miR-299-3p suppresses cell progression and induces apoptosis by downregulating PAX3 in gastric cancer

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Abstract: Gastric cancer (GC) is ranked the fourth leading cause of cancer-related death, with an over 75% mortality rate worldwide. In recent years, miR-299-3p has been identified as a biomarker in multiple cancers, such as acute promyelocytic leukemia, thyroid cancer, and lung cancer. However, the regulatory mechanism of miR-299-3p in GC cell progression is still largely unclear. Cell viability and apoptosis tests were performed by CCK8 and flow cytometry assay, respectively. Transwell assay was recruited to examine cell invasion ability. The interaction between miR-299-3p and PAX3 was determined by the luciferase reporter system. PAX3 protein level was evaluated by western blot assay. The expression of miR-299-3p was down-regulated in GC tissues and cell lines (MKN-45, AGS, and MGC-803) compared with the normal tissues and cells. Besides, overexpression of miR-299-3p significantly suppressed proliferation and invasion and promoted apoptosis in GC. Next, we clarified that PAX3 expression was regulated by miR-299-3p using a luciferase reporter system, qRT-PCR, and western blot assay. Additionally, downregulation of PAX3 repressed GC cell progression. The rescue experiments indicated that restoration of PAX3 reversed miR-299-3p-mediated inhibition on cell proliferation and invasion. miR-299-3p suppresses cell proliferation and invasion as well as induces apoptosis by regulating PAX3 expression in GC, representing desirable biomarkers for GC diagnosis and therapy.

Keywords: miR-299-3p, PAX3, gastric cancer, GC biomarkers, GC therapy

1 Introduction

Gastric cancer (GC) has developed into the fourth leading cause of cancer-related deaths with more than 75% mortality rate worldwide [1,2]. Generally, GC is induced by poor dietary habits, Helicobacter pylori infection, gastric mucosal inflammation, and atrophy [3,4]. Advanced diagnostic tools and medical examinations have facilitated early detection of GC. However, most patients are diagnosed at an advanced stage, which vitiated the treatment outcomes, leading to low 5-year survival rate close to 27% [5–7]. Therefore, exploration of the pathogenesis of GC is essential for the development of novel therapy strategies.

MicroRNAs (miRNAs) refer to a class of small non-coding RNAs comprising 18–23 endogenous oligonucleotides [8]. They specifically participate in tumorigenesis, metabolism, proliferation, differentiation, epithelial-mesenchymal transition (EMT), and metastasis by base-pairing their messenger RNA (mRNA) and resulting in posttranscriptional gene regulation, mRNA degradation, and protein translation suppression [9–11]. Thus, the differential expression of miRNA has been observed in various cancers [12,13]. For instance, the abundance of miR-299-3p markedly accelerated cell growth and G1/S transition in acute promyelocytic leukemia through targeting p21Cip1/Waf1 [14]. On the contrary, miR-299-3p showed a low level of expression in thyroid cancer, and upregulation of miR-299-3p significantly hindered cell progression in vitro and in vivo by regulating SHOC2 expression [15]. Interestingly, miR-299-3p promoted chemosensitivity to doxorubicin by directly targeting ATP binding cassette E1 in lung cancer [16]. However, the biological role of miR-299-3p in GC cell growth remains unknown.
Paired box 3 (PAX3), an essential member of the paired box family gene, is a highly conservative transcriptional factor located at the 35–37 region of the chromosome 2 long arm [17,18]. Typically, PAX3 is involved in tissue development during the embryonic stage and the maintenance of stem cell niches by inhibiting b-Tubulin-III expression [19]. More importantly, PAX3 influences cell self-renewal, migration, and differentiation orientation alteration through multiple pathways [20]. For instance, PAX3 was reported to accelerate human glioma cell proliferation through regulating WNT/β-Catenin signaling pathways [21]. PAX3/FOXO1 fusion accelerated PAX3/FOXO1-positive alveolar rhabdomyosarcoma aggregation by regulating PPP2R1A [22]. Conversely, Wei Liu et al. considered that PAX3 served as a tumor suppressor in thyroid cancer by regulating transcription factor FOXO3a [23]. A recent study shows that the expression of PAX3 might be associated with the prognosis of GC [24]. Another research published in GUT indicated that PAX3 binding to NOC3L affects GC cell growth [25]. These results demonstrate that PAX3 might play a key role in GC progression. However, the precise mechanism of PAX3 in GC is not known. Thus, an investigation of the function of PAX3 in GC is necessary.

