LncRNA SNHG7 Functions as an Oncogene in Cervical Cancer by Sponging miR-485-5p to Modulate JUND Expression

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Background: Long non-coding RNA (LncRNA) SNHG7 is involved in the development of multiple cancers. However, its role in cervical cancer (CC) has not been elucidated. This study aimed to explore the function of SNHG7 in CC progression and the underlying mechanisms.

Materials and Methods: The expression levels of SNHG7 and miR-485-5p in CC tissues and cell lines were measured by qPCR. Functional experiments including CCK-8 assay, wound healing assay, transwell assay, flow cytometry, Western blot, luciferases reporter assay and immunoprecipitation (RIP) were performed to explore the SNHG7/miR-485-5p/JUND pathway. Additionally, in vivo study was carried out by establishing tumor xenograft models.

Results: We found that SNHG7 was markedly enhanced in CC tissues and cell lines, and associated with poor clinical characteristics. In vitro, knockdown of SNHG7 inhibited CC cell proliferation, migration and invasion, as well as aggravated cell apoptosis. As to mechanism investigation, rescue experiments revealed that miR-485-5p inhibitor could partially reverse the effects on CC cells induced by SNHG7 knockdown. SNHG7 upregulated JUND expression via miR-485-5p. Moreover, tumor xenograft models were established to confirm the findings in vivo.

Conclusion: SNHG7 promoted CC progression through miR-485-5p/JUND axis. The SNHG7/miR-485-5p/JUND pathway might provide a novel therapeutic target for CC treatment.

Keywords: LncRNA SNHG7, cervical cancer, CC, miR-485-5p, JUND, cell proliferation

Introduction

Cervical carcinoma (CC) is the second most common malignant tumor in females. 1 CC is mainly caused by human papillomavirus (HPV) infection. Upon HPV infection, E6 and E7 oncoproteins begin to express and regulate downstream gene expressions, thus contributing to continuous cell proliferation to malignant transformation. 2 There are about 500,000 women get CC and 260,000 die from it every year. 3 Although the treatment technology of CC has been greatly improved, the 5-year overall survival rate is still poor and metastatic CC is incurable. 4,5 Thus, it is urgent to improve new therapeutic approaches. Understanding the underlying molecular mechanisms of CC development will contribute to CC treatment.

Long non-coding RNAs (LncRNAs) are a subclass of non-coding RNAs with more than 200 nucleotides. 6 Although they were considered as noises with no biological functions, increasing evidences revealed that LncRNAs play key roles in many processes, such as chromatin modulating, RNA processing and protein
expression regulation. As to cancers, more and more lncRNAs are reported to play as oncogenes or suppressors in multiple tumor progressions. For example, GHET1 could drive prostate cancer progression by regulating KLF2 thereby activating HIF-1alpha/Notch-1 signaling pathway. TTN-AS1 promoted colorectal cancer proliferation via sponging miR-376a-3p and upregulating KLF15. Small nucleolar RNA host gene 7 (SNHG7), with a length of 2157 bp, is a newly discovered lncRNA which was reported to enhance the progression of a series of cancer types, including pancreatic cancer, lung cancer, bladder cancer, colorectal cancer and esophageal cancer. However, whether SNHG7 also play roles in CC development has not been elucidated.

In lncRNAs are likely to function via sponging miRNAs, thereby regulating downstream target genes expression. miR-485-5p was predicted as a target of SNHG7 by informatics tool. Moreover, miR-485-5p was identified as a tumor suppressor in a series of cancers, including cervical cancer. In this study, we aimed to investigate the effects on CC progression and the possible mechanisms.

Materials and Methods

Human Tissues

A total of 45-paired CC tissues and normal tissues were obtained from the First Affiliated Hospital of Guangxi Medical University. Patients had not received systemic or local treatment prior to surgical resection. Table 1 shows the clinicopathologic characteristics. Tissue samples were immediately treated with liquid nitrogen and stored at −80°C. Informed consent was signed by every patient. Written informed consent was obtained from every involved patient. The Ethics Committee of the First Affiliated Hospital of Guangxi Medical University approved this study.

