

# Downregulation of miR-575 Inhibits the Tumorigenesis of Gallbladder Cancer via Targeting p27 Kip1

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**Background:** Gallbladder cancer (GBC) is the most common biliary tract malignant cancer worldwide. It has been reported that microRNA-575 (miR-575) was involved in the tumorigenesis of many cancers. However, the role of miR-575 during the progression of GBC remains largely unknown.

**Methods:** The expression of miR-575 in GBC cells was detected by quantitative real-time polymerase chain reaction. The proliferation of GBC cells was examined by CCK-8 assay and Ki-67 staining. Apoptosis of GBC cells was measured by flow cytometry, and cell invasion was tested by transwell assay. Moreover, protein expressions in GBC cells were evaluated using Western blot. The target gene of miR-575 was predicted using Targetscan and miRDB. Finally, xenograft tumor model was established to verify the function of miR-575 in GBC in vivo.

**Results:** Our findings indicated that miR-575 antagonist decreased the proliferation and invasion of GBC cells. In addition, miR-575 antagonist significantly induced apoptosis of GBC cells via inducing G1 arrest. Meanwhile, p27 Kip1 was found to be a direct target of miR-575 with luciferase reporter assay. Moreover, miR-575 antagonist significantly decreased the expressions of CDK1 and cyclin E1 and upregulated the levels of cleaved caspase3 and p27 Kip1 in GBC cells. Finally, miR-575 antagonist notably suppressed GBC tumor growth in vivo.

**Conclusion:** Downregulation of miR-575 significantly inhibited the tumorigenesis of GBC via targeting p27 Kip1. Thus, miR-575 might be a potential novel target for the treatment of GBC.

**Keywords:** miR-575, p27 Kip1, gallbladder cancer

## Introduction

Gallbladder cancer is the most common biliary tract malignant cancer worldwide.<sup>1</sup> Patients with GBC are always been diagnosed at an advanced stage due to the ignorance of early symptoms.<sup>2</sup> Nowadays, surgery or chemotherapy is regarded as the major treatment of GBC, while only 10% patients with GBC have good prognosis.<sup>3</sup> Moreover, large number of patients with GBC have suffered from severe side effects after chemotherapy and surgical operation.<sup>4</sup> Therefore, it is necessary to explore novel therapeutic methods for the treatment of GBC.

MicroRNAs (miRNAs) are endogenous noncoding small RNAs which are generous in body. Aberrant miRNA expression has found to be related with the progression of multiple diseases recently.<sup>5</sup> In addition, it has been confirmed that miRNAs play key roles in the tumorigenesis of malignancy.<sup>6</sup> MiR-575 is one of the miRNAs

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which has been firstly reported in 2009.<sup>7</sup> MiR-575 was involved in the progression of Kawasaki disease.<sup>8</sup> Meanwhile, a previous study has indicated that BH3-like motif-containing protein (BLID) was a direct target of miRNA-575 in the tumorigenesis of non-small cell lung cancer (NSCLC) *in vitro*,<sup>9</sup> indicating its key role during the progression of malignancies. Yao et al revealed that miR-575 together with the other miRNAs were significantly overexpressed in tumor tissues of patients with GC.<sup>7</sup> However, the role of miR-575 during the progression of GBC remains unclear.

Cyclin-dependent kinase (CDK) inhibitor p27 Kip1 was found to be a member of the CIP/KIP family.<sup>10</sup> In addition, p27 Kip1 has been mostly studied for its roles in inhibiting G1 progression and maintaining cell quiescence in response to anti-proliferative signals or terminal differentiation.<sup>11–13</sup> It can be regarded that p27 Kip1 may act as an important regulator in the growth of malignant tumors by regulation of CDK2 and cyclin E1.<sup>14</sup> Besides, downregulation of p27 Kip1 has been regarded as an independent prognostic factor in GBC.<sup>15</sup> In the current study, we aimed to investigate the function of miR-575 during the tumorigenesis of GBC and explore the correction between miR-575 and p27 Kip1.

## Materials and Methods

### Cell Culture

GBC-SD and G415 cell lines were purchased from Cell Bank of the Chinese Academy of Science (Shanghai, China) and RIKEN Cell Engineering Division-Cell Bank (Tokyo, Japan), respectively. All GBC cells were cultured in Dulbecco's Modified Eagle's Medium (DMEM, Thermo Fisher Scientific) with 10% FBS (Thermo Fisher Scientific), 1% penicillin and streptomycin (Thermo Fisher Scientific) at 37°C, 5% CO<sub>2</sub>.

### Quantitative Real-Time Polymerase Chain Reaction (qRT-PCR)

Total RNA was extracted from GBC-SD or G415 cell lines using TRIzol reagent (TaKaRa, Tokyo, Japan) according to the manufacturer's protocol. cDNA was synthesized using the reverse transcription kit (TaKaRa, Ver.3.0). Real-Time qPCRs were performed in triplicate under the following protocol: 2 min at 94°C, followed by 35 cycles (30 s at 94°C and 45 s at 55°C). The primers for miR-575 and U6 were obtained from GeneCreate Biological Engineering Co., Ltd (Wuhan, China). miR-575: forward, 5'-GTCCACCGCAAATGCTTCTA-3' and reverse 5'-CCATCAGTCCCGTCT

TGAAAC-3'. U6: forward, 5'-CTCGCTTCGGCAGCACAT-3' and reverse 5'-AACGCTTCACGAATTTGCGT-3'. The relative fold changes were calculated using the 2<sup>-ΔΔCt</sup> method by the formula: 2<sup>-(sample ΔCt - control ΔCt)</sup>, where ΔCt is the difference between the amplification fluorescent thresholds of the gene of interest and the internal reference gene (U6) used for normalization.

