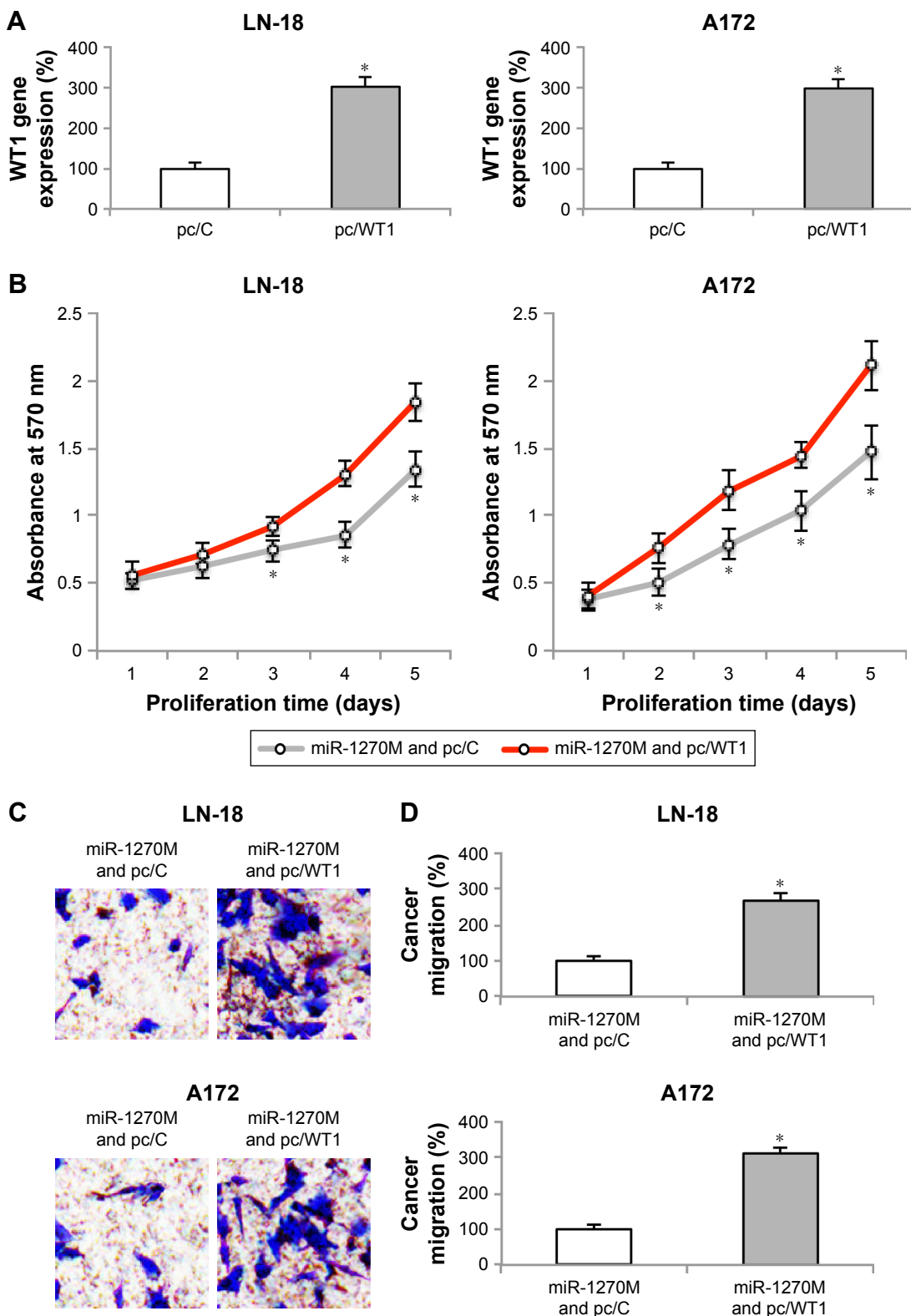
**Notes:** (A) Diagrams are shown for the binding of hsa-miR-1270 on two DNA sites on 3'-UTR of human WT1. The sites were also MU to void hsa-miR-1270 binding. (B) HEK293T cells were transduced with lentiviruses of miR-MC or miR-1270M. Then, they were secondarily transfected with luciferase plasmids of WT1-WT or WT1-MU, followed by a dual-luciferase reporter assay (\* $P < 0.05$ ). (C) In lentiviral-transduced LN-18 and A172 cells, we applied quantitative real-time PCR to examine endogenous *WT1* expressions (\* $P < 0.05$ ). (D) In lentiviral-transduced LN-18 and A172 cells, we applied Western blot to examine *WT1* protein expressions.