Cell Culture

CC cell lines (Hela, SiHA, C-33A and HT-3), human normal cervical cell line (Ect1/E6E7) and HEK-293T cells were purchased from Cell Bank of the Chinese Academy of Sciences (Shanghai, China) and incubated in DMEM (Gibco, USA) plus with 10% FBS and 1% penicillin-streptomycin (Beyotime, China). Cells were cultured in a humidified atmosphere with 5% CO₂ at 37°C.

RT-qPCR

Total RNA was extracted from tissues or cells using Trizol (Invitrogen, USA) following the protocol. The isolated RNA was then reverse-transcribed into cDNA with the help of MMLV (Promega, Wuhan). cDNA was taken as templates and qPCR was conducted on a PikoReal Real-Time PCR System (ThermoFisher Scientific, USA) with SYBR GREEN SMART KIT (Tiangen, Nanjing, China). The expression levels of target genes were calculated by 2^−ΔΔCt method. U6 was used to normalize the relative expression level of miR-485-5p, while β-Actin was used for normalization of SNHG7 and JUND expressions. Specific primers are listed in Table 2.

CCK-8 Assay

CCK-8 assay was conducted to assess the viability of CC cells using CCK-8 Assay Kit (Dojindo, Toyoto). In brief, cells were seeded into 96-well plates with a concentration of 2×10^3 cells/well. After incubation of indicated hours, we added 100μL fresh medium and 10μL CCK-8 reagent to each well of 96-well plates (Corning, USA). Two hours later, optical density was measured at 450 nm daily using a microplate reader (ThermoFisher Scientific, USA).

Flow Cytometry

Cell apoptosis was detected using a FITC/PI Apoptosis Kit (Thermo, USA). Forty-eight hours after transfection, Annexin V binding buffer was taken to suspend cells.

| Table 1 Characteristics of Cervical Cancer Patients |
|-----------------------------------------------|
| Characteristics                           | N 45 | SNHG7 Expression | P value |
|-----------------------------------------------|------|------------------|---------|
| Age                                          |      |                  |         |
| ≥50                                          | 25   | 8                | 0.512   |
| <50                                          | 5    | 7                |         |
| Tumor Size                                   |      |                  |         |
| ≥4                                           | 18   | 8                | 0.012*  |
| <4                                           | 7    | 12               |         |
| FIGO Stage                                   |      |                  |         |
| III–IV                                      | 10   | 8                | 0.026*  |
| I–II                                        | 8    | 19               |         |
| Lymph-Node Metastasis                        |      |                  |         |
| Yes                                          | 15   | 7                | 0.018*  |
| No                                           | 9    | 15               |         |
| Histological Grade                          |      |                  |         |
| Well                                         | 12   | 14               | 0.518   |
| Moderately/poorly                           | 12   | 7                |         |

Note: *P < 0.05 represents statistical differences.
Then, Annexin V/FITC was added in the dark. Five minutes later, PI was added and apoptosis was analyzed by FACS Aria I/II flow cytometer (BD Biosciences, USA).

Wound Healing Assay
Wound healing assay was applied to estimate cell migration rate. When transfected cells reached 90% confluence, a new micropipette tip was taken to make a scratch across the surface. PBS was used to wash the suspended cells, and FBS-free DMEM (Gibco, USA) was added. Twenty-four hours later, images of cells were taken by a light microscope (Olympus, Japan).

Transwell Assay
Transwell assay was conducted to determine the cell invasion ability. The upper chamber of a transwell plate (Corning, USA) was coated with Matrigel (BD, USA). Complete medium with 10% FBS was added into the lower chamber. Twenty-four hours after transfection, the lower chamber was added with 70% ethanol and 0.1% crystal violet. The invaded cells were stained and counted from five visual fields which were randomly chosen.

Luciferase Reporter Assay
miR-NC or miR-485-5p mimics was co-transfected with SNHG7 mut or SNHG7 wt into HEK-293T cells for 48 h. Dual-luciferase reporter assay (Promega) was carried out according to the protocol. Similarly, miR-NC or miR-485-5p mimics was co-transfected with JUND mut or JUND wt, and dual-luciferase reporter assay (Promega, USA) was conducted subsequently.