### MiR-575 Transfection

MiR-575 agonist, miR-575 antagonist, or negative control RNAs (all at 10 nM) were transfected into GBC cells using Lipofectamine 2000 (Invitrogen, Carlsbad, CA).<sup>16</sup> MiR-575 agonist, antagonist, and negative control RNAs were purchased from GenePharma (Shanghai, China). The sequences of miR-575 agonist/antagonist were as follows: miR-575 agonist, GAGCCAGUUGGACAGGAGC and miR-575 antagonist, CUCGGUCAACCUGUCCUCG.

### CCK-8 Assay

GBC cells were seeded in 96-well plates (5×10<sup>3</sup> per well) overnight. Then, cells were treated with negative control (NC) or miR-575 antagonist for 0, 24, 48, and 72 h, respectively. Ten microliters of CCK-8 reagent (Beyotime, Shanghai, China) were added to each well and further incubated for 2 h at 37°C. Finally, the absorbance of GBC cells was measured at 450 nm using a microplate reader (Thermo Fisher Scientific).

### Immunofluorescence

GBC-SD cells were seeded in 24-well plates overnight. Then, cells were treated with NC or miR-575 antagonist for 72 h. Next, cells were blocked with 10% goat serum for 30 min at room temperature and then incubated with anti-Ki67 antibody (Abcam, Cambridge, MA, USA; 1:1000) at 4°C overnight, followed by incubation with goat anti-rabbit IgG (Abcam; 1:5000) at 37°C for 1 h. Then, the nuclei were stained with DAPI (Beyotime, Shanghai, China) for 5 min. Finally, cells were observed under a fluorescence microscope (Olympus CX23, Tokyo, Japan).

### Cell Apoptosis Analysis

GBC-SD cells were trypsinized, washed with phosphate-buffered saline and resuspended in Annexin V Binding Buffer, followed by staining with 5 μL FITC and 5 μL propidium (PI) in the system for 15 min. Cells were analyzed using flow cytometer (BD, Franklin Lake, NJ, USA) to test the cell apoptosis rate.

## Cell Cycle Detection

GBC-SD cells were treated with NC or miR-575 antagonist for 72 h, respectively. Then, the cell cycle distribution was detected by flow cytometry (BD) as previously described.<sup>17</sup> Briefly, cell cycle detection was performed using Cycletest Plus DNA Reagent Kit (BD, NJ, USA). GBC-SD cells were harvested by accutase treatment and counted with a hemocytometer. A total of  $5 \times 10^5$  cells were fixed, permeabilized, and stained in accordance with the manufacturers' instructions, and the sample was analyzed by flow cytometry using a FACSCalibur measuring FL2 area versus total counts. The data were analyzed using ModFit (<http://mycyte.org/>) and FlowJo (<http://mycyte.org/>) softwares to generate the percentages of cells in G1, S, and G2 to M phases of the cell cycle.

## Transwell Assay

For cell invasion analysis, transwell assay was performed in this study. The upper chamber is pre-treated with 100  $\mu$ L of Matrigel. GBC-SD cells were seeded into the upper chamber in media with 1% FBS, and the density was adjusted to about  $1.0 \times 10^6$  cells per chamber. DMEM with 10% FBS was added in the lower chamber. After 48 h of incubation at 37°C, the non-invading cells in the upper chamber were removed with a cotton swab. Then, cells in the lower chamber were stained with 0.1% crystal violet and counted at 3 different fields under a microscope (LEICADMLB2, Frankfurt, Germany).

## Western-Blot Detection

Total protein was isolated from tissue or cell lysates by using RIPA buffer and quantified by BCA protein assay kit (Beyotime, Shanghai, China). Proteins were resolved on 10% SDS-

PAGE and transferred to PVDF (Bio-Rad) membranes. After blocking with 5% skim milk, the membranes were incubated with primary antibodies at 4°C overnight. Then, the membranes were incubated with secondary anti-rabbit antibody (Abcam; 1:5000) at room temperature for 1 h. After that, membranes were scanned by using an Odyssey Imaging System and analyzed with Odyssey v2.0 software (LICOR Biosciences, Lincoln, NE, USA). The primary antibodies used in this study: anti-CDK2 (Abcam, Cambridge, MA, USA; 1:1000), anti-cyclin E1 (Abcam; 1:1000), anti-p27 Kip1 (Abcam; 1:1000), anti-cleaved caspase3 (Abcam; 1:1000) and anti- $\beta$ -actin (Abcam; 1:1000).  $\beta$ -actin was used as an internal control.

## Luciferase Reporter Assay

MiR-575 targeted gene prediction was performed using 2 publicly available programs (miRDB and TargetScan). The results were selected for further analysis. Then, the wild type (wt) or mutate type (mt) of p27 3'-UTR was built based on pmiRGLO vector (Promega, Fitchburg, WI, USA) and named as wt-p27 or mt-p27. Cells were co-transfected with synthetic miRNAs and wt or mt-p27 using Lipofectamine 2000. After 48 h of transfection, luciferase activity was measured using dual-luciferase reporter assay system (Promega).

