

LncSNHG3/miR-139-5p/BMI1 axis regulates proliferation, migration, and invasion in hepatocellular carcinoma

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Objective: Emerging evidence has revealed that lncRNA small nucleolar RNA host gene 3 (SNHG3) is involved in cell proliferation, migration, and invasion in various tumors. However, the underlying molecular mechanism of SNHG3 in hepatocellular carcinoma (HCC) is still not fully explored.

Methods: Quantitative reverse transcriptase PCR was employed to detect the expression of SNHG3, miR-139-5p, and BMI1. Colony assay and MTT assay were used to detect the proliferation. Transwell assay was introduced to measure the migration and invasion ability. Bioinformatics analysis and luciferase reporter assay were used to confirm the relationship between SNHG3, miR-139-5p, and BMI1. An animal experiment was adopted to detect the function of SNHG3 in vivo.

Results: SNHG3 and BMI1 were upregulated in HCC, while miR-139-5p was downregulated. Knockdown of SNHG3 or BMI1 and overexpression of miR-139-5p could inhibit cell proliferation, migration, and invasion in HCC. miR-139-5p was a target of SNHG3 and BMI1 was a direct target mRNA of miR-139-5p. Silencing SNHG3 could impair the tumor progression in vivo.

Conclusion: The lncRNA SNHG3/miR-139-5p/BMI1 axis plays an important role in cell proliferation, migration, and invasion in HCC.

Keywords: SNHG3, miR-139-5p, BMI1, hepatocellular carcinoma

Introduction

Hepatocellular carcinoma (HCC) is commonly diagnosed and identified as a leading cause of cancer death. According to the cancer statistics in China in 2015,¹ the incidence was about 466,100 and mortality was 422,100 annually, which makes it the third most deadly cancer. In the USA, it was estimated that about 41,000 cases and 29,000 deaths were caused by HCC in 2017.² Although great efforts were made in HCC treatment and they revealed that many genes were involved in the progression, migration, invasion, and metastasis of HCC, the molecular mechanism of HCC progression is still poorly understood. Therefore, it is important to explore the underlying mechanism of HCC progression.

Recent studies have highlighted that long non-coding RNAs (lncRNAs) are related to cancer progression. Long non-coding RNAs are endogenous cellular RNAs which are more than 200 nucleotides in length. LncRNAs may regulate the cell processes by mediating transcription or post-transcription. It was reported that lncRNAs could interact with DNA, RNA, and protein to play the regulatory roles. Their dysregulation may lead to many diseases including cancers. For example, lncRNA p10247 overexpression is

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correlated with metastasis in breast cancer.³ Downregulated lncRNA-HOTAIR suppressed colorectal cancer proliferation, invasion, and migration.⁴ LncRNA MIR4435-2HG promoted lung cancer progression.⁵ LINC00844 regulates cell migration and invasion in prostate cancer.⁶ LncRNA TBILA may promote the epithelial–mesenchymal transition of non-small cell lung cancer.⁷ Besides the above-mentioned cancer types, lncRNAs also play roles in HCC. LncRNA FLVCR-AS1,⁸ HNF1A-AS1,⁹ HOXA11-AS,¹⁰ and lncRNA OGFRP1¹¹ were upregulated, while XLOC006926,¹² CASC2,¹³ and lncRNA NKILA¹⁴ were downregulated in HCC. However, there are still many lncRNAs involved in HCC tumorigenesis and progression that need to be explored.

It has been underscored that lncRNAs act as competing endogenous RNAs for miRNAs in biological processes. miRNAs are small non-coding RNAs which are about 18–25 nucleotides in length. Previous studies demonstrated that miRNAs may mediate gene expression by binding the 3'UTR of specific mRNA¹⁵ and showed that miRNAs were involved in many physiological and pathological processes such as tumors. For instance, miR-214 suppressed ovarian cancer,¹⁶ serum miR-196a could be a biomarker in gastric cancer,¹⁷ miR-34a could inhibit liver cancer cell growth,¹⁸ and miRNA-139 downregulation was associated with HCC risk and survival.¹⁹ But the function of miRNA in HCC is not fully understood.

BMI1 encodes a ring finger protein which is a major component of the polycomb group complex 1. BMI1 and other components are involved in multiple regulatory mechanisms through chromatin remodeling. Some studies reported that BMI1 is an oncogene and aberrant expression in numerous cancers including non-small cell lung cancer,²⁰ colorectal cancer,²¹ breast cancer,²² endometrial cancer,²³ and HCC.²⁴ Yet, the detail mechanism of how BMI1 was regulated in cancers is unclear.

In the present study, we aimed to explore the biological role of lncRNA small nucleolar RNA host gene 3 (SNHG3) in HCC. We found that SNHG3 could negatively regulate miR-139-5p. Meanwhile, BMI1 was a downstream target mRNA of miR-139-5p. So, we proposed that SNHG3 may regulate proliferation, migration, and invasion via regulating miR-139-5p and BMI1 in HCC.

Materials and methods

Patient samples

The ethical approval was obtained from the Research Ethics Committee of the First Affiliated Hospital of Sun-Yat-sen

University (Guangzhou, China). Thirty-one specimens of cancerous tissue and 31 specimens of paracancerous tissue were taken from the HCC patients, and all patients signed the informed consent form, and this study was performed in accordance with the ethical standards detailed in the Declaration of Helsinki. No patients had undergone chemotherapy or radiotherapy before surgery. The samples were stored in liquid nitrogen for subsequent laboratory testing.

Cell culture

HCC cell lines Hep3B and HepG2 and immortalized human liver cell line L-02 were obtained from RiboBio Co. (Guangzhou, China). Cells were cultured in Dulbecco's Modified Eagle's Medium DMEM (Gibco, Carlsbad, CA, USA) with 10% FBS (HyClone, Logan, UT, USA) and 1% penicillin and streptomycin (Sigma-Aldrich, St Louis, MO, USA) at 37°C with 5% of CO₂. Medium was changed every 2 days.