**Abbreviations:** MU, mutated; WT, wild type; *WT1*, Wilms' tumor gene.

## Discussion

It is well documented that miRNAs are aberrantly expressed, and play important roles, such as prognostic biomarker and functional regulators in human glioma or GBM.<sup>16–18</sup> A very recent report showed that hsa-miR-1270 was particularly downregulated in grade IV astrocytoma or GBM.<sup>25</sup> Thus, in this study, we first probed the gene expression of hsa-miR-1270 in both GBM cell lines and GBM tumors. We found that hsa-miR-1270 was universally

downregulated in examined GBM cell lines. In addition, in nine of eleven paired clinical samples of GBM patients, hsa-miR-1270 was found to be downregulated in GBM tumors compared with adjacent normal tissues. Interestingly, in the other two paired clinical samples, although it was not significant, hsa-miR-1270 expression levels did seem to be lower in GBM tumors than in adjacent normal tissues (Figure 1B). Thus, the meaning of our results could be twofold. First, hsa-miR-1270 may be predominantly



**Figure 5** Effects of *WT1* upregulation on hsa-miR-1270-correlated GBM proliferation and migration in vitro.

**Notes:** (A) In LN-18 and A172 cells with stable hsa-miR-1270 upregulation, they were secondary transfected with pc/WT1 or pc/C. After transduction was stabilized, we applied quantitative real-time PCR to examine endogenous *WT1* expressions in secondarily transfected LN-18 and A172 cells (\* $P < 0.05$ ). (B) In an in vitro proliferation assay, secondary-transfected LN-18 and A172 cells were allowed to grow for five consecutive days. Their proliferation rates were measured and compared at an absorbance of 570 nm (\* $P < 0.05$ ). (C) In an in vitro migration assay, secondary-transfected LN-18 and A172 cells were given 24 hours to migrate in a Transwell plate. GBM cells attached to the bottom of Transwell plate were stained by 0.1% crystal violet and imaged. (D) For imaging result in (C), in hsa-miR-1270-upregulated LN-18 and A172 cells, cancer migration was compared between pc/WT1- and pc/C-transfected cells (\* $P < 0.05$ ).

**Abbreviations:** GBM, glioblastoma multiforme; WT, wild type; *WT1*, Wilms' tumor gene.



downregulated in both immortal GBM cell lines and GBM tumors, which makes it a potential candidate to be used as a prognostic biomarker in GBM diagnosis. Second, the exact hsa-miR-1270 expression profile in GBM patients could be heterogeneous. There could be subtypes of GBM tumors with less-downregulated hsa-miR-1270 expression levels. Or, hsa-miR-1270 may not be downregulated, or even upregulated in GBM tumors. Therefore, more clinical examinations with a large size of patient pool would be necessary to elucidate the exact hsa-miR-1270 expression profile in GBM.

Second, we applied lentiviral transduction to create two GBM cell lines, LN-18 and A172 cells, with stable hsa-miR-1270 overexpression. Then, by comparing them with GBM cells without hsa-miR-1270 overexpression, we were able to discover that hsa-miR-1270 upregulation had significant antitumor effect in GBM by suppressing cancer cell proliferation and migration in vitro, as well as tumor growth of xenograft in vivo. In several previous cancer studies regarding miR-1270, researchers were able to identify its aberrant expression profile among human cancers, but failed to reveal its functional mechanism.<sup>19,21</sup> Only in a very recent report, Zhong et al showed that inhibiting miR-1270 could yield anticancer mechanisms in osteosarcoma.<sup>20</sup> Thus, the results of our study were among the very first groups to identify functional mechanisms of hsa-miR-1270 in human cancers. Specifically, our data suggested that upregulating hsa-miR-1270 might be an effective biologic target for GBM molecular therapy.

Also in our study, we discovered that human *WT1* gene was actively involved in hsa-miR-1270-mediated cancer-suppressing effects in GBM. Through a dual-luciferase reporter assay, qRT-PCR, and Western blot analyses, *WT1* was shown to be directly bound by hsa-miR-1270, and inversely downregulated by hsa-miR-1270 upregulation in GBM cells. Most importantly, we demonstrated that *WT1* overexpression reversed the anticancer effects of hsa-miR-1270 upregulation on GBM proliferation and migration in vitro. In a previous report, *WT1* was shown to be strongly expressed in glial tumors, hinting its direct correlation with glioma proliferation.<sup>27</sup> In another report, immunotherapy utilizing *WT1* peptide vaccine was demonstrated to be very effective among *WT1*-positive GBM patients, further suggesting an oncogenic nature of *WT1* in GBM.<sup>28</sup> While the results of our study were generally in line with those previous reports, our study was the first one to correlate *WT1* with microRNA epigenetic regulation in GBM.

## Conclusion

Overall, our study presented strong evidence showing that hsa-miR-1270 was predominantly downregulated in GBM. Upregulating hsa-miR-1270 had significant tumor-suppressing effects on GBM functions, possibly through inverse regulation on *WT1* gene. Thus, the miR-1270/*WT1* signaling pathway may be a potential therapeutic target for GBM treatment. Moreover, given the fact that GBM contains subsets of GBM stem-like cells, it would be interesting to learn, possibly through future study, any possible effects of miR-1270/*WT1* signaling pathway in neurogenesis or tumorigenesis in human brain.

## Disclosure

The authors report no conflicts of interest in this work.

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