LINC01224 accelerates malignant transformation via MiR-193a-5p/CDK8 axis in gastric cancer

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Abstract

Background: Gastric cancer (GC) is a malignant tumor with a significantly high mortality rate, yet, its pathogenesis is not fully understood. Bioinformatics predicted that LINC01224 is highly expressed in stomach adenocarcinoma (STAD), and showed that LINC01224 adsorbed miR-193a-5p to target CDK8. Therefore, this study intended to verify the effect of the LINC01224/miR-193a-5p/CDK8 axis on the biological behavior of gastric cancer.

Methods: Expressions of LINC01224, miR-193a-5p, CDK8, apoptosis-, and EMT-related genes were analyzed using the GEPIA website, RT-qPCR, in situ hybridization, and Western blot as needed. Bioinformatics and dual luciferase assay were used to evaluate the relationship between LINC01224, miR-193a-5p, and CDK8. Functional experiments and rescue experiments (MTT assay, flow cytometry, wound healing assay, and Transwell) were conducted to detect the effects of the above genes on the biological characteristics of GC cells. Tumorigenesis assay was used to verify the results of in vitro experiments.

Results: LINC01224 adsorbed miR-193a-5p to target and upregulate CDK8. The expressions of LINC01224 and CDK8 were increased, while the expression of miR-193a-5p was decreased in GC. Overexpressed LINC01224 promoted cell viability, migration and invasion, accelerated tumor formation, attenuated apoptosis, inhibited the expressions of apoptosis-related proteins, and promoted the expressions of EMT-related proteins, whereas silenced LINC01224 led to the opposite effect. MiR-193a-5p inhibitor partially offset the effect of silenced LINC01224; interestingly, siCDK8 significantly reversed the effect of miR-193a-5p inhibitor on GC cells.

Conclusion: LINC01224 affects the biological behavior of gastric cancer by mediating miR-193a-5p to regulate CDK8.

Keywords

biological characteristics, cyclin-dependent kinase 8, gastric cancer, LINC01224, miR-193a-5p
1 | INTRODUCTION

Gastric cancer (GC) is a kind of malignant tumor that originates in gastric epithelial cells.1,2 Despite the decline in incidence in the past few decades, GC is still a major problem affecting global health.3 Hence, it is necessary to understand the pathogenesis of GC and make breakthroughs in the diagnosis, treatment, and prognosis of this disease. At present, a large amount of evidence has confirmed that lncRNA plays a vital role in the growth and metastasis of GC, thus revealing the potential role of lncRNA in the clinical treatment of GC and the improvement of GC patient survival.4

Recently, accumulating evidence has shown that abnormal expression of lncRNA is related to the occurrence and metastasis of GC and the prognosis of GC patients.5,6 For instance, Hu et al. found that lncRNA GAPLINC is able to upregulate the expression level of CD44 by binding to miR-311-3p to weaken its inhibitory effect on CD44, thereby regulating the migration path and proliferation process of cells, which indicates that lncRNA GAPLINC acts as a molecular bait of miR-311-3p and a ceRNA of oncogene CD44 in GC.7 LncRNAs can bind to specific miRNAs through base complementation, thereby interfering with the regulation of downstream target genes by miRNAs.8 This mechanism is called the “sponge effect.” Elucidating this new gene interaction mode would provide a new perspective for the study of the mechanism underlying GC tumor formation and the development of antitumor therapy.9,10

LINC01224 is a newly identified lncRNA whose exact biological function and mechanism of action in tumors, especially GC, remains unclear. According to the bioinformatics database, lncRNA LINC01224 is upregulated in GC, and thus, we chose LINC01224 as the target lncRNA for this study. Next, we predicted the downstream miRNAs that might bind to LINC01224 using a target gene prediction website. According to the data retrieved from the website, miR-193a-5p may specifically bind to LINC01224. MiR-193a-5p has been widely reported to play a role in a variety of diseases and pathological processes.11,12 For example, miR-193a-5p downregulated SRR to inhibit osteosarcoma cell migration, and its expression in highly metastatic osteosarcoma cell lines was suppressed.13 Besides, miR-193a-5p was significantly attenuated apoptosis by targeting Bach2 in prostate cancer.14 In addition, through bioinformatics prediction, we found that miR-193a-5p can further target cyclin-dependent kinase 8 (CDK8), which has been reported to have a cancer-promoting effect in various tumors.15,16 Therefore, the main purpose of the current research is to study the roles and interaction of LINC01224, miR-193a-5p, and CDK8 in regulating the biological characteristics of GC cells.

