Has-miR-875-5p Inhibits Tumorigenesis by Targeting USF2 via TGF- β Signaling Pathway in Gastric Cancer

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Research

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Abstract

Background: Gastric cancer (GC) is one of the most common malignancies, and more and more evidences show that the pathogenesis is regulated by various miRNAs. In this study, we investigated the role of miR-875 in GC.

Methods: The expression of miR-875-5p was detected in human GC specimens and cell lines by miRNA RT-PCR. The effect of miR-875-5p on GC proliferation was determined by CCK-8 proliferation assay and EDU assay. Migration and invasion were examined by transwell migration and invasion assay and wound healing assay. The interaction between miR-875-5p and its target gene USF2 was verified by a dual luciferase reporter assay. The effects of miR-875-5p *in vivo* were studied in xenograft nude mice models. Related proteins were detected by Western blot.

Results: The results showed that miR-875-5p inhibited the proliferation, migration and invasion of gastric cancer cells *in vitro*, and inhibited tumorigenesis *in vivo*. USF2 proved to be a direct target of miR-875-5p. Knockdown of USF2 partially counteracts the effects of miR-875-5p inhibitors. Overexpression of miR-875-5p can inhibit proliferation, migration, and invasion through the TGF- β signaling pathway by down-regulation of USF2 in GC, providing a new research direction for the diagnosis and targeted therapy of GC.

Conclusions: MiR-875-5p can inhibit the progression of GC by directly targeting USF2 and negatively regulating TGF- β signaling pathway. In the future, miR-875-5p is expected to be used as a potential therapeutic target for GC therapy.

Background

Gastric cancer is one of the most common malignant tumors of the digestive tract due to excessive proliferation of gastric epithelial cells. Worldwide, gastric cancer ranks fourth in morbidity and second in mortality(1). Due to the lack of effective biomarkers for early diagnosis, gastric cancer patients often develop to an advanced stage when they are diagnosed. and their 5-year survival rate is less than 30%(2). Early diagnosis of GC can avoid this deterioration and improve survival in patients with GC. In order to improve the prognosis of GC patients, more effective biomarkers for early diagnosis are needed. Therefore, there is an urgent need to understand the basic mechanism of CG tumorigenesis and development and to find effective biological targets. Although changes in oncogenes and tumor suppressor genes have been reported in gastric cancer, the pathogenesis of gastric cancer, the complex molecular mechanisms and signal transduction pathways involved in the progression and development of gastric cancer have not been fully studied, which limits effective biological targets and clinical treatment(3). Therefore, the exact molecular mechanisms of gastric cancer pathogenesis still need to be fully clarified.

MicroRNAs (miRNAs) are a class of small, highly conserved, non-coding single-stranded RNA molecules encoded by endogenous genes, 18–22 nucleotides in length, that directly target the 3' untranslated region

(3'-UTR) of genes by binding to certain sequence specific sites(4). The silencing complex induced by miRNA prevents the translation or direct degradation of mRNA, resulting in a decrease in the expression of these genes (5). There are various regulations between miRNA and mRNA, affecting epigenetics, RNA stability and translation(6). MiRNA plays an important role in cell proliferation, metastasis, differentiation, apoptosis and development(7). It has been shown that miRNAs are abnormally expressed in cancer tissues compared to normal tissues(8, 9). There are still many unknown details about the role of miRNAs in human cancer and further research is needed.

MiR-875-5p is dysregulated in many diseases, including gestational diabetes mellitus(10), liver fibrosis(11), lung cancer(12), esophageal cancer(13), hepatocellular carcinoma(14), prostate cancer(15), colorectal carcinoma(16) and thyroid cancer(17). These results suggest that miR-875-5p plays a role in the occurrence and development of tumors, in the previous study, we found that the expression of miR-875-5p was decreased in gastric cancer(18), but its role in gastric cancer remains unclear. This study aims to explore the effect of miR-875-5p on gastric cancer and its potential mechanism.

Upstream stimulator (USF) was first identified in HeLa nuclear extract because of its ability to bind to elements in the major late promoter of adenovirus (E-box) and stimulate gene transcription(19, 20). Two USF peptides with apparent molecular weights of 43 and 44 kDa were obtained from HeLa cells, called USF1 and USF2, respectively(21, 22). USF-2 is basic helix-loop-helix (BHLH) transcription factors encoded by heterodimeric and overlapping genes(23). Isoforms produced by splicing show different transcriptional activities in certain promoter environments(24, 25). Although it is widely believed that USF2 expression and relative abundance vary (26), it interact synergistically with other factors in tissue and stimulus-specific transcriptional regulation(27, 28). Studies have shown that USF2 plays a promoting role in the development of breast cancer, colorectal cancer and lung cancer(29–32).