## In vivo Experiments

Eighteen BALB/c nude mice (6–8 weeks old) were purchased from Vital River (Beijing, China). The mice were housed within a dedicated SPF facility (six mice per group). GBC-SD cells were subcutaneously transplanted into mouse according to the previous reference.<sup>18</sup> When the tumor volume reached about 150 mm<sup>3</sup>, vector-control or miR-575 antagonist (50 nM) was injected intra-tumor twice weekly. The tumor volume was measured weekly according to the formula: Length $\times$ Width $\times$ Width/2.<sup>19</sup> At the end of the experiments, mice were sacrificed and the tumors were collected and weighted. All in vivo experiments were performed in accordance with National Institutes of Health guide for the care and use of laboratory animals, following a protocol approved by the Ethics Committees of Clinical Medical College, Jiangsu Vocational College of Medicine (No. 20190516030).

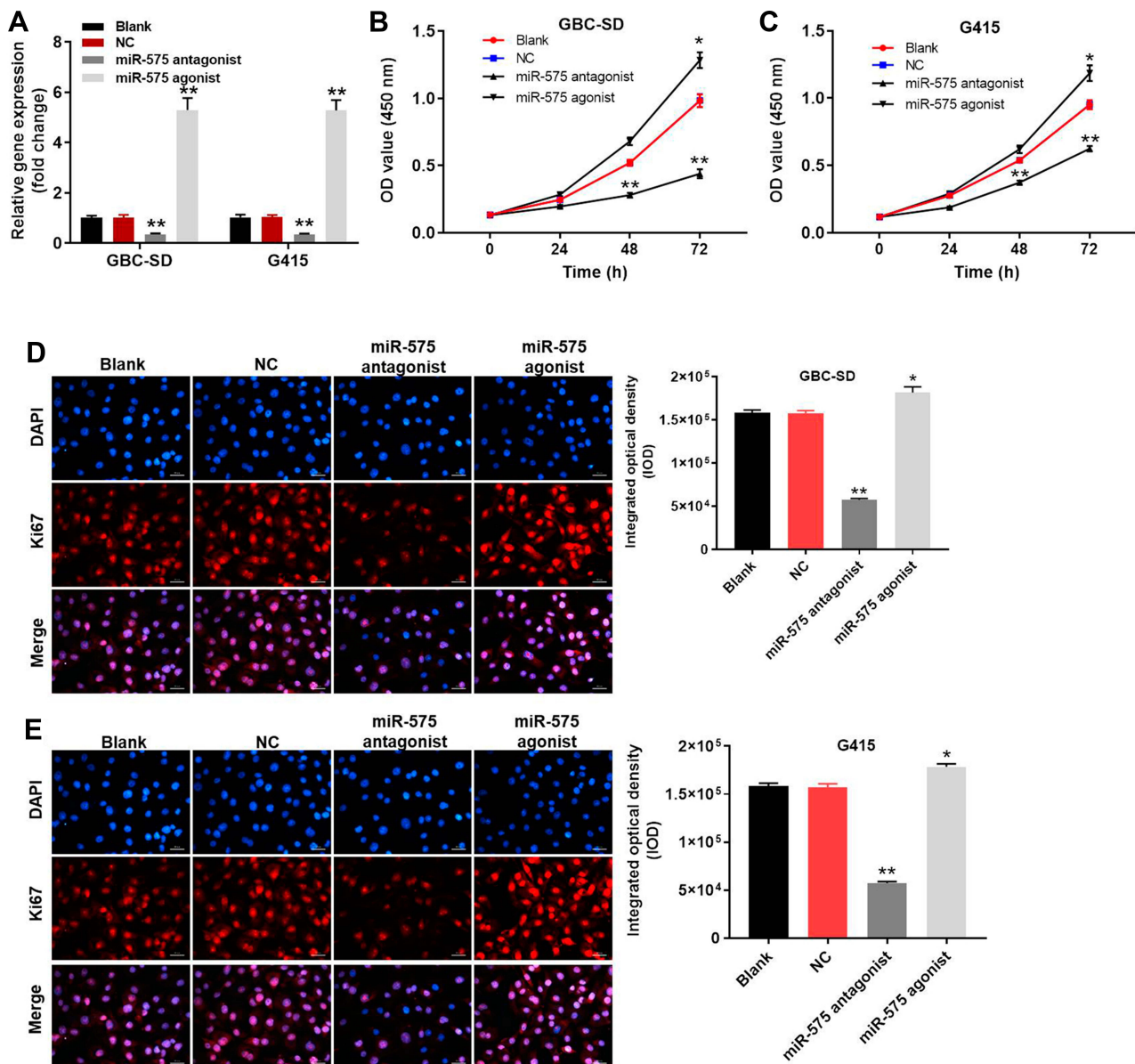
## Statistical Analysis

Each group were performed at least three independent experiments and all data were expressed as the mean  $\pm$  standard deviation (SD). Differences were analyzed using Student's *t*-test (only 2 groups) or one-way analysis of variance (ANOVA) followed by Tukey's test (more than 2 groups, Graphpad Prism7).  $P < 0.05$  was considered to indicate a statistically significant difference.