2 | MATERIALS AND METHODS

2.1 | Ethics statement, specimens, and animals

The clinical and pathological specimens of gastric cancer (N = 40) used in this study were obtained from Weifang People’s Hospital, and the study was approved by the Ethics Committee of our hospital. All patients agreed that their pathological specimens could be used for this study. The animal experiment had been reviewed and supervised by the Institutional Animal Care and Use Committee of Weifang People’s Hospital. Fifty male C57 mice (6 weeks old) were purchased from Jiangsu ALF Biotechnology Co., LTD. (China). All mice were placed in a SPF level animal house at 22°C under the conditions of 12 hours of light/12 hours of darkness with free access to food and drink.

2.2 | Cell culture

Human gastric epithelial cells GES-1 cells (CL-0563) and GC cell lines HGC-27 (CL-0107), AGS (CL-0022), SNU-1 (CL-0474), and SNU-5 (CL-0444) produced by Wuhan Procell Life Science & Technology Co., Ltd. (China) were cultured in DMEM (PM150210A, Procell, China) containing 10% of FBS (164210, Procell) in a 5% of CO2 incubator (SCO6WE-2, SHELLAB, USA) at 37°C.

2.3 | Transfection

Overexpressed LINC01224 (LINC01224, vector: pcDNA3.1), shRNA-targeted LINC01224 (sh-LINC01224, target sequence: GGAAAGGCACCTGAGTAATTT, vector: pLKO.1), and siRNA-targeted CDK8 (siCDK8, target sequence: CTGTGACAATGGACTATGACTTT) recombinant plasmids were synthesized by Guangzhou GENESEED Biotechnology Co., Ltd. MiR-193a-5p inhibitor (B03001) and inhibitor control (B04003) were purchased from Shanghai GenePharma Company (China). The above plasmids or siRNAs were transfected into HGC-27 or SNU-1 cells (1 × 10^6 cells/ml) using Lipofectamine 3000 reagent (L3000001, Invitrogen, USA) according to the manufacturers’ instructions. After 24 hours (h) of cell transfection, real-time quantitative PCR (RT-qPCR) or Western blot was performed to detect the transfection rate.

2.4 | RT-qPCR

MRNA or miRNA from clinical specimens or GC cell lines was extracted using a TaKaRa MiniBEST Universal RNA Extraction Kit (9767, Japan) or RNAiso for Small
RNA (9753A, Japan). The complementary DNA synthesis was conducted with a PrimeScript RT reagent Kit (RR047A, TaKaRa, Japan). The RT-qPCR reaction was performed using LightCycler 480 II (D10003, Roche, Switzerland) and TB Green Premix Ex Taq II (RR820L, TaKaRa, Japan) or a Mir-X miRNA qRT-PCR TB Green Kit (638314, Clontech, Japan). mRNA expressions was calculated by the 2−ΔΔCt method, and normalized to GAPDH or U6. Primers used were as follows (5’-3’): LINC01224 (GCCTGAGCTTTGCTCATAGAA, TGTGCGTGCAAAATCACACCT); miR-193a-5p (CTTTGCGGGAGATGAGT, TCGTATCCAGTGCGTGTCGT); CDK8 (ACCTGTTTGAATACGAGGGCT, TGCCGACACGATGAGATCCCAGT); GAPDH (ACCTGACCTGCCGTCCTAGAA, TCCACCACCTGTTGCTGTA); U6 (CTCGTTTCGGCAGCACA, AACGCTTCACGAATTTGCGT).