This study aims to explore the biological function of miR-875-5p in gastric cancer and its potential mechanism. We demonstrate for the first time that miR-875-5p directly targets and regulates the 3'-UTR of human USF2 mRNA, which is upregulated in many cancers, including gastric cancer. Here, we report that miR-875-5p is indeed inhibited in primary gastric cancer compared to matched adjacent normal gastric tissues and find that the 3'-UTR of human USF2 mRNA is actually the target of miR-875-5p. Further experiments showed that miR-875-5p inhibited the expression of USF2 at the protein level by directly binding to the 3'untranslated region of USF2, and thus affected the TGF- β signaling pathway, thus inhibiting the proliferation, migration and invasion of gastric cancer cell.

Materials And Methods

Tissue samples

Human GC tissues and paired adjacent normal tissues were collected from 36 patients with GC who underwent radical gastrectomy at the Department of General Surgery, Shandong Provincial Hospital Affiliated Cheeloo College of Medicine of Shandong University, China. After surgical excision, the sample

was quickly frozen in liquid nitrogen for experiments. This study was approved by the ethics committee of Shandong Provincial Hospital. Written informed consent is signed prior to specimen collection.

Cell culture

Six gastric cancer cell lines (BGC-823, HGC-27, MGC-803, SGC-7901, MKN-45, AGS) and GES-1 cell lines were purchased from Institute of Biochemistry and Cell Biology, Chinese Academy of Sciences (Shanghai, China). Cells were cultured in RPMI 1640 supplemented with 10% fetal bovine serum (10% FBS), 100 U/mL penicillin, and 100 mg/mL streptomycin (Gibco, Grand Island, NY, USA) was incubated in moist air at 37°C and 5% CO₂.

Cell transfection

Hsa-miR-875-5p mimics and mimics negative control, hsa-miR-875-5p inhibitor and inhibitor negative control were purchased from Gemma. (Shanghai, China). At least 24 hours before transfection, the cells were cultured in complete medium without antibiotics. Cells were planted in six-well plates, when the cell fusion degree reached 50%-70%, the cells were washed with 1x PBS (PH7.4), 50nM miR-875-5p mimics or miR-mimic NC and 100nM miR-875-5p inhibitors or miR-inhibitor NC were transfected into AGS and MKN-45 cells through Lipofectamine™2000 (Invitrogen). Si-USF2 and si-NC were purchased from Genome (Shanghai, China) and transfected as above.

RNA extraction and RT-qPCR

Total RNA was extracted from GC tissues and cells using Trizol reagent (Takara, Japan) according to the manufacturer's instructions. The cDNA was synthesized using the Evo M-MLV RT Premix for qPCR reagent (AG, China). QRT-PCR was performed on Roche LightCycler 480 II fluorescent quantitative PCR instrument with qPCR SYBR Green Pro Taq HS Master Mix(AG). β -actin was used as the endogenous control. The primers used in this study are as follows: USF2 forward: 5'-AAAGGAGGGATCCTGTCCAA-3', USF2 reverse: 5'-CAGGGCGTTCTCATTCTTCA-3'; β -actin forward: 5'-GCATCGTCACTGGGGAC-3' and β -actin reverse: 5'-ACCTGG CCGTCAGGCAGCTC-3'. In addition, unchained curves were used to evaluate nonspecific amplification. QRT-PCR reaction procedures are as follows: 95°C for 30s, 40 cycles at 95°C 5 s and 60°C 30 s. All procedures were carried out in triplicate and relative expression was calculated by the $2^{-\Delta\Delta CT}$ method.

miRNA RT-PCR

Total RNA was extracted as mentioned above. According to the manufacturer's instructions, using the Mir-X miRNA first-Strand Synthesis Kit (Takara, Japan), RNA (2 μ g) was converted to cDNA. PCR reaction was performed using TB Green® Premix Ex Taq™ II (Takara, Japan), and U6 was used as the endogenous control. The primers of Mir-875-5p and U6 were purchased from RiboBio (RiboBio Co., Ltd, Guangzhou, China), the primer sequences used in this study were as follows; has-miR-875-5p forward: 5'-GCGGGCGGTATACCTCAGTTTTAT-3', reverse 5'-ATCCAGTGCAGGGTCCGAGG-3'; U6 forward: 5'-

CTCGCTTCGGCAGCACA-3'; U6 reverse: 5'-AACGCTTCACGAATTTGCGT-3'. $2^{-\Delta\Delta Ct}$ method was used to calculate the relative expression level. All procedures were also performed in triplicate. Each experiment was performed at least three times.