Cell transfection

For transient transfection, siSNHG3, miR-139-5p, pcDNA-SNHG3, miR-139-5p inhibitor, siBMI1, and the corresponding negative control were purchased from GenePharma Co., Ltd. (Shanghai, China). All oligos and plasmids were transfected into Hep3B or HepG2 cells using Lipofectamine 3000 (Invitrogen, Carlsbad, CA, USA) according to the manufacturer's instructions.

For stable transfection, shRNA targeting lnc SNHG3 and negative control were synthesized by GenePharma Co., Ltd. (Shanghai, China). These fragments were inserted into the lentiviral vector pLVXshRNA2 (Clontech, Mountain view, CA, USA). Then the lentiviral vectors were transfected into HEK293T cells with the Lenti-X HTX packaging system (Clontech) according to the protocol. The medium containing lentivirus was collected at 48 hours after transfection. Then the lentiviruses were added to the medium containing HepG2 cells. After that, the transfected cells were sorted with FITC fluorescence using a BD FACSMelody Cell Sorter (BD Biosciences, San Jose, CA, USA).

Quantitative real-time PCR (qRT-PCR)

Total RNA was extracted from tissues and cells using TRIzol (Invitrogen, Carlsbad, CA, USA) and RNA concentration was detected by a NanoDropND-1000 spectrophotometer (Thermo Fisher Scientific, Rockford, IL, USA). cDNA was reverse transcribed using the TaqMan miRNA Reverse Transcription Kit (Applied Biosystems, Foster City, CA, USA) for miRNA and Reverse

Transcription Reagents (Applied Biosystems) for mRNA and lncRNA. SNHG3, miR-139-5p, and BMI1 expression were determined using SYBR[®] Green (Promega, Madison, WI, USA) according to the manufacturer's protocol. A U6 small nuclear RNA (snRNA) was used as a reference gene for miR-139-5p, and 18S rRNA was used as a reference gene for SNHG3 and BMI1. Fluorescence was detected in the iQTM5 Multicolor Real-Time PCR Detection System (Bio-Rad, Hercules, CA, USA). The relative expression of SNHG3, miR-139-5p, and BMI1 was calculated by the $2^{-\Delta\Delta Ct}$ method. Primer sequences: SNHG3, forward 5'-AGTGGTCGCTTCTTCTCCTTG-3' and reverse 5'-GATTGTCAAACCTCCCTGTTA-3'; miR-139-5p, forward 5'-ACACTCCAGCTGGGTCTACAGTGCACGTG-3' and reverse 5'-CTCAACTGGTGTGCGTGGA-3'; BMI1, forward 5'-GGTACTTCATTGATGCCACAAC-3' and reverse 5'-CTGCTGGGCATCGTAAGTATC-3'; 18S rRNA, forward 5'-GGCCCTGTAATTGGAATGAGTC-3' and reverse 5'-CCAAGATCCAACACTACGAGCTT-3'; U6, forward 5'-GTTGACATCCGTAAGACC-3' and reverse 5'-GGAGCCAGGGCAGTAA-3'.

Western blot

Cells were washed with Tris-buffered saline (TBS), and then added to RIPA buffer and maintained on ice for 30 minutes, accompanied with vortexing 30 seconds per 10 minutes. They were then centrifuged for 15 minutes at 4°C with 12,000 rpm and the supernatant was transferred to a new tube for next experiments. The protein concentrations were measured using Nanodrop2000 (Thermo Fisher Scientific, San Jose, CA, USA). Then the loading buffer was added to the protein solution and heated to 95°C for 5 minutes. Proteins (10 µg) were separated using SDS-PAGE and then transferred to polyvinylidene fluoride membranes (PVDF, Millipore, Bedford, MA, USA). After being blocked with 5% dried skimmed milk in TBS, the membranes were incubated with specific primary antibodies including anti-BMI1 (1:100 solution, and all of these antibodies were purchased from Santa Cruz Biotechnology Inc., CA, USA) and anti-β actin antibody (1:2,000 dilution, Santa Cruz Biotechnology Inc.) at 4°C overnight. The PVDF membrane was washed five times for 10 minutes each with Tris-buffered saline with Tween-20 (TBST), following incubation with horseradish peroxidase (HRP)-conjugated anti-mouse IgG secondary antibody (1:2,000 dilution, Santa Cruz Biotechnology Inc.) at room temperature for 1 hour. After being washed with TBST five times, the protein signals were detected using Pierce[™] ECL Western blotting substrate (ThermoFisher Scientific).

Colony formation assay

Transfected cells were trypsinized after 24 hours, counted, and replated at a density of 500 cells/6 cm dish. Ten days later, colonies resulting from the surviving cells were fixed with 3.7% methanol, stained with 0.1% crystal violet, and counted. Colonies containing at least 50 cells were scored. Each assay was performed three times.

MTT assay

Cell viability was measured with MTT assay kit (Sigma-Aldrich) according to the manufacturer's protocol. In brief, cells (2×10^3) were seeded into 96-well plates (Corning Costar, Corning, NY, USA). Then 30 µL serum-free media with MTT solution was added into each well and incubated for 4 hours at 37°C. After discarding the media, 150 µL MTT solvent (4 mM HCl, 0.1% NP40 in isopropanol) was added into each well and then incubated for 3 hours at 37°C. Then the absorbance (A) value of each well was measured at OD=450 nm using the Microplate Reader (MG LABTECH, Durham, NC, USA).

Cell invasion and migration

Cell invasion analysis was performed using Transwell chamber with Matrigel (BD Biosciences, San Jose, CA, USA). Cell migration was displayed using a Transwell chamber without Matrigel. 1×10^5 cells were cultured in non-serum medium in the Transwell upper chamber. The Transwell lower chamber was added in DMEM medium with 10% FBS. After incubated for 24 hours at 37°C, cells on the upper chamber membrane were moved. Then the membrane was fixed with 90% ethyl alcohol and was stained with 0.1% crystal violet for 30 minutes. The number of cell migrations and invasions were counted using a phase contrast microscope.