2.5 | In Situ Hybridization (ISH)

RNAscope 2.5 HD Assay-BROWN (322300, Advanced Cell Diagnostics, China) was used to perform ISH. The brief steps were as follows: After fixation, tissue sections were subjected to antigen repair in Biocare Decloaker for 3 minutes, and then, protease was added for 30 minutes for pre-osmosis treatment. Then, a RNA-specific probe was used to hybridize with the target RNA at 40°C, and the signal was amplified through a multistep process. The expression levels of all probes were observed using DAB solution. The probes used from Advanced Cell Diagnostics (China) were LINC01224 (536619), hs-PPIB (positive control, 313901), and dapB (negative control, 310043). The clear chromophore precipitate spot formed by each single RNA transcript was observed under a BX53M microscope at 100 times magnification (Olympus, Japan).

2.6 | Cell viability assay

The viability of HGC-27 or SNU-1 cells was analyzed using an MTT Cell Proliferation and Cytotoxicity Assay Kit (M1020) from Solarbio (China). HGC-27 or SNU-1 cells were cultured routinely for 24, 48, or 72 hours in a 96-well plate, and then, 10 μl of fresh medium and 90 μl of MTT reagent were added to each well to further culture the cells for 4 hours. Afterward, 110 μl of Formazan solution was added and shaken for 10 minutes. Finally, the absorbance was measured at 490 nm using a microplate absorbance reader (E0228) from Beyotime (China). After 48 hours of transfection in each group, cells were made into a cell suspension with a density of 1 × 10⁵ cells/ml using 1× Binding Buffer. Then, 5 μl of FITC Annexin V and 5 μl of propidium iodide (PI) were added to 100 μl of cell suspension, and the cells were subsequently incubated in the dark for 15 minutes. Finally, a CytoFLEX Flow cytometer (Beckman Coulter, USA) was used to detect the apoptosis of cells in each group and calculate the apoptosis rate.

2.7 | Flow cytometry

Cell apoptosis was analyzed with an Annexin V-FITC/PI Apoptosis Detection Kit (CA1020) produced by Solarbio (China). Total protein of HGC-27 or SNU-1 cells was lysed and extracted using RIPA buffer containing 1% of PMSF (R0010) from Solarbio (China), and its concentration was determined using a BCA kit (PC0020, Solarbio). Twenty μg of protein samples were used to electrophoresis on SDS-PAGE gels for 2 h, and subsequently transferred to a PVDF membrane (ISEQ00010, Millipore, USA), and then, blocked for 1 h, and incubated with primary antibody at 4°C overnight, following further incubated with secondary antibody for 1 h, and developer (SW2040, Solarbio, China) was added dropwise for exposure and development. Following are the antibodies used in this experiment, which were obtained from Abcam (UK): E-cadherin (ab40772, 1/10000, 97 kDa), N-cadherin (ab18203, 1 μg/ml, 100 kDa), Vimentin (ab92547, 1/1000, 54 kDa), Snail (ab229701, 1/1000, 21 kDa), Cleaved-caspase-3 (ab2302, 1 μg/ml, 17 kDa), Cyto C (ab133504, 1/5000, 14 kDa), CDK8 (ab224828, 1/2000, 36 kDa), and Goat Anti-Rabbit (1/5000, ab205718).

2.8 | Western blot

The cells were digested with trypsin, and then, resuspended in serum-free medium (1 × 10⁵ cells/ml). Next, 150 μl of cell suspension of each group was added to the upper chamber
of the Transwell chamber (3428, Corning, USA) pre-coated with Matrigel (354230, BD, USA), and 500 μl of complete medium containing 10% of FBS was added to the lower chamber. After 48 hours of incubation, the chamber was removed, and then, the cells on the membrane were fixed with 950 ml/L of ethanol and stained with 0.1% of crystal violet solution (C8470, Solarbio, China). Finally, the invasive cells from five randomly chosen fields of each sample were counted under an inverted microscope (magnification ×200).

2.11 | Bioinformatics assay

The GEPIA website (http://gepia.cancer-pku.cn/) was used to analyze the expression of LINC01224 in stomach adenocarcinoma (STAD) (T = 408, N = 211). The relationship between LINC01224, miR-193a-5p, and CDK8 was predicted using starBase v2.0 (http://starbase.sysu.edu.cn/) and the TargetScan v7.2 website (http://www.targetscan.org/vert_72/), and confirmed by dual luciferase assay.