Protein extraction and Western blot

72h after transfection, the cells were flushed with cold PBS, then RIPA buffer (Beyotime, Shanghai, China) containing PMSF (SolarBio, Beijing, China) and phosphatase inhibitors (SolarBio, Beijing, China) was used for cracking. The protein concentration was calculated using the BCA Protein Assay Kit (Beyotime, Shanghai, China) and the protein was separated by SDS-PAGE using a 10% and 12% polyacrylamide gel (20 µg per sample). Proteins are transferred to the immobilon-NC membrane by electrotransfer. The imblotted membrane was sealed in 5% skimmed milk diluted with TBST, and then incubated with appropriate primary antibodies (anti-USF2, anti-p21, anti-p57, anti-E-cadherin, anti-Vimentin, anti-smad2, anti-phospho-smad2, anti-smad3, anti-phospho-smad3 and anti-GADPH obtained from CST) at 4°C for 12 h. Each experiment was performed at least three times.

Dual-luciferase reporting assay

Dual luciferase assay was used to further verify the targeting relationship of miR-875-5p and USF2. Bioinformatics analysis predicted that the possible binding site of miR-875-5p was the 3' UTR site 595-601 of USF2 mRNA. AGS was grown in 1640 medium containing 10% FBS and transfected with psicheck2-Husf2-3' UTR reporter plasmid and human miR875-5p mimics using Lipofectamine™2000. The dual luciferase reporter gene system was used to detect whether human microRNAs bind and regulate USF2. Each experiment was performed at least three times.

CCK8 assay

Cell growth was measured using cell proliferation reagent CCK-8 (MCE). Cells were inoculated into a 96-well plate (Corning Costar, Corning, NY) at a concentration of 2.0×10^3 per well, as per manufacturer's instructions, and 10 µL CCK8 was added to each well at harvest. One hour after CCK8 was added, cell viability was determined by measuring the absorbance of the transformed dye at 450 nm. Each experiment was performed at least three times.

Edu experiment

The transfected cells were seeded into a 96-well plate at a concentration of 1.0×10^4 /well. The EDU solution (Reagent A) was diluted with cell complete medium in a ratio of 1000:1 to prepare an appropriate amount of 50 µM EDU medium according to the manufacturer's instructions. Add 100 µL 50 µM EDU medium to each well and incubate for 2 hours, then discard the medium; PBS washed the cells twice, 5 minutes each time. Add 50 µL cell fixation solution (PBS containing 4% paraformaldehyde) to each well and incubate at room temperature for 30 min, then discard the fixation solution; Add 50 µL 2 mg/mL glycine to each well, and incubate on decolorizing shaker for 5 min, then discard the glycine solution; Add

100µl PBS to each well, wash with decolorizing shaker for 5min, discard PBS; Add 100µl penetrant (0.5% TritonX-100 PBS) to each well and incubate in decolorizing shaker for 10 min. PBS cleaning once, 5 minutes; Add 1X Apollo® dyeing reaction solution of 100µl to each well, and incubate for 30 min in dark, at room temperature, decolorizing shaker, then discard the dyeing reaction solution. Add 100µl penetrant (0.5% TritonX-100 PBS) to decolorizing shaker, wash it 3 times, 10 min each time, discard penetrant; Add 100µl methanol to each well and clean it twice, 5min each time; PBS cleaning once, 5 minutes each time; Add 100µl 1X Hoechst 33342 reaction solution to each well, and incubate in dark, at room temperature and decolorized shaker for 30 min. Then discard the dyeing reaction solution. Add 100µl PBS to each well and clean 3 times; Add 100µl PBS to each well, save. The results were observed using a high-content imaging system. Each experiment was performed at least three times.