## Results

### Downregulation of miR-575 Promoted the Proliferation of GBC Cells

To detect the efficacy of transfections, the expressions of miR-575 in GBC cells were measured with qRT-PCR. As indicated in Figure 1A, the level of miR-575 in GBC cells was significantly increased in the presence of miR-575 agonist. In contrast, miR-575 antagonist notably



**Figure 1** Downregulation of miR-575 significantly upregulated the proliferation of GBC cells. GBC-SD or G415 cells were transfected with miR-575 agonist, miR-575 antagonist, negative control (NC) or nothing (Blank) for 24 h. (A) The expression of miR-575 in GBC-SD or G415 cells was detected by using qRT-PCR. (B) After 0, 24, 48, or 72 h of incubation, the OD value of GBC-SD cells was determined using CCK-8 assay. (C) After 0, 24, 48, or 72 h of incubation, the OD value of G415 cells was determined using CCK-8 assay. (D) After 48 h of incubation, the expression of Ki-67 in GBC-SD cells was detected by immunofluorescence staining. The expression of Ki-67 in GBC-SD cells was quantified by integrated optical density (IOD). (E) After 48 h of incubation, the expression of Ki-67 in G415 cells was detected by immunofluorescence staining. The expression of Ki-67 in G415 cells was quantified by integrated optical density (IOD). Each group were performed at least three independent experiments. \* $P < 0.01$  vs control group; \*\* $P < 0.01$  vs control group.

downregulated the expression of miR-575 in GBC cells. This data revealed that GBC cells were stably transfected with miR-575 agonist or antagonist. Next, to explore the role of miR-575 during the tumorigenesis of GBC, CCK-8 assay was performed. As revealed in Figure 1B and C, the OD value of GBC cells was significantly decreased by miR-575 antagonist but increased in the presence of miR-575 agonist. In Ki-67 immunofluorescence assay,

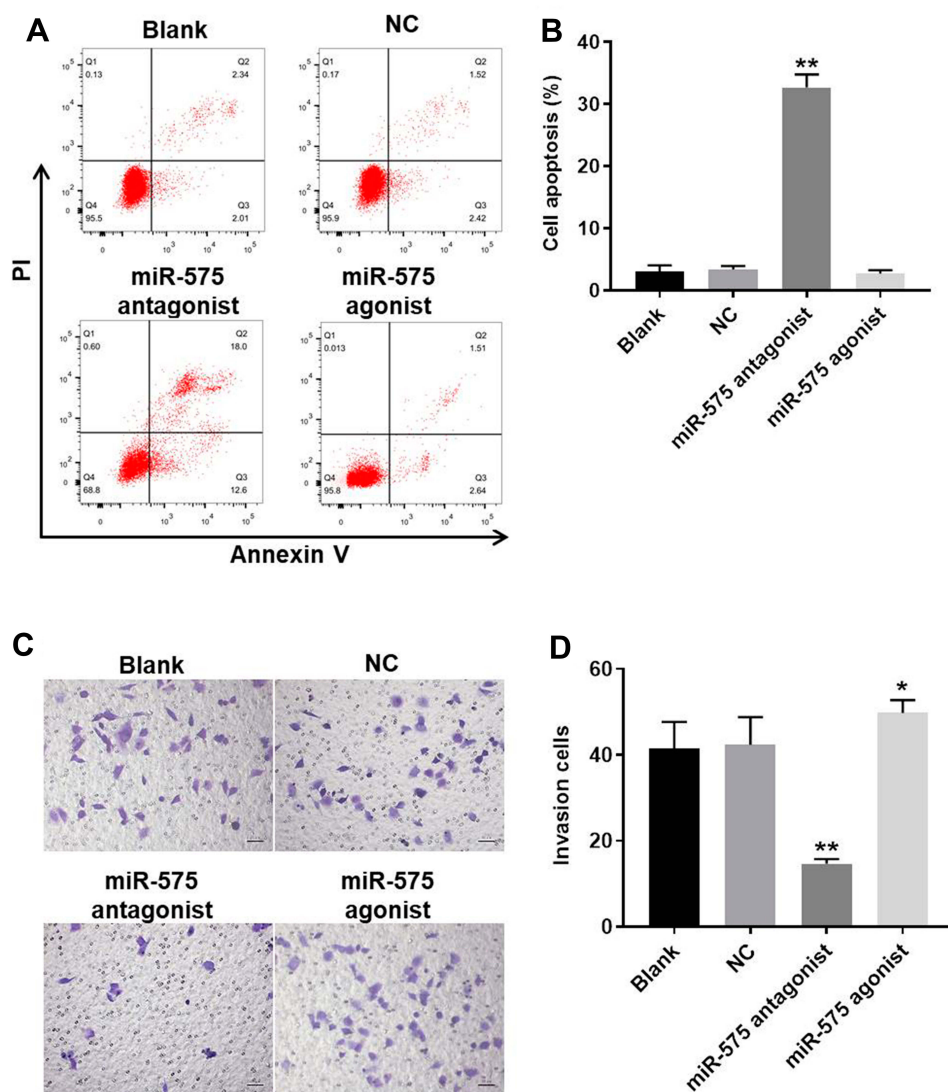
downregulation of miR-575 obviously inhibited the proliferation ability of GBC cells compared with control. However, overexpression of miR-575 significantly promoted the proliferation of GBC cells (Figure 1D and E). Thus, these results revealed that downregulation of miR-575 could inhibit the proliferation of GBC cells, while upregulation of miR-575 exhibited the opposite effect.

## Downregulation of miR-575 Significantly Induced Apoptosis of GBC Cells

Next, to assess the effect of miR-575 on cell apoptosis of GBC-SD, flow cytometry was performed. As showed in [Figure 2A](#) and [B](#), miR-575 antagonist notably induced apoptosis of GBC-SD cells. As expected, upregulation of miR-575 had no effect on cell apoptosis. Consistently, as showed in [Supplementary Figure 1A](#) and [B](#), downregulation of miR-575 notably induced the apoptosis of G415 cells. These data suggested that downregulation of miR-575 could induce apoptosis of GBC cells.