2.12 | Dual luciferase assay

We verified the binding of LINC01224 and miR-193a-5p, miR-193a-5p and CDK8 through dual luciferase experiments. The specific steps were as follows: The sequences combined with the wild-type or mutant site of LINC01224 or CDK8 were reorganized to pmirGLO vectors (CL414-01, Biomed, China). Next, the above report vectors were co-transfected with miR-193a-5p.
(A) Relative LINC01224 expression level in HGC-27 and SNU-1 cells treated with Blank, Vector, LINCO1224, and sh-LINC01224.

(B) Relative cell viability in HGC-27 and SNU-1 cells treated with Blank, Vector, LINCO1224, and sh-LINC01224 at 24, 48, and 72 hours.

(C) Apoptosis rates in HGC-27 and SNU-1 cells treated with Blank, Vector, LINCO1224, and sh-LINC01224.

(D) Western blot analysis of Bcl-2, Bax, Cleaved caspase-3, Cyto c, and GAPDH in HGC-27 and SNU-1 cells treated with Blank, Vector, LINCO1224, and sh-LINC01224.

(E) Relative Bcl-2 protein expression levels in HGC-27 and SNU-1 cells treated with Blank, Vector, LINCO1224, and sh-LINC01224.

(F) Relative Bax protein expression levels in HGC-27 and SNU-1 cells treated with Blank, Vector, LINCO1224, and sh-LINC01224.

(G) Relative Cleaved caspase-3 protein expression levels in HGC-27 and SNU-1 cells treated with Blank, Vector, LINCO1224, and sh-LINC01224.

(H) Relative Cyto c protein expression levels in HGC-27 and SNU-1 cells treated with Blank, Vector, LINCO1224, and sh-LINC01224.
inhibitor or inhibitor control into HGC-27 or SNU-1 cells using Lipofectamine 3000. At last, the luciferase activity was determined using a dual luciferase analysis kit (E1910, Promega, USA) and a GloMax 20/20 Luminometer (Promega, USA).

2.13 | Tumorigenesis assay

C57 mice (n = 10) were divided into five groups. The Control group received no treatment, and the other groups were subjected to subcutaneous injection. HGC-27-Vector group: HGC-27 cells transfected with pcDNA3.1 were injected into C57 mice. HGC-27-LINC01224 group: HGC-27 cells transfected with LINC01224 were injected into C57 mice. SNU-1-Vector group: SNU-1 cells transfected with pcDNA3.1 were injected into C57 mice. SNU-1-LINC01224 group: SNU-1 cells transfected with LINC01224 were injected into C57 mice. The xenotransplantation tumor length (L) and width (W) were scientifically measured every 5 days until day 20. After 20 days, the mice underwent deep anesthesia by intraperitoneal injection of 150 mg/kg sodium pentobarbital (B5646-50 mg, ApexBio, USA), and the tumors were taken and photographed and analyzed.

2.14 | Statistical analysis

Statistical analysis was performed using GraphPad prism 8.0. The measurement data were expressed as mean ± SD. The differential expression of genes in normal tissues and cancer

![Figure 3](image-url)

**FIGURE 3** The effect of LINC01224 on gastric cancer cell migration and invasion and EMT-related proteins. A, Wound healing assay results demonstrated that overexpressed LINC01224 facilitated gastric cancer cell migration, while silent LINC01224 had the opposite effect (magnification ×100). B, Transwell assay results indicated that overexpressed LINC01224 accelerated gastric cancer cell migration, while silent LINC01224 had the opposite effect (magnification ×200). C–G, Western blot was performed to detect the effect of LINC01224 on the protein expressions of E-cadherin, N-cadherin, Vimentin, and Snail. GAPDH was set as control. *p < 0.05, **p < 0.01, ***p < 0.001 vs Vector. EMT: epithelial-mesenchymal transition

![Figure 4](image-url)