Transwell migration/invasion assay

MKN-45 and AGS cells grew to approximately 70% confluence in RPMI 1640 containing 10% fetal bovine serum and then transfected. After 24 hours, the cells were digested by trypsin and then washed once with PBS. To measure cell migration, culture inserts with an 8 mm aperture (Transwell; CoStar, High Wycombe, UK) were placed in the holes of the 24-well plate and the upper and lower chambers were separated. In the lower chamber, add RPMI 1640 containing 20% FBS, 600µl. Then, serum-free medium containing 5×10^4 cells was added to the upper compartment for migration assay, and 1×10^5 cells were used for matrix gel invasion assay. After incubation at 37°C and 5% CO₂ for 24 h, the cells in the upper compartment were removed with a cotton swab. The cells that invaded the base membrane of the inserts were fixed in 100% methanol for 10 min, stained with 1% crystal violet for 20 min, rinsed in PBS, and examined microscopically. Each experiment was performed at least three times.

Wound healing assay

To evaluate the motility of GC cells. A total of 1×10^6 cells/Wells were inoculated in a 6-well plate and cultured overnight, then transfected with miR-875-5p mimics or NC, miR-875-5p inhibitor or NC and si-USF2 or NC. Eight hours later, the confluent cell monolayers were scraped with a sterile pipette head and the plates were washed twice with PBS and cultured for 24 hours before adding fresh serum-free medium. An image of the plate is taken under a microscope. The clearance size was analyzed with Image J software. Each experiment was performed at least three times.

Tumorigenesis in nude mice

BALB/c nude mice (male, 4-6 weeks old, 16-20 g) were purchased from Beijing Vital River Laboratory Animal Technology Co., Ltd (Beijing, China). All animal experiments were conducted in accordance with the Guidelines for the Care and Use of Laboratory Animals of Provincial Hospital Affiliated to Shandong University. MiR-875-5p NC and miR-875-5p mimics were transfected into MKN-45 cells, and 5×10^5 MKN-45 cells in logarithmic growth phase were suspended in 100µL phosphate buffer, and then seeded subcutaneously into the right axillary of nude mice. The experiments were divided into two groups on average (miR-875-5p NC group and miR-875-5p mimics group, n =6), tumor size was monitored by

measuring length (L) and width (W) with a vernier caliper every 4 days, and volume was calculated using the following formula: $(L \times W^2) / 2$. The mice were fed for 28 days. Tumors were sacrificed and collected. Tumor volume and weight was measured for analysis.

Immunohistochemical staining of xenograft tumors tissue

Tumor sections were incubated overnight with commercial rabbit polyclonal antibodies against USF at 1:50 dilution at 4°C. Then, the slices were diluted with horseradish peroxidase (HRP) antibody (1:100; Invitrogen, Thermo Fisher, US) was conjugated at room temperature for 2 h and then covered with DAB (SP kit (rabbit streptavidin-biotin method detection system), ZSGB-BIO, China). After rinsing the colored plates with water for a period of time, they were soaked in hematoxylin and dyed, then dehydrated and sealed. Subsequently, all fields of view were observed under a light microscope (Tissue FAXS Systems, Austria).

Statistical Analysis

Quantitative values were expressed as the means $\pm$ SD. SPSS 19.0 software (SPSS Inc., Chicago, IL, USA) was used for all statistical analyses. Statistical significance was measured by multiple comparisons using Student's t-test with a significance level of $p < 0.05$.

Results

miR-875-5p is down-regulated in human GC tissues and cell lines

In order to confirm the abnormal expression of miR-875-5p in GC tissues, 36 pairs of GC tissues and adjacent normal tissues were collected and the relative expression of miR-875-5p was examined by miRNA RT-PCR. As shown in Fig.1A, the expression of miR-875-5p in human GC tissues was lower than that in paired adjacent tissues ($P < 0.05$). The expression of miR-875-5p was further examined by miRNA RT-PCR in normal gastric mucosal epithelial cells (GES-1) and GC cell lines (AGS, BGC-823, MKN-45, HGC-27, MGC-803 and SGC-7901). As shown in Fig.1B, the expression of miR-875-5p in GC cell lines was lower than that in GES-1 cells ($P < 0.05$). These data suggest that miR-875-5p is downregulated in GC tissues and cell lines.