Animal experiments

Animal experiments were carried out according to the National Institutes of Health guidelines to the Care and Use of Laboratory Animals of the First Affiliated Hospital of Sun-Yat-sen University. shSNHG3 and their negative control (shNC) were stably transfected into HepG2 cells using lentiviruses. Then 5×10^6 cells were subcutaneously injected in the flank of female nude mice (Beijing Vital River Laboratory Animal Technology Co., Ltd., China). There were five nude mice in each group. Tumor volumes were measured every 7 days until 35 days. All mice were sacrificed to collect tumor xenografts and tumor xenografts were weighted.

Immunohistochemistry

The tumor tissue from mice was fixed with 12 mL 4% paraformaldehyde (PFA) in PBS followed by being embedded with paraffin. Then the tissues were made into 4 μ M sections and de-waxed in xylene, rehydrated in ethanol. Peroxidase blocking reagent was used to quench the endogenous peroxidase activity for 10 minutes. The sections were incubated overnight at 4°C with primary antibody (BMI1, ab38295, Abcam, Cambridge, MA, USA). Then washed with PBS and treated with HRP-conjugated secondary antibody at room temperature for 1 hour and developed with liquid diaminobenzidine substrate chromogen system.

Statistical analysis

The data are presented as mean \pm SD (standard deviation) from three biological replicated experiments. The results of analysis were shown and plotted using GraphPad Prism 7.0 (GraphPad Software, San Diego, CA, USA). All group comparisons were carried out using Student's *t*-test. *P*-values less than 0.05 were regarded as statistically significant.

Results

SNHG3 was upregulated and miR-139-5p was downregulated in HCC

Firstly, we obtained HCC tumor tissues and adjacent tissues from 31 patients without any treatment except surgery. qRT-PCR was used to detect the expression of SNHG3 and the results showed that SNHG3 was upregulated significantly in HCC tissues than in normal tissues (Figure 1A). Then we continued to check SNHG3 expression in HCC cell lines. Consistent with that in tissues, SNHG3 was upregulated in Hep3B and HepG2 cells (Figure 1B). Meanwhile, we also checked miR-139-5p expression in HCC tissues and cell lines. The qRT-PCR results showed that miR-139-5p was downregulated in HCC tissues (Figure 1C), Hep3B and HepG2 cells (Figure 1D). Our finding suggested that upregulation of SNHG3 and downregulation of miR-139-5p might be involved in HCC.

Knockdown of SNHG3 inhibited cell proliferation, migration, and invasion in HCC

To better understand the function of SNHG3 in HCC, we downregulated SNHG3 via small interfering RNA in Hep3B

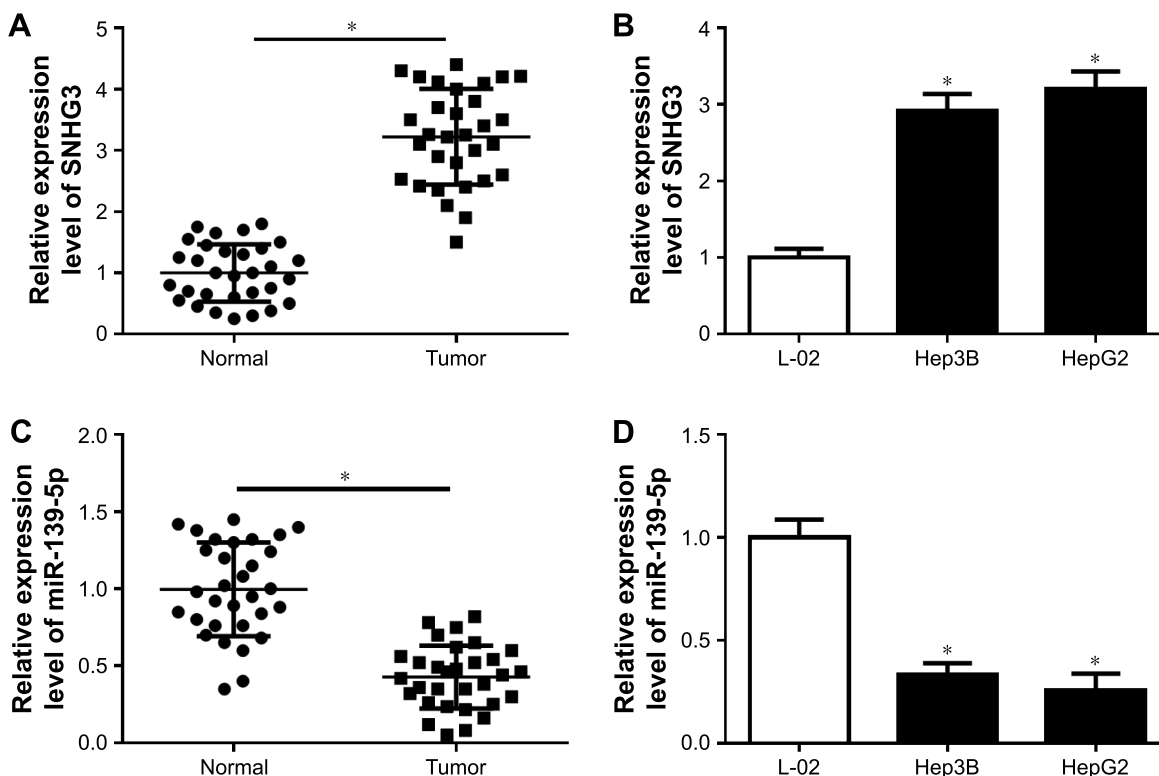


Figure 1 Expression of SNHG3 and hsa-mir-139-5p in HCC tissue and cells. (A) SNHG3 expression in normal and HCC tissues. (B) SNHG3 expression in L-02, Hep3B, and HepG2 cells. (C) miR-139-5p expression in normal and HCC tissues. (D) miR-139-5p expression in L-02, Hep3B, and HepG2 cells. **P*<0.05.