**FIGURE 4** LINC01224-targeted miR-193a-5p, which was downregulated in gastric cancer. A, StarBase v2.0 predicted the target miRNA of LINC01224. B, Dual luciferase assay results verified that LINC01224 bound to miR-193a-5p. C, RT-qPCR results showed that the expression of miR-193a-5p in gastric cancer samples was downregulated (N = 40). D, The expression of miR-193a-5p in GES-1, HGC-27, AGS, SNU-1, and SNU-5 cells was assessed by RT-qPCR. U6 was set as control. E, The expression of LINC01224 in GES-1 cell was assessed by RT-qPCR. GAPDH was set as control. &&&p < 0.001 vs IC; △△△p < 0.001 vs Normal; +++p < 0.001 vs GES-1
(A) Relative miR-192a-5p expression level

(B) Relative cell viability (%)

(C) Apoptosis rates (%)

(D) Western blot analysis of Bcl-2, Bax, Cleaved caspase-3, Cyto c and GAPDH

(E) Relative Bcl-2 protein expression levels

(F) Relative Bax protein expression levels

(G) Relative Cleaved caspase-3 protein expression levels

(H) Relative Cyto c protein expression levels
tissues was analyzed using paired sample t-test. One-way analysis of variance was used for comparison among multiple groups, and post hoc pairwise comparisons were performed using Bonferroni or Dunnott or Tukey tests. \( p < 0.05 \) was considered as statistically significant.

### 3 | RESULTS

#### 3.1 Abnormally elevated LINC01224 was found in GC samples and cell lines

LINC01224 was highly expressed in STAD, which attracted our attention (\( p < 0.05 \), Figure 1A). Next, we verified that LINC01224 expression was increased in GC tissues (\( p < 0.001 \), Figure 1B). In order to understand the function of LINC01224 in GC, we then confirmed this result by ISH experiments. The expression of LINC01224 was upregulated in the tumor group, and there was a clear positive reaction (brown) (magnification \( \times100 \), Figure 1C). Similarly, the expression of LINC01224 in HGC-27, AGS, SNU-1, and SNU-5 cell lines was higher than that of GES-1 cell lines (\( p < 0.01 \), Figure 1D). Among them, the expression of LINC01224 in SNU-1 and HGC-27 cells was the most significant and was selected for later experiments (\( p < 0.001 \)).

#### 3.2 Overexpressed LINC01224 facilitated the malignant progression of GC cells, while silencing LINC01224 had the opposite effect

Exogenous upregulation or downregulation of LINC01224 was used to study its effect on the biological characteristics of GC cells (\( p < 0.01 \), Figure 2A). It was found that overexpressed LINC01224-enhanced cell viability and slowed down the rate of apoptosis, while silent LINC01224 weakened cell viability and facilitated cell apoptosis in GC cells (\( p < 0.05 \), Figure 2B–C). Mechanistically, in comparison to the Vector group, the expression of Bcl-2 was elevated in the LINC01224 group, while the expressions of Bax, Cleaved caspase-3, and Cyto C were suppressed (\( p < 0.01 \), Figure 2D–H). Contrarily, the expression of Bcl-2 was downregulated in the sh-LINC01224 group, while the expressions of Bax, Cleaved caspase-3, and Cyto C were augmented (\( p < 0.01 \), Figure 2D–H). As shown in Figure 3A–B, overexpressed LINC01224 accelerated GC cell migration and invasion, while silent LINC01224 had the opposite effect (\( p < 0.05 \)). Besides, EMT-related proteins were detected to evaluate the possible mechanism of LINC01224 in GC cells. It was found that the expression of E-cadherin in the LINC01224 group was suppressed, and the expressions of N-cadherin, Vimentin, and Snail were promoted, while the sh-LINC01224 group displayed an opposite result (\( p < 0.01 \), Figure 3C–G).

#### 3.3 MiR-193a-5p was directly targeted by LINC01224 and was downregulated in GC

We analyzed the potential of LINC01224 to interact with miRNA and found that LINC01224 may directly bind to miR-193a-5p, which was confirmed by dual luciferase reporter gene assay (\( p < 0.001 \), Figure 4A–B). In addition, the expression of miR-193a-5p in GC clinical samples was suppressed (\( p < 0.001 \), Figure 4C). Similarly, miR-193a-5p expression was also significantly inhibited in GC cell lines (\( p < 0.001 \), Figure 4D). To be on the safe side, we also tested the effect of overexpressed LINC01224 and miR-193a-5p inhibitor (I) on the expression of LINC01224 in normal GES-1 cells. The results were the same as we expected: Overexpression of LINC01224 significantly increased the mRNA level of LINC01224, and whereas miR-193a-5p inhibitor downregulated LINC01224 (\( p < 0.05 \), Figure 4E). Subsequently, we examined the transfection efficiency of miR-193a-5p and found that silencing LINC01224 promoted miR-193a-5p expression (\( p < 0.001 \), Figure 5A).