## Downregulation of miR-575 Notably Inhibited the Invasion of GBC Cells

To examine the effect of miR-575 on invasion of GBC cells, transwell was performed. As expected, downregulation of miR-575 significantly inhibited the invasion of GBC-SD cells, while miR-575 agonist promoted cell invasion ([Figure 2C](#) and [D](#)). Similarly, miR-575 antagonist significantly inhibited the invasion of G415 cells ([Supplementary Figure 1C](#) and [D](#)). These data revealed that downregulation of miR-575 significantly downregulated the invasion of GBC cells, while upregulation of miR-575 exhibited the opposite effect in vitro.



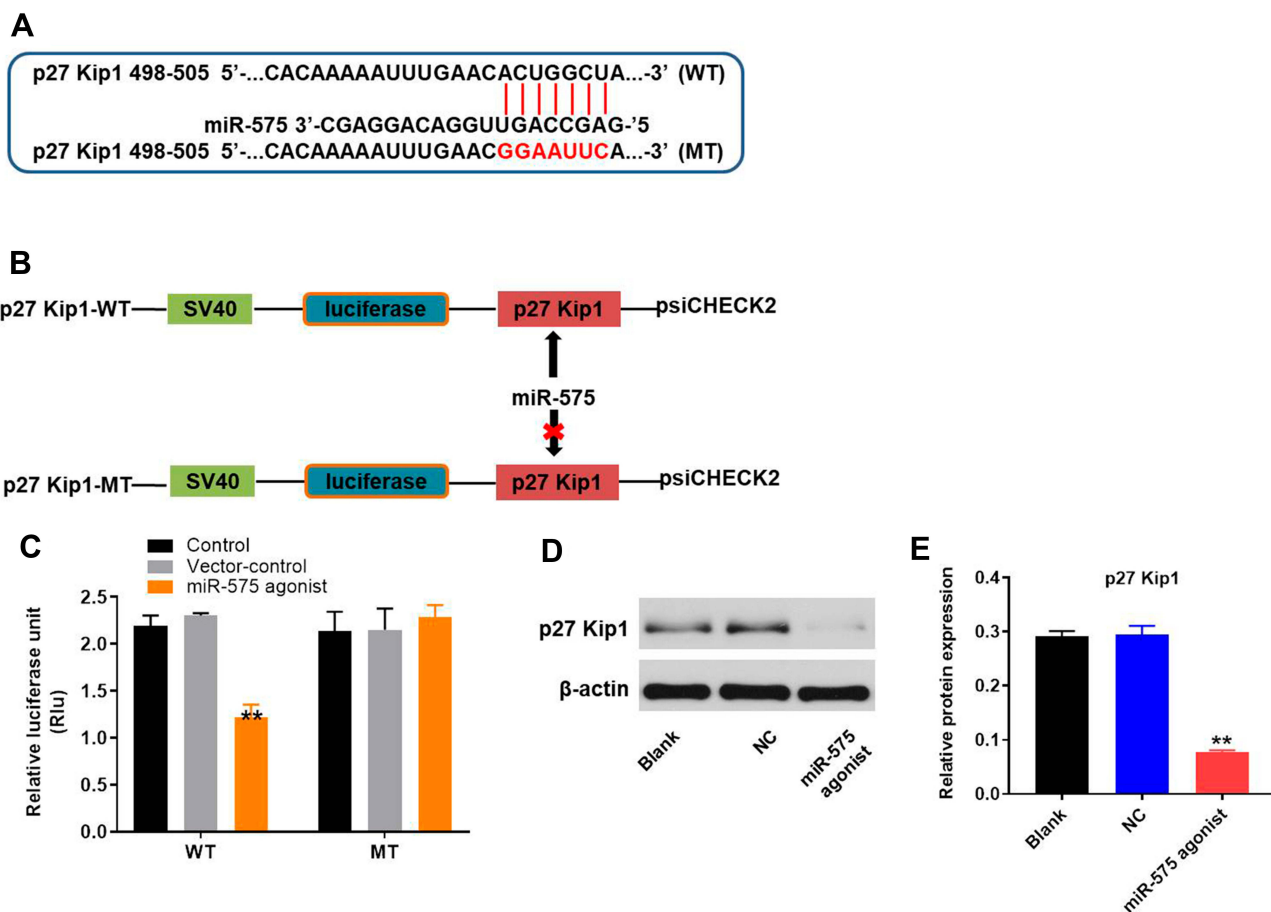
**Figure 2** Downregulation of miR-575 significantly induced the apoptosis and inhibited the invasion in GBC-SD cells. GBC-SD cells were transfected with miR-575 antagonist, miR-575 agonist, or NC for 24 h. Then, cells were incubated for another 48 h. **(A, B)** The rate of apoptotic cells was detected by FACS after double staining with Annexin V and PI. X-axis: the level of Annexin-V FITC fluorescence; Y-axis: the PI fluorescence. **(C, D)** The invasion of GBC-SD cells was detected with transwell assay. Each group were performed at least three independent experiments. \* $P < 0.01$  vs control group; \*\* $P < 0.01$  vs control group.