In the present study, we explored the function of miR-299-3p during GC cell progression. Examination of miR-299-3p expression by qRT-PCR showed that miR-299-3p was downregulated in GC tumors and cell lines, suggesting the suppressive role of miR-299-3p. Moreover, we demonstrated that PAX3 is a target of miR-299-3p. Besides, miR-299-3p regulates cell progression by targeting PAX3. Our research provides promising targets for GC treatment.

2 Materials and methods

2.1 Patient tissues

Fresh GC tumor tissues and the corresponding normal tissues were collected from 48 GC patients who underwent surgery in Hainan General Hospital. Then, the tissues were transferred and stored at −80°C immediately until use. All the participants have not received preoperative treatment before surgery.

Informed consent: Informed consent has been obtained from all individuals included in this study.

Ethical approval: The research related to human use has been complied with all the relevant national regulations, institutional policies, and in accordance with the tenets of the Helsinki Declaration and has been approved by the Ethics Committee of Hainan General Hospital.

2.2 Cell culture

GC cell lines MKN-45, AGS, and MGC-803 and human gastric mucosal epithelial cell line GES-1 were purchased from Cell Bank of Chinese Academy of Sciences (Shanghai, China). All the cells were maintained in DMEM (Gibco, Grand Island, NY, USA) supplemented with 10% FBS and 0.05% penicillin/streptomycin (Invitrogen, CA, USA) at 37°C in a 5% CO₂ incubator.

2.3 Cell transfection

The miR-299-3p mimics and miRNA negative control (miR-NC) were purchased from RIBOBIO (Guangzhou, China). Small interfering RNA (siRNA) targeting PAX3 (si-PAX3), siRNA negative control (si-NC), and PAX3 were synthesized by GenePharma (Shanghai, China). These plasmids were transfected in MKN-45 and AGS cells using Lipofectamine 2000 (Invitrogen).

2.4 Quantitative reverse transcription polymerase chain reaction (qRT-PCR)

Total RNA was isolated from GC tissues and cells using TRIzol reagent (Invitrogen) by following manufacturers’ instructions. The cDNA for miR-299-3p and PAX3 was synthesized using RNA by All-in-One™ First-Strand cDNA Synthesis Kit (FulGenGen, Guangzhou, China). qRT-PCR was performed using SYBR green (Applied Biosystems, Foster City, CA, USA) according to the standard procedure. The primers for miR-299-3p and PAX3 were as follows: miR-299-3p (Forward, 5’-TTCAGTGTAAACATCCTCGACTG-3’; Reverse, 5’-TGCCAATGGTCTGGAGTGG-3’); PAX3 (Forward, 5’-GTGGGGAATCCCGAGACA-3’; Reverse, 5’-CCTCCTCCTCTTTACCTTT-3’).

2.5 CCK8 assay

CCK8 assay was used to evaluate the cell proliferation ability of GC cells. Briefly, transfected MKN-45 and AGS cells (5,000 cells/well) were seeded onto 96-well plates. After continuous incubation for 24, 48, and 72 h at 37°C in 5% CO₂ incubator, 10 μL of CCK8 reagent (Beyotime, Shanghai,
China) was added to each well for another 2 h. The optical density (OD) value at 450 nm was read by a spectrophotometer (Thermo Fisher Scientific, Waltham, MA, USA).

2.6 Flow cytometric analysis

Transfected MKN-45 and AGS cells were seeded on a 24-well plate and continuously incubated for 48 h. The cells were then collected and stained using Annexin V-FITC/PI Apoptosis Detection Kit (Vazyme, Nanjing, China) for 20 min. The apoptotic rate was counted by BD FACS Canto II (BD Biosciences, Franklin Lakes, NJ, USA) flow cytometry.