#### 3.4 LINC01224 interacted with miR-193a-5p and affected the biological behavior of GC cells

Functional experiments revealed that miR-193a-5p inhibitor increased cell viability, restrained apoptosis, and interfered with the expression of apoptosis-related proteins, and partially offset the effect of sh-LINC01224 on cell viability and apoptosis (\( p < 0.05 \), Figure 5B–H). Meanwhile, miR-193a-5p inhibitor facilitated cell migration and invasion, and notably overturned the effect of sh-LINC01224 (\( p < 0.01 \), Figure 6A–B). In addition, compared with the Vector+inhibitor control (IC) group, E-cadherin expression was repressed in the Vector+inhibitor (I) group and strengthened in the sh-LINC01224+IC group (\( p < 0.001 \), Figure 6C–D).
Figure 6C–D). Conversely, the expressions of N-cadherin, Vimentin, and Snail were increased in the Vector+I group and suppressed in the sh-LINC01224+IC group \(p < 0.001\), Figure 6C–E–G). Furthermore, the regulation of the above genes in the Vector+I group or the sh-LINC01224+IC group was significantly reversed in the sh-LINC01224+I group \(p < 0.001\), Figure 6C–G).

### 3.5 MiR-193a-5p bound to CDK8, which was upregulated in GC

Eight regions (position 457–464) were found to interact with miR-193a-5p in the CDK8 3’UTR region (Figure 7A). In the CDK8-WT+ miR-193a-5p inhibitor group, the luciferase activity increased significantly, confirming the binding relationship of miR-139a-5p and CDK8 \(p < 0.001\), Figure 7B). Next, we also detected the expression of CDK8 in GC tissues and cell lines, and found that CDK8 expression was significantly increased \(p < 0.001\), Figure 7C–D). Afterward, SNU-1 and HGC-27 cell lines were transfected with siCDK8 and miR-193a-5p inhibitor alone or in combination. As shown in Figure 7E–G, compared with the IC+siNC group, CDK8 expression was prominently raised in the I+siNC group and visibly decreased in the IC+siCDK8 group \(p < 0.001\). Moreover, the regulation of CDK8 expression in the I+siNC group or the IC+siCDK8 group was evidently overturned in the I+siCDK8 group \(p < 0.001\), Figure 7E–G).

### 3.6 MiR-193a-5p-targeted CDK8 and regulated malignant transformation of GC cells

As shown in Figure 7H, siCDK8 reduced cell viability and partially reversed the effect of miR-193a-5p inhibitor on the viability of GC cells \(p < 0.05\). However, siCDK8-induced apoptosis, and the regulatory effect of miR-193a-5p inhibitor on cell apoptosis was evidently reversed in the I+siCDK8 group compared with the I+siNC group \(p < 0.001\), Figure 8A). Meanwhile, Bcl-2 expression was decreased yet the expressions of Bax, Cleared-caspase-3, and Cyto C were increased in the IC+siCDK8 group; interestingly, siCDK8 also reversed the regulatory effect of miR-193a-5p inhibitor on the above genes \(p < 0.001\), Figure 8B–F). In addition, siCDK8 attenuated cell migration and invasion, increased E-cadherin expression, and diminished N-cadherin, Vimentin, and Snail expressions; moreover, the effect of miR-193a-5p inhibitor on GC cells was offset in the siCDK8 group \(p < 0.01\), Figure 9A–G).

#### 3.7 Regulation of the LINC01224/miR-193a-5p/CDK8 axis in vivo

In vitro experiments indicate that LINC01224 adsorbed miR-193a-5p and regulated CDK8 as a ceRNA to participate in the malignant transformation of GC cells. In xenograft tumor experiments, we found that overexpressed LINC01224 significantly increased the size of the tumor and promoted tumor growth \(p < 0.001\), Figure 10A–B). Subsequently, we verified the expressions of LINC01224, miR-193a-5p, and CDK8 in vivo. As expected, in the HGC-27-LINC01224 and SNU-1-LINC01224 groups, the expressions of LINC01224 and CDK8 elevated while that of miR-193a-5p declined \(p < 0.001\), Figure 10C–E).