miR-875-5p inhibits the proliferation of GC cell lines

To investigate the potential role of miR-875-5p in GC tumorigenesis, miR-875-5p mimics and inhibitors were transfected into AGS and MKN-45 cells. Using CCK8 kit to measure cell growth *in vitro*, the results showed that compared with their negative control, overexpression or silencing of miR-875-5p significantly inhibited or promoted the proliferation of GC cells, respectively ($P < 0.05$, Fig.2A). The EDU experiment further confirmed the role of miR-875-5p in inhibiting GC cells proliferation ($P < 0.05$, Fig.2B). In addition, we also demonstrated that overexpression and silencing of miR-875-5p caused up-regulation or down-

regulation of proliferation-related proteins P21 and P57 ($P < 0.05$, Fig.2C). These results suggested that miR-875-5p could inhibit GC cell proliferation.

miR-875-5p inhibits the migration and invasion of GC cell lines

The inhibitory effect of miR-875-5p in GC cells was further investigated. The effect of miR-875-5p on GC cell migration and invasion was examined by cell migration and matrix invasion assay. As shown in Fig.3, the number of penetrating cells in miR-875-5p mimics group in AGS and MKN-45 cells was also reduced compared to the control group, while silencing of miR-875-5p showed the opposite effect in AGS and MKN-45 cells. In wound healing assay, overexpression of miR-875-5p reduced the migration rate of AGS and MKN-45 cells. In contrast, knockdown of miR-875-5p significantly accelerated AGS and MKN-45 cells migration (Fig.4A). In addition, we also demonstrated that overexpression and silencing of miR-875-5p caused up-regulation or down-regulation of E-cadherin and Vimentin proteins (Fig.4B). These results suggested that miR-875-5p inhibited cell migration and invasion in GC cells.

USF2 is a predicted target of miR-875-5p

The target of miR-875-5p was predicted by TargetScan (http://www.targetscan.org/vert_71/), miRmap (<https://mirmap.ezlab.org/>) and Pictar (<https://pictar.mdc-berlin.de/>). According to the predictions of three miRNA bioinformatics websites, we found that USF2, an oncogene in many malignancies, may be one of the target genes of miR-875-5p.

USF2 is upregulated in human GC tissues

In order to determine the difference in the expression of USF2 in gastric cancer tissues and normal tissue adjacent to cancer, we obtained the expression of USF2 in gastric cancer tissues and normal tissue adjacent to cancer through biological information analysis and database. As shown in the Fig.5A, the expression of USF2 in gastric cancer tissues was higher than that in the corresponding adjacent normal tissues.

The USF2 3'-UTR is a target of miR-875-5p

Dual luciferase assay was used to verify the targeting relationship between miR-875-5p and USF2. Bioinformatics analysis predicted that the possible binding site of miR-875-5p was the 3' UTR site 595-601 of USF2 mRNA (Fig.5B). AGS was grown in 1640 medium containing 10% FBS and transfected with psicheck2-Husf2-3' UTR reporter plasmid and human miR-875-5p simulant or inhibitor using Lipofectamine™ 2000. After 48h of culture, human microRNA was detected by the dual luciferase reporter gene system to determine whether the human microRNA binds and regulates USF2. Compared with the control group, luciferase activity was significantly reduced in AGS cells transfected with miR-875-5p mimics and psicheck2-USF2WT 3'-UTR. Luciferase activity in AGS cells transfected with miR-875-5p inhibitor and psicheck2-USF2WT 3'-UTR was significantly increased. The overexpression and knockdown of miR-875-5p did not affect luciferase activity in cells transfected with psicheck2-usf2mut3' UTR (Fig.5B). After AGS and MKN-45 cells were transfected with miR-875-5p mimics, the mRNA and

protein expression of USF2 were detected by RT-qPCR and Western Blot, and the mRNA and protein expression of USF2 were decreased. The results were reversed after transfection with miR-875-5p inhibitor. In addition, all of these results indicate that the 3'-UTR of USF2 is targeted by miR-875-5p, and point mutations in this sequence cancel this interaction.

USF2 promote proliferation, migration, and invasion in GC cells

In order to detect the potential effect of USF2 on the proliferation, migration and invasion of AGS and MKN-45 cells. AGS and MKN-45 cells were transfected with USF2 siRNA or negative control. CCK-8 and EDU experiments showed that USF2 knockdown significantly inhibited the proliferation of AGS and MKN-45 cells (Fig.S9). We also demonstrated that knockdown of USF2 caused changes of proliferation-associated proteins (Fig.S9C). The wound healing experiment showed that knockdown of USF2 significantly inhibited the migration and invasion of AGS and MKN-45 cells (Fig.S10A). In addition, transwell assay showed that knockdown of USF2 significantly inhibited the migration and invasion of AGS and MKN-45 cells (Fig.S10B). We also demonstrated that knockdown of USF2 caused changes of migration and invasion related proteins (Fig.S10C).