2.7 Transwell assay

Cell invasion ability was examined by transwell assay. The upper chamber was coated with Matrigel (Becton Dickinson, Franklin Lakes, NJ, USA) for 4 h. Then, transfected MKN-45 and AGS cells were seeded on the upper chamber coated with Matrigel and continuously incubated for 48 h. Afterward, noninvasive cells were removed from the upper chamber using a PBS-soaked cotton swab. The invasive cells were fixed with 4% paraformaldehyde and stained with 0.1% crystal violet for 10 min, respectively. The visible cells were counted manually under the microscope.

2.8 Dual-luciferase reporter assay

PAX3 sequences harboring wild-type or mutant-type miR-299-3p binding sites were amplified and inserted into the downstream of the stop codon of psiCHECK2 dual-luciferase reporter plasmids (Cat. no C8021; Promega, Madison, WI, USA) and named as PAX3-Wt or PAX3-Mut (Geneseed, Guangzhou, China). Then the wild-type and mutant-type luciferase vectors (PAX3-Wt and PAX3-Mut) were co-transfected with miR-299-3p or miR-NC into MKN-45 and AGS cells using Lipofectamine 2000 transfection reagent. Luciferase activities were evaluated by dual-luciferase assay system (Promega, Madison, WI, USA).

2.9 AGS xenograft model

1×10⁶ AGS cells stable transfected with miR-299-3p were injected into athymic BALB/C mouse (6 weeks old) purchased from Beijing Vital River Laboratory Animal Technology Co., Ltd. (Beijing, China). Then, tumor volume (length × width × width/2) was measured every 5 days. Twenty-eight days later, the mice were sacrificed, and the tumor weight was analyzed. Tumor tissues are snap frozen at −80° for gene expression analysis.

Ethical approval: The research related to animal use has been complied with all the relevant national regulations and institutional policies for the care and use of animals.

2.10 Statistical analysis

All the experiments were conducted at least three times, and data were presented as mean ± standard deviation (SD). Statistical analysis was carried out using SPSS 13.0 software (Chicago, IL, USA) and GraphPad Prism 7 (GraphPad Inc., San Diego, CA, USA). A p-value of <0.05 was considered statistically significant.

3 Results

3.1 miR-299-3p is downregulated in GC tumor tissues and cell lines

The relative expression of miR-299-3p in 48 pairs of GC tumors and normal tissues was measured by qRT-PCR. As illustrated in Figure 1a, there is a variable 2 to 4-fold difference in miR-299-3p expression in GC tumors compared with the corresponding normal tissues. Consistently, miR-299-3p expression was relatively lower in GC cell lines (MKN-45, AGS, and MGC-803) compared with human gastric mucosal epithelial cell lineGES-1 (Figure 1b). From these data, it is speculated that miR-299-3p might play a role of the tumor inhibitor in GC progression.

3.2 Overexpression of miR-299-3p suppresses cell proliferation and invasion and promotes apoptosis in GC

Evaluation of the regulatory effect of miR-299-3p on GC cell proliferation, invasion, and apoptosis was carried out by qRT-PCR, CCK8, flow cytometry, and transwell assay, respectively. The expression of miR-299-3p was elevated
significantly in MKN-45 and AGS cells transfected with miR-299-3p mimics compared with the miR-NC group, indicating that the transfection efficiency was extremely high (Figure 2a and b). CCK8 results revealed that the abundance of miR-299-3p obviously hindered cell proliferation ability (Figure 2c and d). Meanwhile, we noticed that the apoptotic rate was enhanced in MKN-45 and AGS cells transfected with miR-299-3p mimics compared with the miR-NC group (Figure 2e and f). Moreover, upregulation of miR-299-3p suppressed cell invasion in GC (Figure 2g and h).

Figure 1: The expression of miR-299-3p in GC tumors and cell lines. (a) The expression of miR-299-3p in GC tumor tissues compared with the corresponding adjacent normal tissues measured by qRT-PCR. (b) The expression of miR-299-3p in GC cell lines (MKN-45, AGS, and BGC-823) compared with human gastric mucosal epithelial cell line GES-1. *P < 0.05.