## p27 Kip1 Was a Direct Binding Target of miR-575

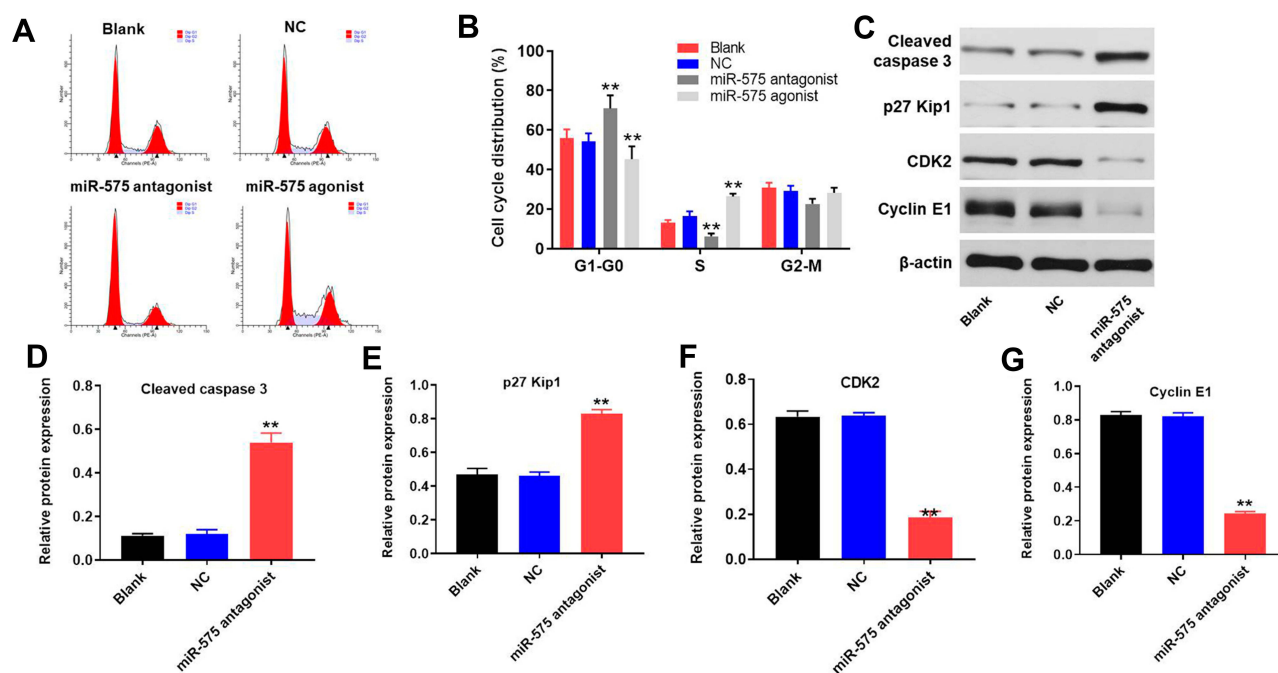
TargetScan and miRDB were used to search target genes of miR-575. The result suggested that p27 Kip1 might be a potential target of miR-575 (Figure 3A and B). In addition, we applied dual-luciferase reporter assay to evaluate whether p27 Kip1 was a binding target for miR-575. The results demonstrated that miR-575 agonist inhibited the luciferase activity of psiCHECK-2-p27 Kip1-WT; however, it did not affect the luciferase activity of psiCHECK-2-p27 Kip1-MT (Figure 3C). Additionally, the result of Western blot showed that the protein expression of p27 Kip1 in GBC-SD cells was notably decreased in the presence of miR-575 agonist (Figure 3D and E). All these data suggested that p27 Kip1 was a direct downstream target of miR-575.

## Downregulation of miR-575 Obviously Induced G1 Arrest in GBC Cells

Next, flow-cytometry was applied to detect the cell cycle distribution in GBC-SD. As illustrated in Figure 4A and B, downregulation of miR-575 notably induced G1 arrest in GBC-SD cells, while miR-575 agonist exhibited the opposite effect. Since miR-575 antagonist exhibited significantly regulatory effect on cell apoptosis and cycle of GBC-SD, apoptosis and cell cycle-related proteins were detected by Western blot. The results indicated that miR-575 antagonist markedly enhanced the expressions of cleaved caspase3 and p27 Kip1 and decreased the expressions of CDK2 and cyclin E1 in GBC cells (Figure 4C–G). Moreover, miR-575 antagonist greatly induced G1-arrest in G415 cells (Supplementary Figure 1E and F). Taken together, downregulation of miR-575 could significantly induce G1 arrest in GBC-SD cells.



**Figure 3** MiR-575 directly targeted p27 Kip1. (A, B) Gene structure of p27 Kip1 at the position of 498–505 indicates the predicted target site of miR-575 in its 3'UTR, with a sequence of GGAAUUC. (C) The luciferase activity was measured after co-transfecting with WT/MT p27 Kip1 3'-UTR plasmid and miR-575 agonist in GBC-SD cells using the dual-luciferase reporter assay. The results were normalized to Renilla luciferase. (D) GBC-SD cells were transfected with miR-575 agonist, NC, or nothing (Blank) for 24 h and the cells were incubated for another 48 h. Protein expression of p27 Kip1 in GBC-SD cells was detected with Western blotting. (E) The relative expression of p27 Kip1 was quantified via normalization to  $\beta$ -actin. Each group were performed at least three independent experiments. \*\* $P < 0.01$  vs control group.



**Figure 4** Downregulation of miR-575 induced G1 arrest in GBC-SD cells. GBC-SD cells were transfected with miR-575 antagonist, miR-575 agonist, nothing, or miR-NC (NC) for 24 h. Then, cells were incubated for another 48 h. After transfection (**A**, **B**) the cell cycle distribution in G0/G1, S, and G2 phase after propidium iodide staining of GBC cells were determined by FACS. (**C**) Expressions of cleaved caspase3, p27 Kip1, CDK2, and cyclin E1 in GBC-SD cells were detected with Western blotting. (**D–G**) The relative expressions of cleaved caspase3, p27 Kip1, CDK2, and cyclin E1 in GBC-SD cells were quantified via normalization to  $\beta$ -actin. Each group were performed at least three independent experiments. \*\* $P < 0.01$  vs control group.