Western Blot
Total protein was isolated from tissues or cells by RIPA (Sangon, Shanghai, China). Equal amounts of each sample were loaded and separated by 10% SDS-PAGE. Proteins were transferred from gel to PVDF membranes (Millipore, USA) and blocked with 5% silk milk for 1 h. at room temperature. The membranes were then incubated with specific primary antibodies: anti-JUND (1:1000, CST, Beijing); anti-β-Actin (1:2500, CST, Beijing) overnight at 4°C. Subsequently, the membranes were incubated with HRP-conjugated antibody for 2 h at normal temperature. The blot signals were washed by TBST and visualized with an ECL Kit (Solarbio, Beijing, China). Images were taken by an ImageQuant LAS4000 Biomolecular Imager (GE Healthcare).

Immunohistochemical Staining
Ki67 expression was estimated via incubation in monoclonal antibody against Ki67 (1:500, CST, Beijing) overnight at 4°C. Subsequently, the samples were washed by PBS for three times and incubated with biotin-labeled IgG secondary antibody (1:500, SPT, USA). After 1 h, the samples were treated with HRP-conjugated streptavidin (CST, USA) and developed by DAB. Signals were visualized by EnVision system (Dako, Glostrup, Denmark).

Animal Experiments
Four-week-old BALB/C nude mice were purchased from Shanghai LAC Laboratory (Shanghai, China) and randomly divided into 2 groups (6 mice/group). Hela cells that were transfected with sh-NC or sh-SNHG7 stably were injected into flanks of mice. Then, the tumor volumes

Table 2 Primers of qRT-PCR

| Gene    | Primers                        |
|---------|--------------------------------|
| SNHG7   | **Forward**<br>5′-CGATACCCATTGAACACGCTGC-3′<br>**Reverse**<br>5′-GGTGGAGGTCGCCCGAGT-3′ |
| miR-485-5p | **Stem-loop RT primer**<br>**Forward**<br>5′-GTCGATCCAGTGAAGGCTGAGTCCGAGGTTCCAGGATTGACTACGACTGGATACGACGAATTCC-3′<br>**Reverse**<br>5′-AGAGGCTGGGCCGCTGG-3′<br>5′-GTCGAGGGTGCGAGGTTG-3′ |
| JUND    | **Forward**<br>5′-AGGACCTTCAGCACAAGGCGAGAC-3′<br>**Reverse**<br>5′-GTTGGGCTACACGGCCGCTG-3′ |
| β-Actin | **Forward**<br>5′-AGGCCACATCGCTCAGACAC-3′<br>**Reverse**<br>5′-GCCCAATAACGACCAAAATCC-3′ |
| U6      | **Forward**<br>5′-GCTTCGGCGACGACATATACCTAAAATG-3′<br>**Reverse**<br>5′-CGTTCAGCAATTGGCTGT-3′ |
were detected every week. After 5 weeks, mice were sacrificed and the weights of tumors were measured. Then, the tumor tissues were collected for following experiments to detect the expressions of miR-485-5p, JUND and Ki67. Ethical and legal approval was obtained from Ethics Committee of the First Affiliated Hospital of Guangxi Medical University prior to the commencement of the study. All experiments were performed following the guidelines and regulations of Ethics Committee of the First Affiliated Hospital of Guangxi Medical University.

**Statistical Analysis**

Data were presented as mean±standard deviation (SD). Each sample for experiments were repeated at least 3 times. SPSS 17.0 and Graphpad Prim6 was taken for statistical analysis. Student’s t-test was applied for the difference between two groups and one-way ANOVA was for comparisons among multiple groups. P<0.05 suggested statistically significant.

**Results**

**SNHG7 Was Upregulated in CC Tissues and Cell Lines**

To explore the role of SNHG7 in CC development, the first step for us was to detect the expression pattern of SNHG7 in CC tissues and cell lines. As shown in Figure 1A, SNHG7 expressed markedly more in the CC tissues than the normal tissues. Similarly, SNHG7 expressed a higher level in CC cell lines (Hela, SIHA, C-33A and HT-3) than in normal cell line (Ect1/E6E7) (Figure 1B). Additionally, CC patients with higher SNHG7 expression levels have poorer overall survivals (Figure 1C).