Figure 2: miR-299-3p inhibited cell proliferation and invasion and enhanced apoptosis in GC. (a and b) The expression of miR-299-3p in MKN-45 (a) and AGS cells (b) transfected with miR-299-3p and miR-NC. (c and d) Cell viability of MKN-45 (c) and AGS cells (d) transfected with miR-299-3p and miR-NC for 24, 48, and 72 h detected by CCK8 assay. (e and f) The apoptotic rate of MKN-45 (e) and AGS cells (f) 48 h post-transfection evaluated by flow cytometry. (g and h) Transwell assay was utilized to assess cell invasion ability of MKN-45 (g) and AGS cells (h) 48 h post-transfection. *P < 0.05.
Collectively, these results shown that miR-299-3p suppresses proliferation and invasion and facilitates apoptosis in GC.

### 3.3 PAX3 is a target of miR-299-3p

We used online software, StarBase, miRmap, and miRanda, to computationally search target genes on miR-299-3p. Considering that miRNAs bind their target mRNAs via partial base-pairing within the RNA-Induced Silencing Complexes (RISC), and in consequence, promote translational suppression and/or RNA degradation, the predicted genes with GC promotion effect were further screened out. qRT-PCR was conducted to detect the effects of miR-299-3p on the 15 candidate genes' expression. As shown in Figure A1, PAX3 is remarkably downregulated when miR-299-3p was upregulated in GC. Based on bioinformatics prediction by online database StarBase, we found that miR-299-3p could bind to PAX3 3′-UTR (Figure 3a). To validate that,

![Figure 3](image)

**Figure 3**: PAX3 is a downstream target of miR-299-3p in GC. (a) Bioinformatics tool StarBase predicted that PAX3 3′-UTR had the binding sites of miR-299-3p. (b and c) Luciferase activity of MKN-45 (b) and AGS cells (c) co-transfected with PAX3-Wt or PAX3-Mut and miR-299-3p or miR-NC. (d and e) The expression of PAX3 mRNA in MKN-45 (d) and AGS cells (e) transfected with miR-299-3p and miR-NC. (f and g) The expression of PAX3 protein in MKN-45 (f) and AGS cells (g) transfected with miR-299-3p and miR-NC. (h) The expression level of PAX3 was detected by qRT-PCR. (i) The correlation between miR-299-3p and PAX3 levels was measured. *P < 0.05.
we have constructed wild-type PAX3 (PAX3-Wt) and mutant-type PAX3 (PAX3-Mut) vectors, which then were co-transfected with miR-299-3p or miR-NC in MKN-45 and AGS cells to establish a luciferase reporter system. As illustrated in Figure 3b and c, miR-299-3p reduced the fluorescence activity of PAX3-Wt. However, the fluorescence activity of PAX3-Mut was unchanged after miR-299-3p transfection. Moreover, the expression of PAX3 mRNA was decreased in MKN-45 and AGS cells transfected with miR-299-3p (Figure 3d and e). Similarly, upregulation of miR-299-3p repressed PAX3 protein expression (Figure 3f and g). Furthermore, PAX3 was upregulated in tumor tissues compared with normal tissues (Figure 3h), and there was a reverse correlation between miR-299-3p and PAX3 expression levels in GC tissues (Figure 3i). All the data indicated that PAX3 is a downstream target of miR-299-3p in GC.

3.4 PAX3 depletion inhibits cell progression and induces apoptosis in GC

We hypothesized that miR-299-3p exerts its cell regulation function by binding to the target gene PAX3. Thus, we transfected si-PAX3 and si-NC in MKN-45 and AGS cells for the subsequent detection. We observed that PAX3 protein level was significantly lower in MKN-45 and AGS cells after PAX3 knockdown compared with the si-NC group (Figure 4a and b). Moreover, PAX3 silencing significantly attenuated GC cell proliferation at 24, 48, and 72 h post-transfection (Figure 4c and d). Likewise, the number of invasive cells decreased distinctly in si-PAX3 transfection cells compared with si-NC transfection cells (Figure 4g and h). Oppositely, the abundance of PAX3 induced cell apoptosis markedly (Figure 4e and f). Taken together, these results suggest that PAX3 depletion inhibits cell progression and induces apoptosis in GC.