## MiR-575 Antagonist Significantly Inhibited GBC Tumor Growth in vivo

To further evaluate the effect of miR-575 on tumor growth of GBC in vivo, xenograft model was established. As indicated in Figure 5A and B, downregulation of miR-575 markedly inhibited the tumor sizes in mice. In addition, miR-575 antagonist significantly decreased the weight of tumors, compared with control (Figure 5C). Meanwhile, the expression of miR-575 was significantly downregulated in tumor tissues after miR-575 antagonist injection (Figure 5D). All these data revealed that miR-575 antagonist could inhibit tumor growth of GBC in vivo.

## Downregulation of miR-575 Significantly Decreased the Expression of Cell Cycle-Related Proteins in vivo

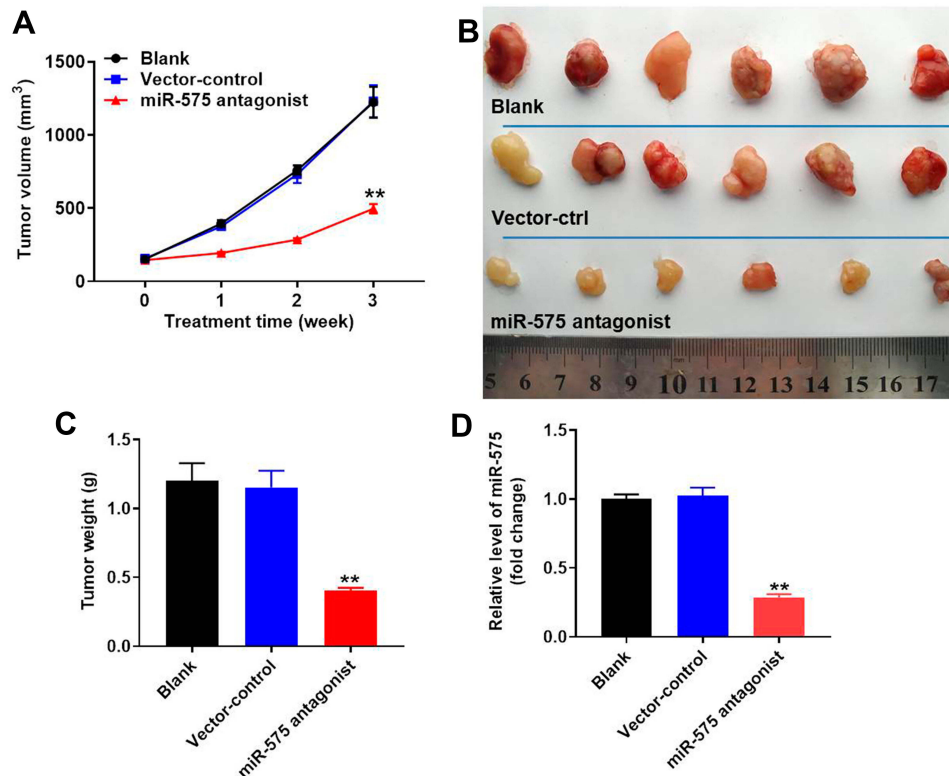
To investigate the effect of miR-575 on the expression of cycle-related proteins in tumor tissues, Western blot assay was performed. As indicated in Figure 6A–D, miR-575 antagonist notably increased the level of p27 Kip1 and decreased the expressions of cyclin E1 and CDK2 in tumor tissues. These data were consistent with in vitro

result and further revealed that miR-575 antagonist inhibited GBC tumor growth via modulating the expression of cell cycle-related proteins.

## Discussion

Recent studies have reported that the miRNA was involved in the growth of GBC, and dysregulation of miRNAs plays a critical role during the tumorigenesis of GBC.<sup>20–22</sup> In this research, we found that the cell growth of GBC was significantly inhibited by miR-575 antagonist. Our study firstly found the effect of miR-575 on GBC, indicating that miR-575 could function as an oncogene in GBC. A previous study found that miR-575 knockdown significantly downregulated the progression of gastric cancer cells.<sup>23</sup> Additionally, miR-575 was notably upregulated in hepatic carcinoma.<sup>24</sup> Our finding was consistent with these data suggesting that miR-575 might act as an oncogene.

It has been confirmed that miRNAs exert their biological functions due to their target genes.<sup>25,26</sup> In this study, luciferase reporter assay indicated that p27 Kip1 was identified as a downstream target of miR-575 in GBC. P27 Kip1 is a cell cycle regulator firstly regarded as a cyclin-dependent kinase antagonist.<sup>27</sup> It has been



**Figure 5** Downregulation of miR-575 inhibited tumor growth of GBC in vivo. GBC-SD cells were subcutaneously injected into nude mice to establish tumor xenograft model. MiR-575 antagonist or vector control was directly injected into the tumors twice a week. **(A)** Tumor volumes of mice were measured weekly. **(B)** At the end of the study, tumor tissues of mice were collected and pictured. **(C)** Tumor weights in each group of mice were calculated. **(D)** The level of p27 Kip1 in tumor tissues was detected using qRT-PCR. Each group were performed at least three independent experiments. \*\* $P < 0.01$  vs control group.