MiR-875-5p inhibits proliferation, migration, and invasion in GC cells by targeting USF2

We demonstrated that miR-875-5p overexpression and USF2 knockdown inhibited proliferation, migration, and invasion in GC cells and inhibited USF2 protein expression through mRNA degradation. The results were opposite when miR-875-5p was silenced. In order to further confirm that miR-875-5p inhibited proliferation, migration and invasion in GC by regulating USF2. MKN-45 and AGS cells were co-transfected with miR-NC + si-NC, miR-875-5p inhibitor + si-NC, and miR-875-5p inhibitor + si-USF2. Through CCK8 assay and EDU assay, we found that down-regulation of USF2 in MKN-45 and AGS cells partially offset the effect of miR-875-5p silencing in proliferation (Fig.6). Proliferation-associated protein was detected by western blot (Fig.6C). Through transwell migration and invasion assay and wound healing assay, we found that down-regulation of USF2 in MKN-45 and AGS cells partially offset the effect of miR-875-5p silencing in migration and invasion (Fig.7). Migration and invasion related protein was detected by western blot (Fig.7C). These findings suggest that miR-875-5p inhibits GC cell proliferation, migration, and invasion by directly targeting USF2.

MiR-875-5p decreases the TGF- β signaling pathway by downregulation of USF2

To examine the mechanisms by which miR-875-5p and USF2 inhibit GC proliferation, migration, and invasion, we investigated whether miR-875-5p mediated these effects by affecting the TGF- β signaling pathway. Western blotting was used to examine USF2 expression levels and downstream proteins Smad2, p-Smad2, Smad3 and p-Smad3. As shown in Fig.6C, the silencing of miR-875-5p showed a significant increase in p-Smad2 and p-Smad3 in GC cells compared to the control group. However, there was no significant difference in Smad2 and Smad3 expression. These effects can be partially counteracted by down-regulating USF2 expression in miR-875-5p inhibitor cells. These results suggest

that USF2 is a downstream functional regulator of miR-875-5p, acting through the TGF- β signaling pathway.

MiR-875-5p suppresses the growth of GC *in vivo*

To demonstrate the effect of miR-875-5p on tumor growth *in vivo*, miR-875-5p-mimics transfected cells were injected into the flanks of nude mice, and miR-NC transfected cells served as negative controls. As shown in Fig.8, compared with control group, the miR-875-5p mimics group showed a significant reduction in tumor volume and weight. In addition, immunohistochemistry was used to investigate the expression levels of USF2. Compared with the control group, USF2 protein levels were significantly reduced in miR-875-5p-mimics group (Fig.8B). In conclusion, miR-875-5p inhibited the growth of GC cell *in vivo*.

Discussion

Abnormal expression of miRNAs plays an important role in the occurrence and development of tumors(33–35). Previous studies have shown that miRNA can bind to specific sites of certain sequences to directly target the 3'-UTR of genes and prevent their translation or direct degradation of mRNA, thus participating in the occurrence and development of tumors(9, 36, 37). As a newly discovered tumor-related miRNA, the role of miR-875-5p in gastric cancer and the specific mechanism of its role are still unclear. In this study, we analyzed the expression level of miR-875-5p in GC tissues and normal tissue adjacent to cancer, gastric cancer cell lines and GES-1 cells, and predicted the target genes. In CCK8 and Edu experiments, the overexpression of miR-875-5p inhibited the proliferation of gastric cancer cells and affected the expression of proliferation-related proteins P21 and P57. The knockdown of miR-875-5p showed opposite results, indicating that miR-875-5p played an inhibitory role in the proliferation of gastric cancer cells. Tumor spread and metastasis are affected by the migration and invasion ability of tumor cells, both of which are inhibited by miR-875-5p *in vitro*. Transwell migration and invasion experiment showed that the overexpression of miR-875-5p inhibited the migration and invasion of gastric cancer cells. In addition, the expression of related proteins E-cadherin and Vimentin was affected, and the results were reversed when miR-875-5p was knocked down. Experiments in nude mice demonstrated that the overexpression of miR-875-5p inhibited the tumorigenicity of gastric cancer cells *in vivo*, and this series of experiments showed that miR-875-5p have an inhibitory effect on gastric cancer, suggesting that miR-875-5p may be a potential therapeutic target for gastric cancer patients. Through a series of experiments, it was found that the overexpression of miR-875-5p inhibited the proliferation, migration, invasion and tumorigenicity of gastric cancer cells, suggesting that miR-875-5p could be a potential therapeutic target for GC patients. In addition, Western blotting and dual luciferase report analysis showed that USF2 is the target gene of miR-875-5p and is negatively correlated with miR-875-5p. Our study also showed that miR-875-5p directly binds to the 3'-UTR region of USF2 to inhibit its expression and down-regulate TGF- β signaling pathway.