Abbreviations: HCC, hepatocellular carcinoma; SNHG3, small nucleolar RNA host gene 3.

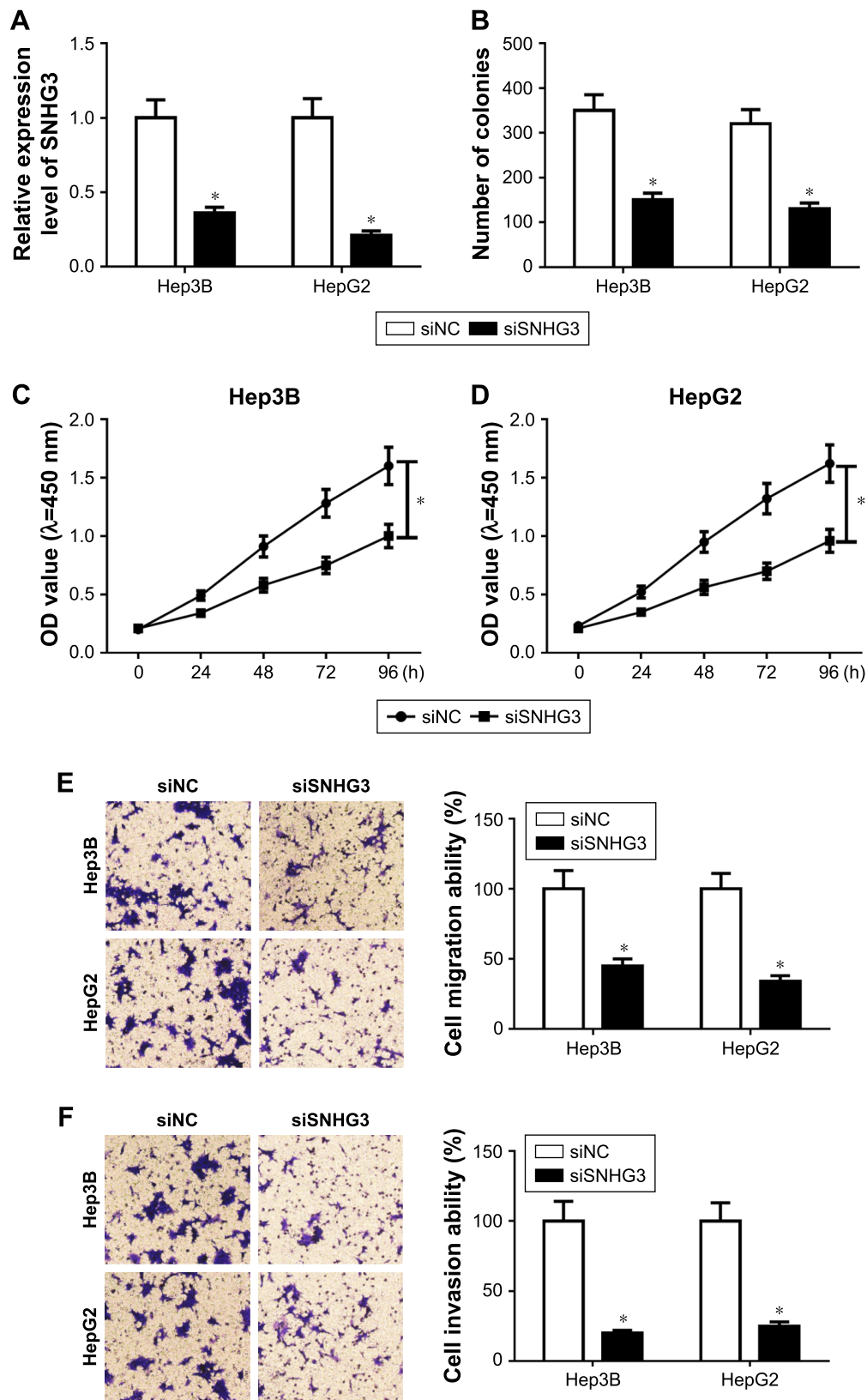


Figure 2 Effect of lncSNHG3 on the cell proliferation, migration and invasion in HCC. (A) SNHG3 knockdown in Hep3B and HepG2 cells. (B) Colony number of Hep3B and HepG2 transfected with siSNHG3. (C and D) MTT assay detected the proliferation of Hep3B (C) and HepG2 (D) after SNHG3 knockdown. (E) Cell migration analysis of Hep3B and HepG2 transfected with siNC and siSNHG3. Magnification 200 \times . (F) Cell invasion analysis of Hep3B and HepG2 transfected with siNC and siSNHG3. Magnification 200 \times . *P<0.05.

Abbreviations: HCC, hepatocellular carcinoma; lnc, long non-coding; SNHG3, small nucleolar RNA host gene 3.

and HepG2 cells. After 48 hours of transfection, the knock-down efficiency was checked and SNHG3 was indeed down-regulated in Hep3B and HepG2 cells (Figure 2A). Then a colony formation assay was performed to detect the proliferation, and the colony number was significantly decreased after SNHG3 silencing (Figure 2B). Meanwhile, an MTT assay confirmed the inhibitory effect of SNHG3 in cell proliferation in HCC (Figure 2C and D). Next, we checked the function of SNHG3 in migration and invasion in HCC, and Transwell assay suggested that knockdown of SNHG3 significantly inhibited the migration (Figure 2E) and invasion (Figure 2F) of Hep3B and Hep2G cells.

Overexpression of miR-139-5p impaired cell proliferation, migration, and invasion in HCC

Considering that miR-139-5p was downregulated in HCC tissues and cells, we overexpressed miR-139-5p in Hep3B and HepG2 cells (Figure 3A). Both colony formation assay (Figure 3B) and MTT assay (Figure 3C and D) suggested that miR-139-5p overexpression inhibited the cell proliferation. In addition, the migration (Figure 3E) and invasion (Figure 3F) ability were impaired after miR-139-5p overexpression.