### 4 DISCUSSION

Studying the relationship between lncRNA and GC is expected to control GC cell proliferation and metastasis and improve the life quality of GC patients. LINC01224 is a newly identified lncRNA, but its exact function and mechanism in GC remain to be revealed. Here, through a bioinformatics analysis, we found that LINC01224 was upregulated in GC, identified miR-193a-5p as the miRNA that binds to LINC01224, and predicted that CDK8 interacts with miR-193a-5p as its target gene. Based on these analyses, we explored the roles of these three genes in the malignant transformation of gastric cancer cells. In present study, we found that LINC01224 adsorbed miR-193a-5p and upregulated CDK8 to accelerate the progression of gastric cancer.

LncRNAs are considered to be a very important class of molecules that regulate tumors and other diseases. Recent studies have shown that compared with normal cells, LncRNAs are often in a "deregulated" state in tumor cells; meanwhile, overexpression or downregulation of specific LncRNAs in tumor cells can often trigger apoptosis or increase the sensitivity of tumor cells to treatments that induce apoptosis. In this study, by analyzing the LINC01224 levels in 40 pairs of GC tissues and four GC cell lines, we found that LINC01224...
FIGURE 7 MiR-193a-5p-mediated CDK8, which was highly expressed in gastric cancer, regulated cell viability in gastric cancer cells. A–B, TargetScan V7.2 predicted that CDK8 was targeted by miR-193a-5p and the prediction was verified by dual luciferase assay. C–D, RT-qPCR results showed that the expression of CDK8 was increased in gastric cancer samples and cell lines. GAPDH was set as control. E–G, Western blot and RT-qPCR were performed to detect the expression of CDK8 in the IC+siNC, I+siNC, IC+siCDK8, and I+siCDK8 groups. GAPDH was set as control. H, MTT results showed that siCDK8 reduced cell viability and partially counteracted the effect of miR-193a-5p inhibitor. Each experiment was repeated three times independently. *p < 0.05, **p < 0.01, ***p < 0.001 vs IC; ΔΔΔp < 0.001 vs Normal; +++p < 0.001 vs GES-1; *p < 0.05, **p < 0.001 vs IC+siNC; ^p < 0.05, ^^^p < 0.001 vs I+siNC; ^p < 0.05, ***p < 0.001 vs IC+siCDK8. NC: negative control. CDK8: cyclin-dependent kinase 8.
FIGURE 8  MiR-193a-5p-targeted CDK8 and interfered with apoptosis and expressions of apoptosis-related proteins in gastric cancer cells. A, Results from flow cytometry showed that siCDK8 accelerated cell apoptosis and partially counteracted the effect of miR-193a-5p inhibitor. B–F, The effect of miR-193a-5p-mediated CDK8 on the expressions of Bcl-2, Bax, C-caspase-3, and Cyto C in gastric cancer cells was measured with Western blot. GAPDH was set as control. *** $p < 0.001$ vs IC+siNC; ^^^ $p < 0.001$ vs I+siNC; ### $p < 0.001$ vs IC+siCDK8
was highly expressed in GC, and knocking down LINC01224 inhibited GC cell viability, migration, invasion, and EMT as well as accelerated cell apoptosis. To our knowledge, the current study is the first to reveal the role of LINC01224 in GC and suggest that it may be a therapeutic target for GC tumors.

As of now, there have been as few as three reports on LINC01224. The expression trend of LINC01224 in GC in our study is similar to previous studies. LINC01224 can be used to predict the prognosis of breast cancer patients. Xing et al. demonstrated that LINC01224 is upregulated in epithelial ovarian cancer, and its high expression is related to the poor prognosis of patients; the underlying mechanism may be that LINC01224 targets miR-485-5p/PAK4 to form a competitive endogenous RNA network and thereby regulate the progression of epithelial ovarian cancer. In addition, in a recent study of hepatocellular carcinoma,
LINC01224 has been found to act as an oncogene to regulate the miR-330-5p/CHEK1 axis. Based on previous studies, we believe that an lncRNA-miRNA-mRNA pathway related to LINC01224 is also implicated in the development of GC tumors.