To elucidate the mechanism of the effect of miR-875-5p on proliferation, migration and invasion. Bioinformatics analysis predicted the putative target of miR-875-5p in GC cells. Among the candidate target genes, we selected USF2(32). The upstream stimulatory factor(USF2) plays an important role in various cellular processes, especially in the genesis and progression of tumors(38–40). However, the role of USF2 in different tumors is contradictory, suggesting that it is either a tumor promoter or an inhibitor(29–31, 41). The role of USF2 in gastric cancer has not been studied. Therefore, it is very necessary to study its expression and role in gastric cancer. Compared with normal gastric tissue, the expression of USF2 in gastric cancer tissue was significantly increased. In CCK8 experiment and Edu experiment, the OD value and the proportion of proliferating cells in si-USF2 group were significantly decreased, indicating that the proliferation of gastric cancer cells would be inhibited after USF2 knockdown. Transwell migration and invasion experiment showed that the migration and invasion ability of gastric cancer cells would be inhibited after USF2 knockout. Dual luciferase experiments further confirmed that miR-875-5p could directly bind to the USF2 3'-UTR region and degrade USF2 mRNA. In addition, knockdown of USF2 partially offset the promoting effect of miR-875-5p silencing on proliferation, migration, and invasion. In conclusion, our results indicate that the inhibitory effect of miR-875-5p in gastric cancer is mediated by down-regulation of USF2.

Previous studies have shown that TGF- β signaling pathway is associated with tumor proliferation and metastasis(42–45). USF2 has been reported to down-regulate Smurf1 and Smurf2, thereby regulating the TGF- β pathway in breast cancer(30). Therefore, we studied the TGF- β pathway changes induced by miR-875-5p expression in GC. Western Blot results showed that the expression of phosphor-SMAD2 and phosphor-SMAD3 increased after knockdown of miR-875-5p, while the expression of SMAD2 and SMAD3 remained unchanged and TGF- β signaling pathway was activated. On this basis, the expression of phosphor-SMAD2 and phosphor-SMAD3 decreased slightly after knockdown of USF2. The expression of SMAD2 and SMAD3 remained unchanged. These results suggest that the decreased miR-875-5p can activate the TGF- β signaling pathway by upregulation of USF2. These results suggest that the decreased miR-875-5p can activate the TGF- β signaling pathway by upregulating USF2. In other words, miR-875-5p can be used as a negative regulator of TGF- β pathway to inhibit the proliferation, migration and invasion of gastric cancer. However, it is not ruled out that miR-875-5p can also influence other signaling pathways to exert anti-cancer effects.

Conclusion

Our study showed that miR-875-5p was significantly down-regulated in gastric cancer tissues and gastric cancer cell lines. As a key inhibitor of the growth and metastasis of gastric cancer cells, miR-875-5p inhibited the progression of GC by directly targeting USF2 and negatively regulating TGF- β signaling pathway. In the future, miR-875-5p is expected to be used as a potential therapeutic target for GC therapy.

Abbreviations

USF2: upstream stimulate factor2; miRNA: Micro RNA siRNA: Smallinterfering RNA; 3'-UTR: 3'-untranslated region; qRT-PCR: Quantitative real-time PCR; FBS: Fetal bovine serum; EdU: 5-ethynyl-2'-deoxyuridine; CCK-8: CellCounting Kit-8; IHC: Immunohistochemistry; TGF- β :transforming growth factor- β

Declarations

Ethics approval and consent to participate

All institutional and national guidelines for the care and use of laboratoryanimals were followed.

Consent for publication

All authors have agreed with publishing this manuscript.

Availability of data and materials

The datasets used and/or analyzed during the current study are availablefrom the corresponding author on reasonable request.

Competing interests

The authors declare no conflict of interest.