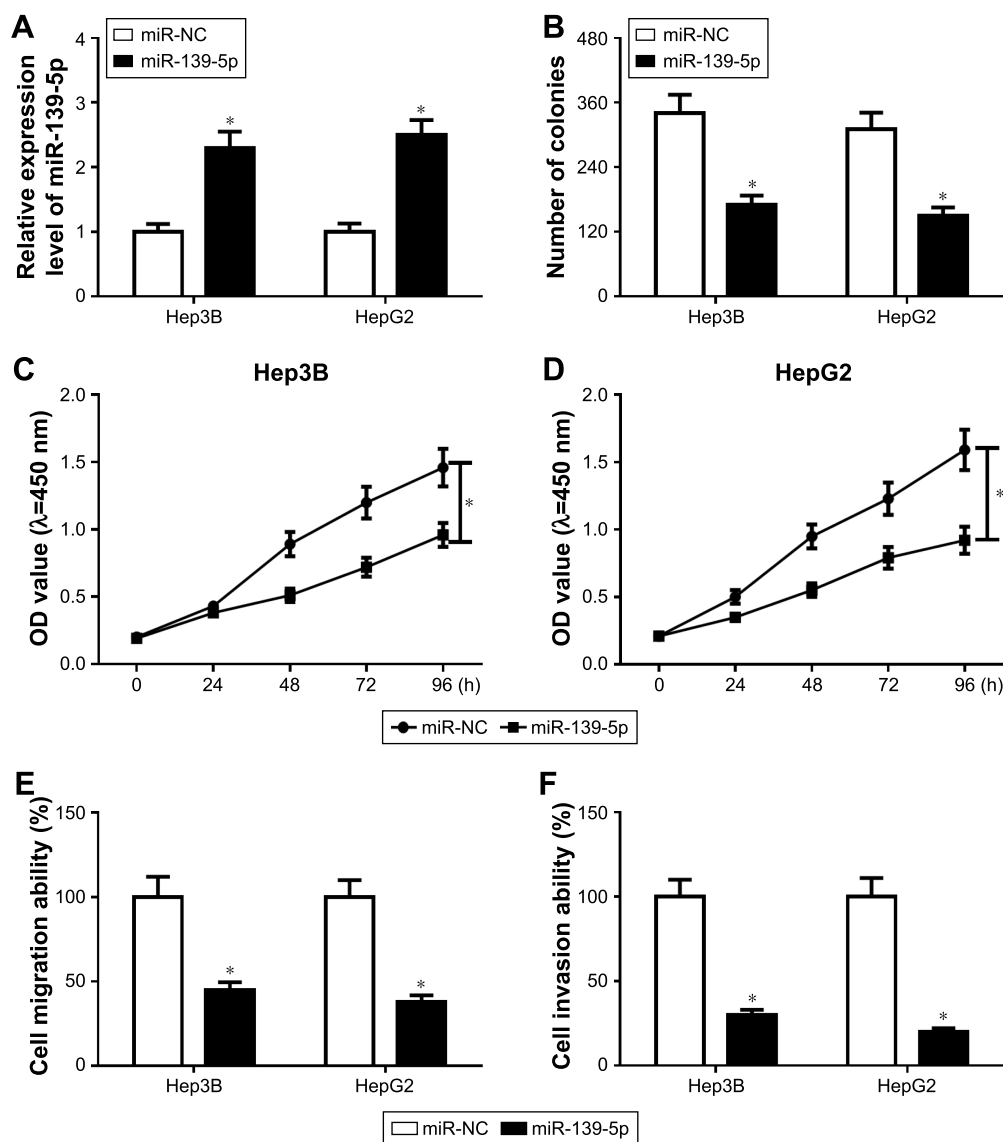


Figure 3 Effect of miR-139-5p on the cell proliferation, migration, and invasion in HCC. (A) miR-139-5p overexpression in Hep3B and HepG2 cells. (B) Colony number of Hep3B and HepG2 transfected with miR-139-5p. (C and D) MTT assay detected the proliferation of Hep3B (C) and HepG2 (D) after miR-139-5p overexpression. (E) Cell migration analysis of Hep3B and HepG2 transfected with miR-NC and miR-139-5p. (F) Cell invasion analysis of Hep3B and HepG2 transfected with miR-NC and miR-139-5p. * $P < 0.05$.

Abbreviation: HCC, hepatocellular carcinoma.