Although we found a mutual relationship between LINC01224 and miR-193a-5p and between miR-193a-5p and CDK8 through bioinformatics analysis, the predicted results need to be verified by specific experiments. Fortunately, our experimental results confirmed the predicted relationship between these genes. At the same time, it was found that miR-193a-5p functions as a tumor suppressor gene in GC, and downregulation of miR-193a-5p can reverse the effect of silencing LINC01224 on GC cells. The role of miR-193a-5p in other diseases has been widely reported. For example, upregulation of miR-193a-5p significantly slowed the proliferation and invasion of hepatoblastoma cells by inhibiting DPEP1. miR-193a-5p can be adsorbed by lncRNA TTN-AS1 and participate in the regulation of prostatic cancer cell proliferation and apoptosis. Besides, in two reports on the role of miR-193a-5p in GC, it was found that miR-193a-5p may inhibit GC cell malignant transformation, which is consistent with our results. However, in our study, we further predicted that CDK8 is directly targeted by miR-193a-5p.

In the study of gene expression regulation, miRNA and lncRNA are a very important link. They not only have an important influence on gene transcription, posttranscriptional regulation, and other gene expression modules, but also more importantly, they participate in the body’s physiological and pathological responses through interaction. One of the widely recognized methods is that lncRNA and miRNA competitively bind to regulate the downstream target genes of miRNA. This regulation mechanism has been widely confirmed in GC: LncRNA MT1JP sponged miR-92a-3p and regulated the downstream FBXW7 gene, which in turn affected the process of GC; LINC01234 acts as the ceRNA of miR-204-5p and prevents the activation of CBFB in GC. Our experimental results supplement the related content of lncRNA-miRNA-mRNA mechanism in GC. In addition, we have also analyzed other mechanisms that may change the physiological functions of cells.

Almost all tumors are characterized by the disablement of cell cycle control mechanisms that leads to uncontrolled cell growth, blocked differentiation, and abnormal apoptosis. CDKs are the core of cell cycle regulation, which together with cyclin, cyclin-dependent kinase inhibitors, etc. constitute a cell cycle regulation network system. Moreover, the role of CDK8 in promoting canceration has also been proven multiple times. As Song et al. reported, CDK8 reversed the effect of miR-107 to promote the proliferation of GC cells. Our research also confirmed that upregulation of CDK8 can lead to malignant phenotype and EMT of GC cells, and in vivo experiments also show that LINC01224 inhibits miR-193a-5p and upregulates CDK8 to accelerate tumor growth, suggesting that the LINC01224/miR-193a-5p/CDK8 axis is an important part of the GC regulatory network.

Given our results, it can be concluded that LINC01224 promotes migration, invasion, and EMT and restrains apoptosis in GC cells through the miR-193a-5p/CDK8 axis. Unfortunately, although existing studies have found that LINC01224 is related to GC and has a regulatory effect on GC cells, it was not clear during the experiment whether LINC01224 was exogenously upregulated or silenced whether it would have a functional effect on normal GES-1 cells. This point requires us to reflect and further analyze in future research.

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Not applicable.

CONFLICT OF INTEREST
The authors declare no conflicts of interest.

AUTHORS’ CONTRIBUTIONS
Substantial contributions to conception and design: Hui Sun. Data acquisition, data analysis and interpretation: Jihong Yan, Guangyu Tian, Xiaojun Chen, Wenbo Song. Drafting the article or critically revising it for important intellectual content: Hui Sun. Final approval of the version to be published: Hui Sun, Jihong Yan, Guangyu Tian, Xiaojun Chen, Wenbo Song. Agreement to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of the work are appropriately investigated and resolved: Hui Sun, Jihong Yan, Guangyu Tian, Xiaojun Chen, Wenbo Song.

DATA AVAILABILITY STATEMENT
The analyzed data sets generated during the study are available from the corresponding author on reasonable request.

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