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Author Contributions

XG designed the project; SGcontributed to performin vitro and in vivo experiments and write article; ZZ, XW and YMcontributed to analyze experiment data; CL, HL, CJ and LL provided advice and critical comments.

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Figures

Figure 1

Expression of miR-875-5p in GC tissues, GC cells and transfected cells. (A) The expression levels of miR-875-5p in 28 pairs of human GC tissues and adjacent normal tissues were detected by miRNA RT-PCR. (B) The expression level of miR-875-5p in GC cells and GES-1 was examined by miRNA RT-PCR. (C) The efficacy of transfection was verified in cells transfected with miR-875-mimics and miR-875-inhibitor. * $P < 0.05$. Data expressed as mean $\pm$ SD.

Figure 2

miR-875-5p inhibited the proliferation of GC cells in vitro. (A) CCK8 assay was used to evaluate the effect of miR-875-5p on the proliferation of AGS or MKN-45 cells. (B) EDU assay showed the effect of overexpression or silencing of miR-875-5p on the growth of GC cells. (C) Western blotting showed the levels of proliferation-related proteins (P21 and P57) in miR-875-5p overexpressed and miR-875-5p silenced AGS or MKN-45 cells and their negative control cells.* $P < 0.05$. Data expressed as mean $\pm$ SD.

Figure 3

miR-875-5p inhibited migration and invasion in vitro and induced associated protein changes. The effects of miR-875-5p on up-regulation and down-regulation of migration and invasion ability of GC cells were estimated by Transwell assay.

Figure 4

miR-875-5p inhibited migration and invasion in vitro and induced associated protein changes. (A) The wound healing test was performed to study the changes of miR-875-5p overexpression and silencing on the migration ability of GC cells. The percentage of wound closure or the number of transmembrane cells were calculated and compared. (B) Western blot analysis showed that miR-875-5p overexpression and or silencing increased or decreased E-cadherin and Vimentin protein expression, respectively. * $P < 0.05$. Data expressed as mean $\pm$ SD.

Figure 5

USF2 was up-regulated in GC tissues and confirmed to be a direct target gene of miR-875-5p. (A) Differential expression of USF2 in cancer and normal tissue adjacent to the carcinoma. (B) Luciferase reporter assay was performed to confirm that miR-875-5p directly bound to the 3'-UTR region of USF2. The results of luciferase activity were analyzed in cells treated with miR-875-5p-mimics or negative control and miR-875-5p-inhibitor or negative control. (C) The expression level of USF2 mRNA in AGS and

MKN-45 cells after transfected was detected by qRT-PCR. (D) The expression level of USF2 protein in AGS and MKN-45 cells after transfection was verified by Western blot. * P<0.05. Data expressed as mean ±SD.

Figure 6

The effect of miR-875-5p silencing on GC cells was partially offset by USF2 knockdown. (A) CCK8 assay was performed to determine that USF2 knockdown could partly counteract miR-875-5p silencing-induced proliferation in MKN-45 and AGS cells. and (B) EDU assay was performed to determine that USF2

knockdown could partly counteract miR-875-5p silencing-induced proliferation in MKN-45 and AGS cells. (C) Western blot showed the changes of related proteins such as USF2, P21, P57, Smad2, p-Smad2, Smad3 and p-Smad3 in AGS and MKN-45 cells after transfection. (D) Proposed model of miR-875-5p regulating USF2 and affecting TGF- β signaling pathway. * $P < 0.05$; Data expressed as mean $\pm$ SD.

Figure 7

The effect of miR-875-5p silencing on GC cells was partially offset by USF2 knockdown (A) Changes in cell migration were examined by wound healing assayed in AGS cells and MKN-45 cells. (B) Transwell assay was used to confirm the influence of USF2 alterations on GC cell migration and invasion.(C) Western blot showed the changes of related proteins such as E-Cadherin and Vimentin in AGS and MKN-45 cells after transfection. * $P < 0.05$; Data expressed as mean $\pm$ SD.

Figure 8

miR-875-5p inhibits tumor formation in vivo. (A) xenograft tumors were obtained from different groups of nude mice injected with MKN-45 cells transfected with miR-875-5p NC and miR-875-5p-mimics, respectively. (B) The expression of USF2 in the xenograft tumors was determined by immunohistochemistry. (C) The tumor growth curve and tumor weight were significantly different between miR-875-5p mimics group and miR-875-5p NC group.* $P < 0.05$; Data expressed as mean $\pm$ SD.

Supplementary Files

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