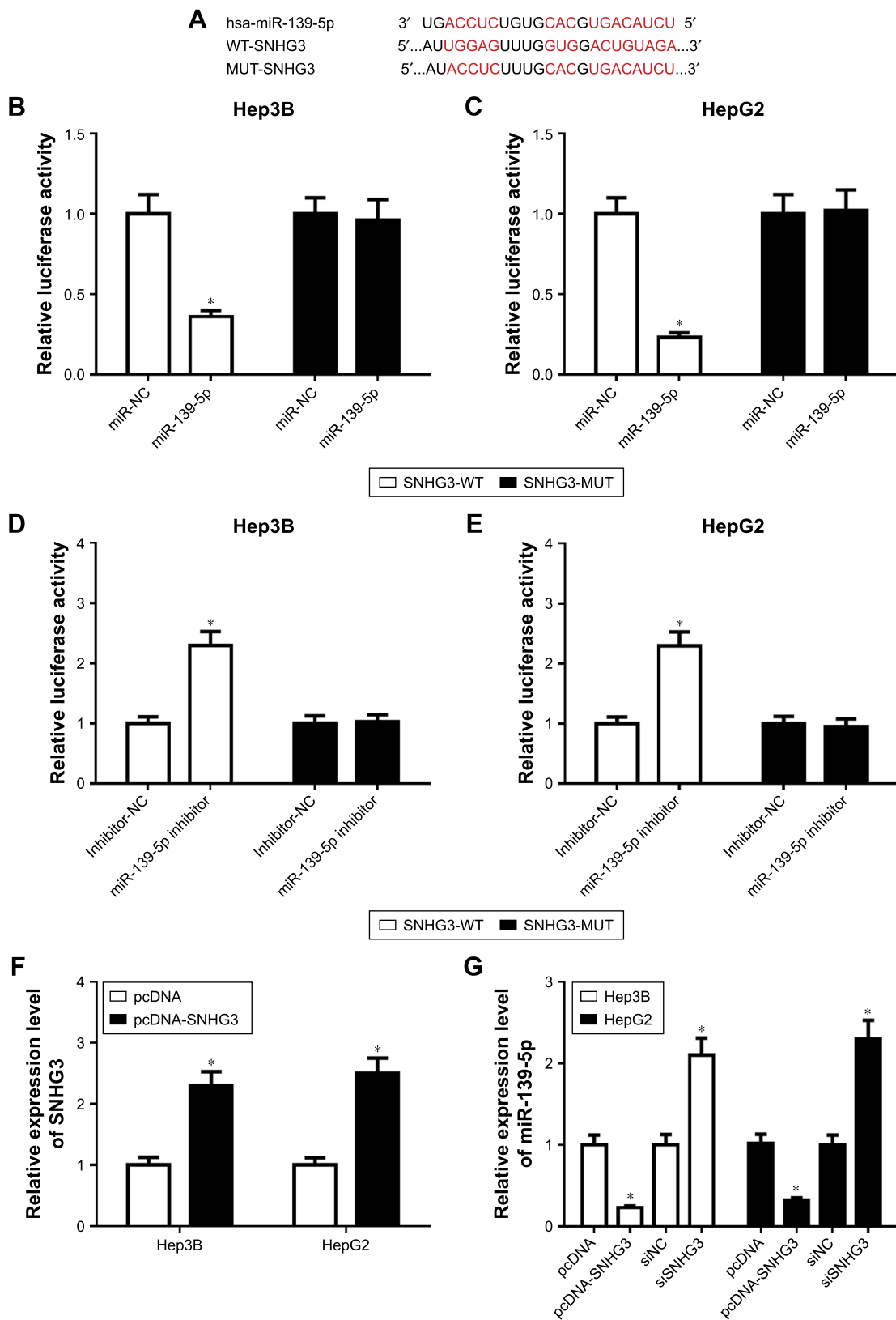


Figure 4 miR-139-5p was a target of SNHG3. (A) The relationship between SNHG3 and miR-139-5p was predicted by starbase. (B and C) SNHG3-WT or SNHG3-MUT was cotransfected into Hep3B (B) and HepG2 (C) cells with miR-139-5p or their negative controls. (D and E) SNHG3-WT or SNHG3-MUT was cotransfected into Hep3B (D) and HepG2 (E) cells with miR-139-5p inhibitor or their negative controls. (F) SNHG3 overexpression in Hep3B and HepG2 cells. (G) miR-139-5p expression in pcDNA, pcDNA-SNHG3, siNC, and siSNHG3. * $P < 0.05$.

Abbreviation: SNHG3, small nucleolar RNA host gene 3.

miR-139-5p was a target of SNHG3

Next, starBase was used to predict the relationship between SNHG3 and miR-139-5p. Bioinformatics analysis revealed that there were complementary sequence between lncSNHG3 and miR-139-5p (Figure 4A). Then we constructed luciferase report vector containing wildtype and mutant SNHG3. The luciferase activity was decreased when wildtype SNHG3 and miR-139-5p were transfected into Hep3B (Figure 4B) and HepG2 (Figure 4C) but not with mutant SNHG3. To further confirm the binding between SNHG3 and miR-139-5p, we introduced miR-139-5p inhibitor and the luciferase activity was upregulated about 2-fold when

cotransfected with wildtype SNHG3 in Hep3B (Figure 4D) and HepG2 cells (Figure 4E). Subsequently, we overexpressed SNHG3 in Hep3B and HepG2 cells (Figure 4F) and miR-139-5p expression was inhibited after SNHG3 overexpression and increased when SNHG3 was silenced (Figure 4G). The results demonstrated that miR-139-5p was a target of SNHG3.

BMI1 was a direct target of miR-139-5p

Then, to investigate the mechanism of miR-139-5p in HCC, the starBase database was used to predict the downstream target. Luciferase reporter plasmids of wildtype BMI1 and mutant BMI1 were constructed (Figure 5A). Cotransfection