3.5 Restoration of PAX3 attenuated miR-299-3p-induced inhibition on GC cell proliferation and invasion

To explore the regulatory mechanism of miR-299-3p/PAX3 axis in GC cell growth, MKN-45 and AGS cells were transfected with miR-299-3p, miR-299-3p + PAX3, miR-299-3p + vector, and miR-NC. Western blot results exhibited that PAX3 protein was reduced by miR-299-3p, and this effect was reversed by PAX3 plasmid transfection in MKN-45 and AGS cells (Figure 5a and b). Moreover, PAX3 reversed miR-299-3p-induced inhibition on cell proliferation in GC (Figure 5c and d). Besides, low levels of PAX3 accelerated apoptosis, while restoration of PAX3 suppressed apoptosis (Figure 5e and f). Cell invasion ability was inhibited by downregulation of PAX3 expression.

![Figure 4](image_url): PAX3 knockdown repressed proliferation and invasion and induced apoptosis in GC. (a and b) The expression of PAX3 protein in MKN-45 (a) and AGS cells (b) transfected with si-PAX3 and si-NC. (c and d) Cell viability of MKN-45 (c) and AGS cells (d) transfected with si-PAX3 and si-NC for 24, 48, and 72 h. (e and f) The apoptotic rate of MKN-45 (e) and AGS cells (f) at 48 h post-transfection. (g and h) Cell invasion ability of MKN-45 (g) and AGS cells (h) at 48 h post-transfection. *P < 0.05.
However, PAX3 rescued the inhibition of miR-299-3p on cell invasion (Figure 5g and h). These findings clarified that restoration of PAX3 could rescue miR-299-3p-induced inhibition on GC cell proliferation and invasion.

### 3.6 Overexpression of miR-299-3p inhibited the growth of GC in vivo

To further investigate the efficacy of miR-299-3p in GC, we constructed GC tumor xenograft models. As shown in Figure 6a and b, the tumor volume and weight of mice injected with AGS cells stable transfected with miR-299-3p were significantly decreased compared with those of miR-NC groups. Furthermore, in tumors from mice injected with miR-299-3p overexpressing cells, miR-299-3p expression was remarkably upregulated, while PAX became downregulated (Figure 6c and d).

### 4 Discussion

It is well acknowledged that miRNAs play pivotal roles in the physiological and pathological processes of multiple cancers, such as hepatocellular carcinoma, nasopharyngeal...
miR-299-3p suppresses cell progression and induces apoptosis

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Growing evidence has validated that miRNA exerts its function through interaction with the specific target gene. According to bioinformatics prediction by StarBase, PAX3 is a target gene of miR-299-3p. It is well acknowledged that PAX3 is closely associated with embryonic tissue development, disease formation, and tumorigenesis [36–38]. For example, activation of PAX3 controlled muscle precursor cell migration and skeletal muscle formation during forelimb muscle development [39]. PAX3 also contributes to glioblastoma tumorigenesis and differentiation by suppressing p53 transcriptionally [40]. In addition, overexpression of PAX3 was reported to induce cell aggregation and interfere with commissural axon projection in vitro and in vivo during embryonic spinal cord development [41]. PAX3 is often considered an oncogene, although there are contradicting reports emphasizing the growth-inhibiting potential of PAX3. The ectopic expression of PAX3 dramatically inhibited thyroid cancer progression in vitro and in vivo through inhibiting the activity of PI3K/Akt and MAPK signaling pathways and promoting the expression and activity of transcription factor FOXO3 [42]. Overexpression of miR-29 and 206 downregulates cell cycle gene expression and induces cell cycle arrest through stabilization of PAX3 in rhabdomyosarcoma, suggesting a tumor suppressor role for PAX3 [43]. Whether miR-299-3p modulates cell progression by targeting PAX3 is still unclear.

In conclusion, we clarified the biological mechanism of the miR-299-3p/PAX3 axis on GC cell proliferation, invasion, and apoptosis (Figure 7). We found that miR-299-3p was closely associated with GC cell progression, and depletion of miR-299-3p reduced cell progression and enhanced apoptosis in GC. Moreover, the results illustrated that miR-299-3p suppressed proliferation and invasion by targeting PAX3, representing the potential biomarkers for novel targeted drug development.

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Data availability statement: The datasets generated during and/or analyzed during the current study are available from the corresponding author on reasonable request.

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Appendix

Figure A1: The expression levels of target genes of miR-299-3p were measured by qRT-PCR in AGS cell line.