**SNHG7 Knockdown Inhibited CC Proliferation, Migration and Invasion**

To investigate the effects of SNHG7 on CC development, we performed functional experiments using SNHG7 siRNA (si-SNHG7). Figures 2A and S1 showed that both the two siRNAs against SNHG7 exerted significant knockdown efficiency, and we chose si-SNHG7#2 for the following experiments. Next, we transfected si-SNHG7 or negative control (si-NC) into CC cell lines. CCK-8 assay suggested that si-SNHG7 inhibited CC cell viability (Figure 2B). Flow cytometry assay showed a significant promotion in CC cell apoptosis after SNHG7 silencing (Figure 2C). Suppression of SNHG7 also inhibited CC cell migration (Figure 2D) and invasion (Figure 2E). These primary results indicated that SNHG7 might act as an oncogenic role in CC progression.

**SNHG7 Acted as a ceRNA for miR-485-5p**

To know the potential mechanisms underlying SNHG7-regulated CC, we investigated the relationship between SNHG7 and miR-485-5p. StarBase 3.0 predicted the interacting sites of SNHG7 and miR-485-5p (Figure 3A), which was confirmed by the following luciferase reporter assay (Figure 3B). As expected, miR-485-5p was upregulated upon SNHG7 knockdown, and SNHG7 was
downregulated with miR-485-5p mimics transfection (Figure 3C and D). Moreover, RIP assay demonstrated that SNHG7 and miR-485-5p directly interacted in Ago2 complex (Figure 3E). Furtherly, SNHG7 was found downregulated in CC tissues (Figure 3F) and associated negatively with miR-485-5p expression (Figure 3G).
Figure 3 SNHG7 sponged miR-485-5p. (A) Predicted binding sites between SNHG7 and miR-485-5p from Starbase. (B) miR-NC mimics or miR-385a-5p mimics was co-transfected with SNHG7 mut or SNHG7 wt, and luciferase reporter assay was performed 48 h later. (C) si-NC or si-SNHG7 was transfected into EC cells, and the relative expression of miR-485-5p was detected by qPCR 48 h later. (D) miR-NC mimics or miR-485-5p mimics was transfected into EC cells, and the relative expression of miR-485-5p was detected by qPCR 48 h later. (E) The direct interaction between SNHG7 and miR-485-5p was detected by RIP with Ago-2 antibody. (F) The relative expression of miR-485-5p in CC tissues and adjacent normal tissues was measured by qPCR. (G) The relationship between SNHG7 and miR-485-5p in CC patients was analyzed by Pearson’s correlation analysis. **P<0.01.
Figure 4 SNHG7-induced EC cell activity changes was mediated by miR-485-5p. (A) miR-485-5p expression was measured by qPCR with miR-485-5p inhibitor transfection. Then, EC cells were transfected with si-SNHG7 in the presence of miR-NC or miR-485-5p mimics, (B) cell viability was measured by CCK-8 assay; (C) cell apoptosis was determined by flow cytometry; (D) migration was detected by wound healing assay; (E) cell invasion was detected by transwell assay; (F) statistical assays were presented in bar graphs. **P<0.01 versus si-NC+miR-NC; ##P<0.01 versus si-SNHG7+miR-NC.
Suppression of miR-485-5p Partially Reversed the Effects of SNHG7 on CC Cells

To determine whether SNHG7 exhibited biological functions via miR-485-5p, we co-transfected miR-485-5p inhibitor or miR-NC with SNHG7 into CC cells. Figure 4A shows the inhibitory property of miR-485-5p inhibitor. Furtherly, miR-485-5p attenuated the suppression of SNHG7 on CC cell viability (Figure 4B), migration (Figure 4D) and invasion (Figure 4E). Also, depletion of miR-485-5p abolished the promotion of SNHG7 on CC cell apoptosis (Figure 4C). Figure 4F displays the quantitative analysis.