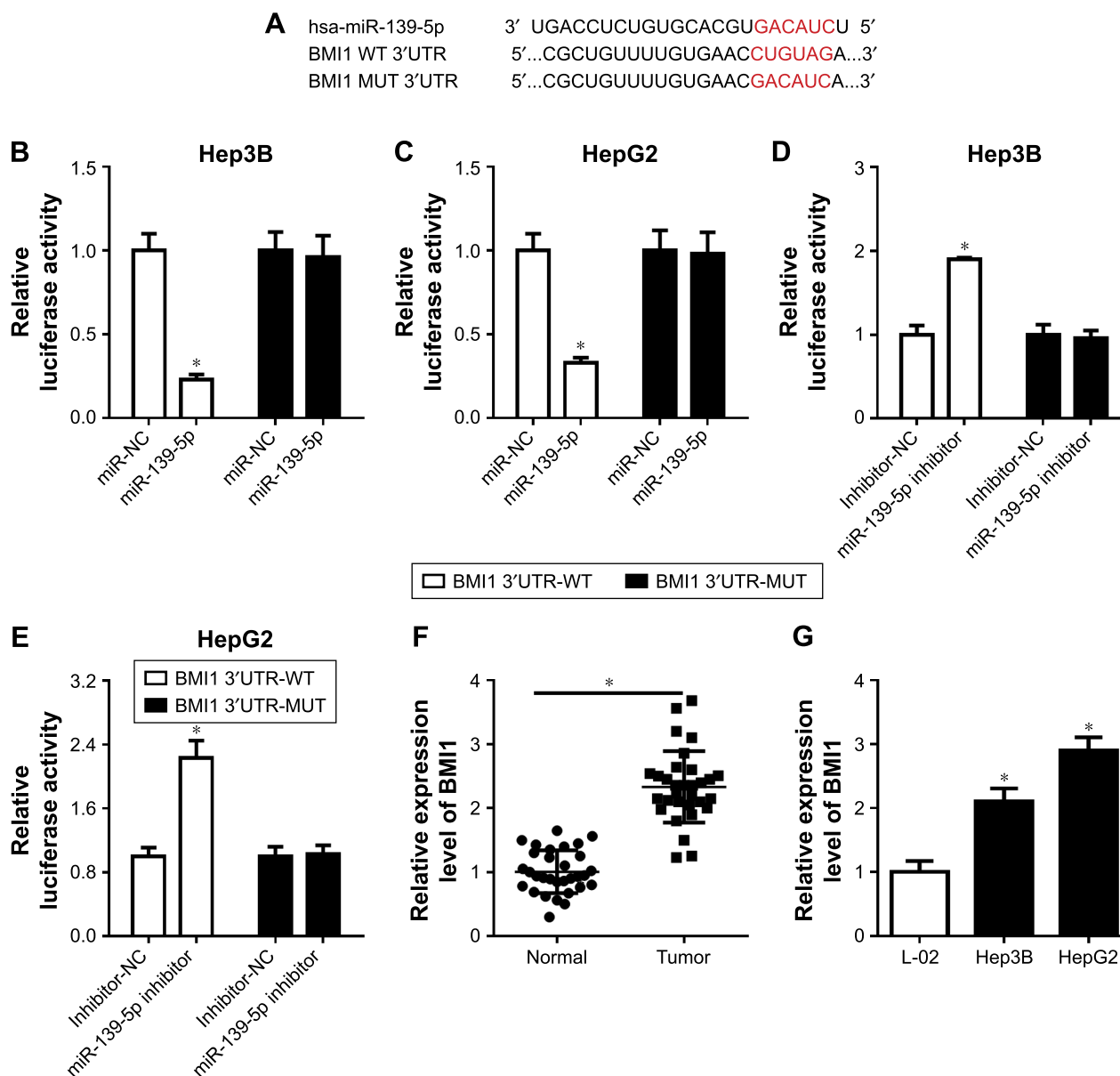


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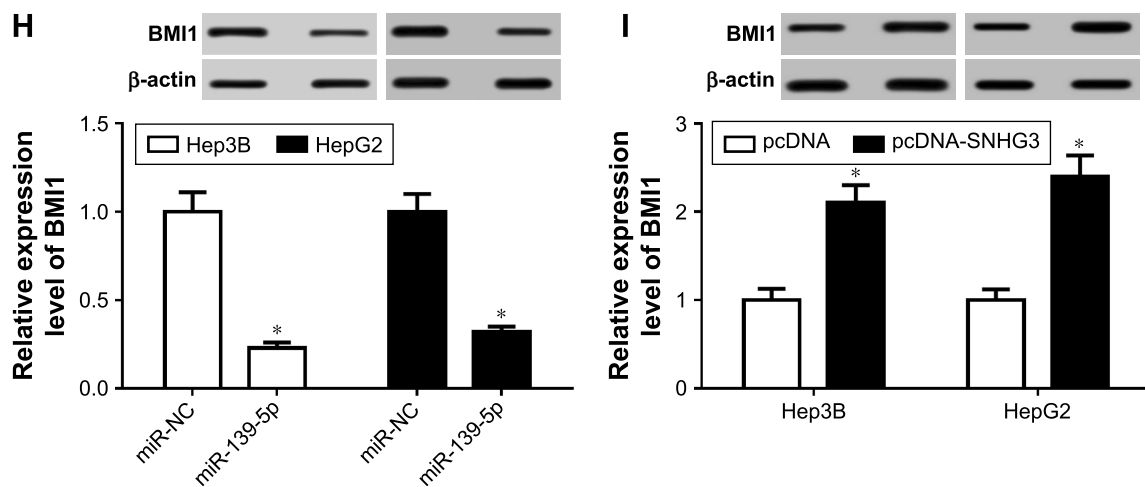


Figure 5 BMI1 was a direct target of miR-139-5p. (A) The relationship of miR-139-5p and BMI1 was predicted by starbase. (B and C) WT-BMI1 or MUT-BMI1 was cotransfected into Hep3B (B) and HepG2 (C) cells with miR-139-3p or their negative controls. (D and E) WT-BMI1 or MUT-BMI1 was cotransfected into Hep3B (D) and HepG2 (E) cells with miR-139-3p inhibitor or their negative controls. (F) BMI1 expression in HCC tissues. (G) BMI1 expression in LC-02, Hep3B, and HepG2 cells. (H) BMI1 expression under the condition of miR-139-5p overexpression in Hep3B and HepG2 cells. (I) BMI1 expression under the condition of SNHG3 overexpression in Hep3B and HepG2 cells. * $P < 0.05$. **Abbreviations:** HCC, hepatocellular carcinoma; SNHG3, small nucleolar RNA host gene 3.

of the luciferase reporter plasmid containing BMI1 3'UTR-WT with miR-139-5p in Hep3B and HepG2 resulted in a decreased luciferase activity, but not with mutant BMI1 (Figure 5B and C). The miR-139-5p inhibitor could increase the luciferase activity in Hep3B (Figure 5D) and HepG2 cells (Figure 5E).