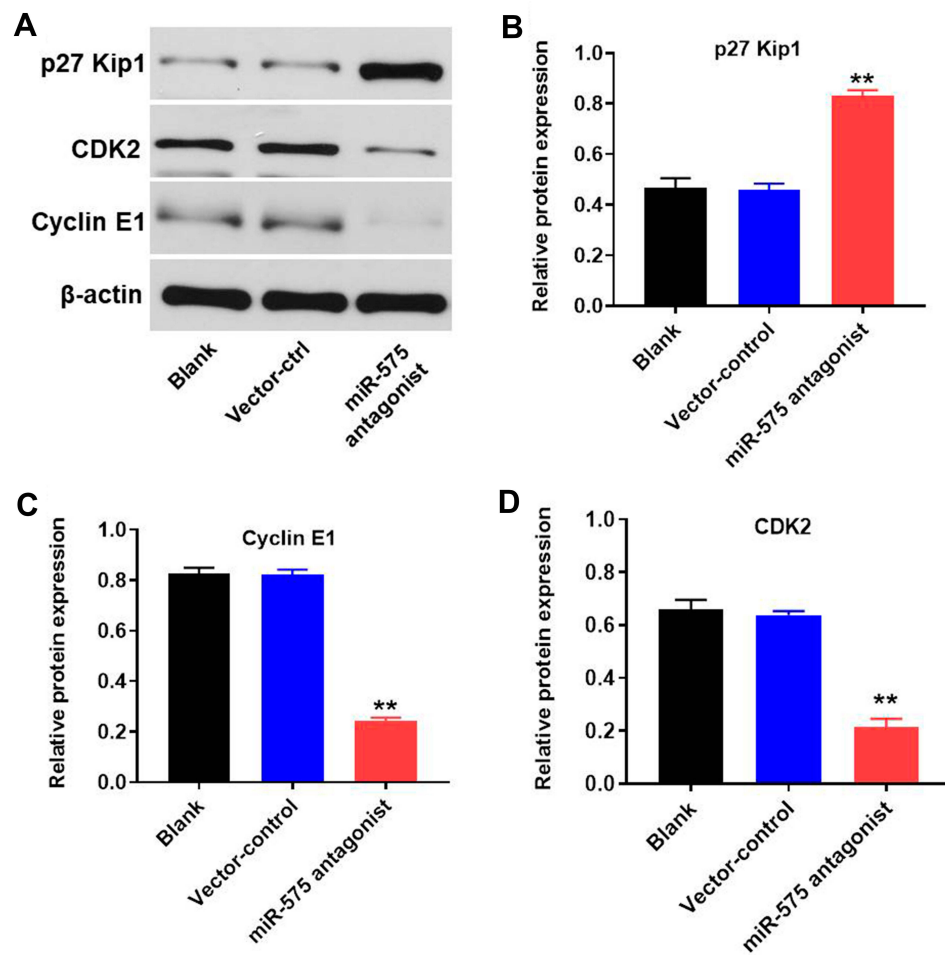
reported that Rooibos decreased the proliferation of castration-resistant prostate cancer cells via upregulation of p27 Kip1.<sup>28</sup> Additionally, Hig-1a inhibited the progression of pancreatic cancer cells through a pERK/p27 Kip1/pRB pathway.<sup>29</sup> In our research, downregulation of miR-575 significantly increased the protein level of p27 Kip1. Our findings were consistent with the previous studies. Collectively, our findings revealed that miR-575 exhibited an oncogenic effect on GBC cells via targeting p27 Kip1. Otherwise, a previous report has suggested that miR-575 could mediate the progression of gastric cancer via targeting PTEN.<sup>23</sup> This difference may be due to the different tumor types.

Additionally, in vitro experiments indicated that downregulation of miR-575 significantly induced G1 arrest in GBC cells via downregulating the expression of CDK2 and cyclin E1. CDK2 and cyclin E1 have been regarded to be cell cycle regulators in malignant tumors.<sup>30–33</sup> Moreover, Liu et al found that NF- $\kappa$ B induced cell cycle arrest in laryngeal squamous cell cancer via downregulation of CDK2.<sup>34</sup> Overexpression

of p38 $\gamma$  could promote the progression of human colorectal cancer via upregulation of cyclin E1.<sup>35</sup> Our research was consistent with these data, revealing that miR-575 antagonist notably induced G1 arrest in GBC cells through inactivating CDK2 and cyclin E1. On the other hand, this study found that downregulation of miR-575 notably activated cleaved caspase 3 in GBC cells. It has been confirmed that cleaved caspase 3 is a pro-apoptotic protein.<sup>36</sup> Our result further confirmed the role of cleaved caspase 3, indicating that miR-575 antagonist induced the apoptosis of GBC cells through upregulation of cleaved caspase 3. Frankly speaking, we only focused on the effect of miR-575 on cell cycle-related proteins so far. Since Lupeol inhibited the tumorigenesis of GBC via suppression of EGFR/MMP-9 signaling pathway,<sup>37</sup> we will further investigate the effect of miR-575 on EGFR/MMP-9 signaling.

Taken together, downregulation of miR-575 significantly suppressed the tumorigenesis of GBC via targeting p27 Kip1, which may be used as a potential target for the treatment of GBC.





**Figure 6** Downregulation of miR-575 inhibited tumor growth of GBC via modulating the expression of cycle-related and pro-apoptosis proteins. **(A)** The protein expressions of p27 Kip1, CDK2, and cyclin E1 in tumor tissues were detected using Western-blot. **(B–D)** The relative expressions of p27 Kip1, CDK2, and cyclin E1 in tumor tissues were quantified via normalization to  $\beta$ -actin. Each group were performed at least three independent experiments. \*\* $P < 0.01$  vs control group.

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## Disclosure

The authors declare no competing interests in this research.

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