miR-485-5p Targeted JUND Directly

The bioinformatics TargetScan software revealed that miR-485-5p had complementary binding sites with JUND, which was verified by luciferase reporter assay (Figure 5A and B). JUND protein expression was inhibited

Figure 5 miR-485-5p directly targeted JUND. (A) Predicted binding sites between miR-485-5p and JUND from TargetScan. (B) miR-NC mimics or miR-485-5p mimics was co-transfected with JUND mut or JUND wt, and luciferase reporter assay was performed 48 h later. (C) miR-NC or miR-485-5p mimics was transfected into EC cells, and the relative expression of JUND was detected by Western blot 48 h later. (D) The relative expression of JUND in CC tissues and adjacent normal tissues was measured by qPCR. (E) The relationship between miR-485-5p and JUND in CC patients was analyzed by Pearson’s correlation analysis. **p<0.01.
upon miR-485-5p overexpression (Figure 5C). Furtherly, we examined the expression profile of JUND in vivo and found that JUND expressed more in CC tissues than in normal tissues (Figure 5D). Additionally, JUND was negatively associated with miR-485-5p in CC tissues from patients (Figure 5E).

**JUND Was Upregulated by SNHG7/miR-485-5p Axis**

Previously found that SNHG7 sponged miR-485-5p (Figure 3) and miR-485-5p targeted JUND (Figure 5). Herein, we wanted to investigate whether SNHG7 could regulate JUND via miR-485-5p. As shown in Figure 6A–C, both mRNA and protein expression levels of JUND were downregulated upon SNHG7 depletion. However, miR-485-5p could mitigate the inhibitory effects. Moreover, SNHG7 and JUND showed positive relationship in CC tissues (Figure 6D).

**SNHG7 Depletion Reduced Tumorigenesis in vivo**

In vivo experiments were further conducted to validate the oncogenic function of SNHG7 in vitro. Xenograft tumor models were established in nude mice by injecting sh-NC or sh-SNHG7 stably transfected Hela cells. We observed that the tumor growth was more slowly in sh-SNHG7 implanted group than sh-NC group (Figure 7A). The tumor weight measured 5 weeks later was lighter in sh-SNHG7 mice, in

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**Figure 6** JUND was regulated by SNHG7/miR-485-5p axis. si-SNHG7 was co-transfected with si-NC or miR-485-5p inhibitor, then the mRNA and protein expression levels of JUND were estimated by qPCR (A) and Western blot (B, C). (D) The relationship between SNHG7 and JUND in CC patients was analyzed by Pearson’s correlation analysis. **P<0.01 versus si-NC+miR-NC; ##P<0.01 versus si-SNHG7+miR-NC.
comparison with sh-NC group (Figure 7B). Moreover, the expression of miR-485-5p was elevated in sh-SNHG7 group (Figure 7C), whereas the JUND protein expression was lowly expressed (Figure 7D). IHC staining furtherly demonstrated that Ki67 was markedly upregulated by SNHG7 depletion (Figure 7E). These results indicated that SNHG7-regulated CC progression via sponging miR-485-5p and thereby upregulating JUND expression (Figure 8).

**Discussion**

Increasing study has reported that dysregulated lncRNAs were involved in the regulation of cancer progression.\(^{17,18}\) A series of lncRNAs showed abnormal expressions in CC tissues, such as MNX1-AS1, HCP5, GAS5 and PVT1.\(^{19-22}\) Also, SNHG7 was observed overexpressed in multiple cancer types and exhibited oncogenic roles in multiple cancer types. For example, SNHG7 enhanced cell viability, as well as inhibited cell apoptosis in gastric cancer.\(^{23}\) Silencing of SNHG7 attenuating cell proliferation ability and driven cell apoptosis in thyroid cancer development.\(^{24}\) SNHG7 knockdown inhibited EMT initiation and tumorigenesis in breast cancer.\(^{25}\) However, whether SNHG7 influence CC progression has not been elucidated. Herein, our data revealed that SNHG7 was highly expressed in CC tissues, and depletion of SNHG7 reduced CC cell viability, migration and invasion, as well as enhanced apoptosis. The findings indicated that SNHG7 play oncogenic roles in CC development.

Acting as ceRNA for miRNAs was an important mechanism of lncRNA-regulating cancer progression. SNHG7 was reported to participate in cancer development by sponging miRNAs. For instance, SNHG7 sponged miR-381 to drive breast cancer.\(^{26}\) Qi et al found SNHG7 acted as ceRNA for miR-503 to accelerate prostate cancer.\(^{27}\) Han et al demonstrated that silencing of SNHG7 suppressed epithelial–mesenchymal transition in prostate cancer by targeting miR-324-3p.\(^{28}\) Also, many lncRNAs were found to regulate CC by interacting with miRNAs. For example, DTRG1 sponged miR-330-5p to enhance CC progression.\(^{29}\) NEAT1 acted as an oncogene in CC by targeting miR-193-3p.\(^{30}\) In the present work, to explore the underlying mechanism of SNHG7-regulated CC development, we predicted miR-485-5p as the target of SNHG7. miR-485-5p exerted tumor-suppressive effects in multiple cancers, including hepatocellular carcinoma,\(^{31}\) gastric cancer,\(^{32}\) papillary thyroid cancer\(^{33}\) and breast cancer.\(^{34}\) Herein, we observed that miR-485-5p inhibitor

![Figure 7](image_url) SNHG7 knockdown suppressed cervical tumor growth via miR-485-5p/JUND axis in vivo. Xenograft models were established by injecting sh-NC or sh-SNHG7 transfected HeLa cells into in the flanks nude mice, then (A) tumor volume was measured weekly; (B) tumor weight was measured 5 weeks later; (C) miR-485-5p mRNA expression was detected by qPCR; (D) JUND protein expression was detected by Western blot; (E) Ki67 was detected by IHC. **P<0.01 versus sh-NC. **P<0.01.
partially abolished the inhibitory effects of si-SNHG7 on CC cell viability, migration and invasion, indicating that SNHG7 promoted CC cell progression via miR-485-5p.

MiRNAs downregulate target mRNA expression through binding to 3’UTR seed region, which is the basis of IncRNA-miRNA-mRNA regulating network. Yu et al revealed that HCP5 enhanced CC development through modulating miR-15a/MACC1 axis. Gao et al demonstrated that SBF2-AS1 driven CC by regulating FOXM1 via miR-361-5p. Ji et al showed that SNHG14 promoted CC development via acting as ceRNA for miR-206 to regulate YWHAZ. In this work, we found jun D proto-oncogene (JUND) as the target gene of miR-485-5p using TargetScan software. JUND, a member of the AP-1 family, functioned as tumor-promoting roles in some cancer types. For example, Elliott B reported that JUND was a crucial modulator in prostate cell cycle progression and thus promoting cancer development. Ishikawa et al found that Butein ameliorated adult leukemia via inhibiting JUND expression. Cheng et al also revealed that JUND was involved in cancer stemness and drug resistance in hepatocellular carcinoma. Herein, luciferase assay confirmed the direct interaction between miR-485-5p and JUND. Moreover, SNHG7 positively regulated JUND via miR-485-5p, suggesting the SNHG7/miR-485-5p/JUND axis involved in the mechanism. Based on the characteristics that one IncRNA can sponge multi-miRNAs, and one miRNA also targets different genes, further study is needed to elucidate whether IncRNA-SNHG7 can regulate other miRNAs and proteins. This would help to understand the underlying mechanism of SNHG7-regulated cervical cancer progression better.

**Conclusions**

In conclusion, IncRNA SNHG7 played as an oncogene which promoted cell proliferation, migration and invasion by sponging miR-485-5p and thereby upregulating JUND (Figure 8). SNHG7.miR-485-5p/JUND axis might provide a novel approach for CC treatment.

**Ethical Approval**

Written informed consent was obtained from every involved patient. The Ethics Committee of the First Affiliated Hospital of Guangxi Medical University approved this study. All experiments were performed following the guidelines and regulations of Ethics Committee of the First Affiliated Hospital of Guangxi Medical University.

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**Disclosure**

The authors declare that they have no competing interests in this work.

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