Next, we checked the BMI1 expression in HCC tissues and cell lines. The mRNA expression level was upregulated in HCC tumor (Figure 5F) and cell lines (Figure 5G). Afterward, we overexpressed miR-139-5p in Hep3B and HepG2 cells and found that BMI1 was decreased (Figure 5H). Meanwhile, BMI1 expression was increased after SNHG3 overexpression

(Figure 5I). These findings indicated that BMI1 can serve as a direct target of miR-139-5p.

BMI1 knockdown inhibited cell proliferation, migration, and invasion in HCC

To better understand the function of BMI1 in HCC, we downregulated BMI1 in Hep3B and HepG2 cells (Figure 6A and B). Cell proliferation was detected by colony formation assay (Figure 6C) and MTT assay (Figure 6D and E). The results showed that proliferation was inhibited by siBMI1. Moreover, cell migration (Figure 6F) and invasion

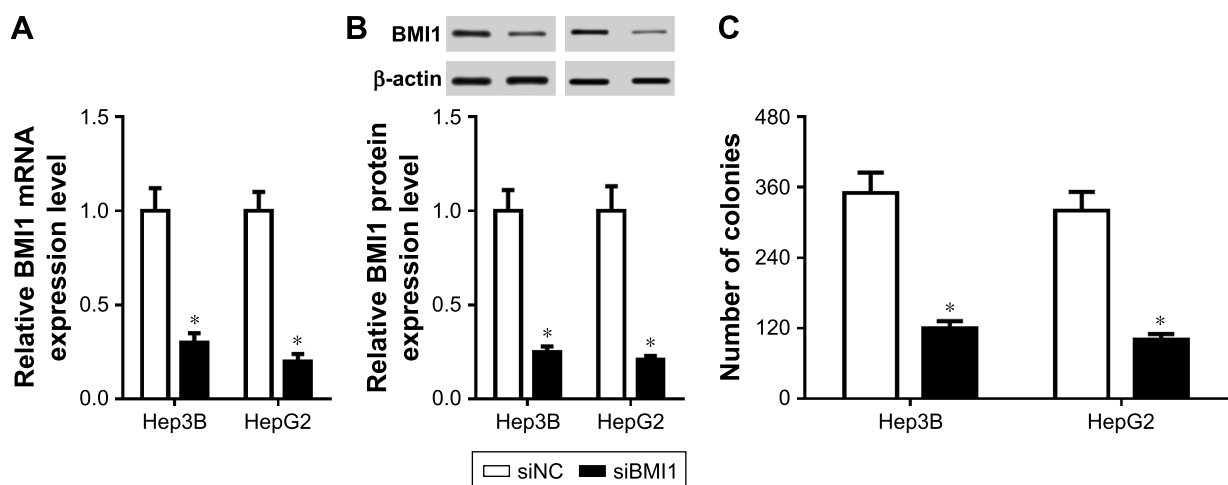


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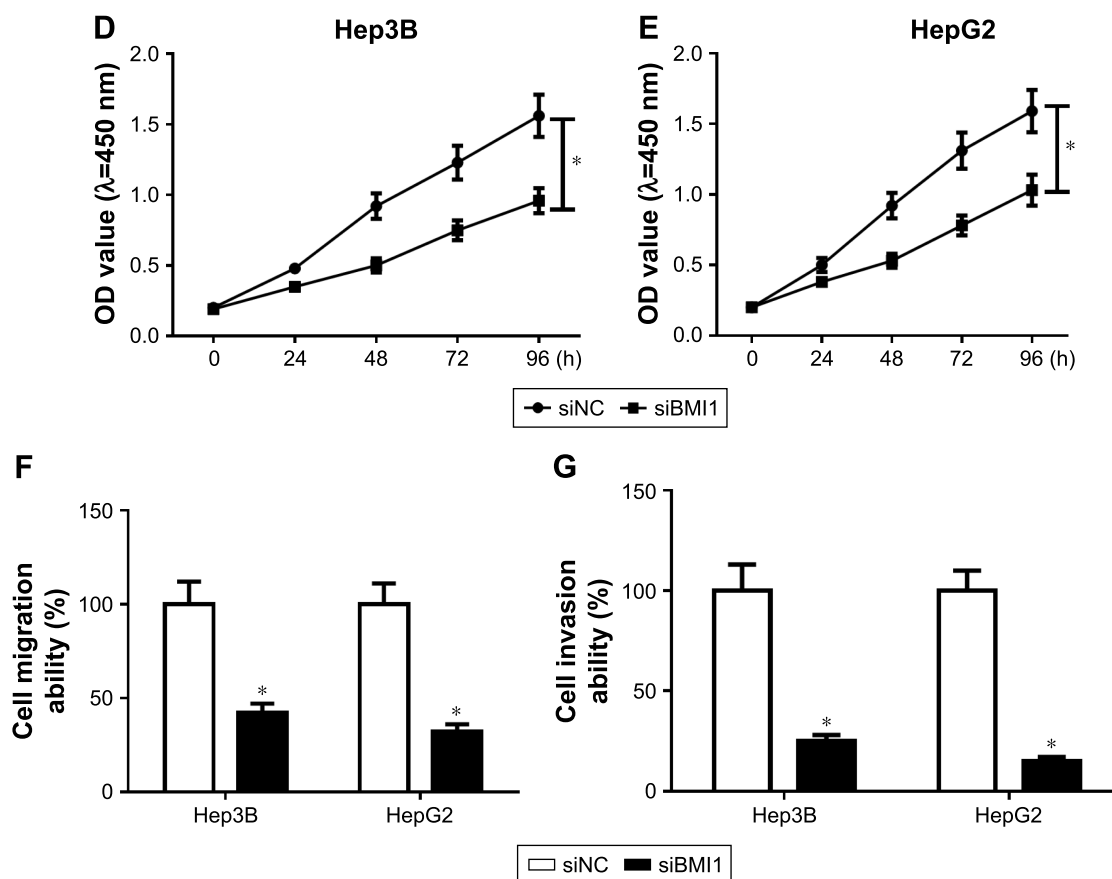


Figure 6 Effect of BMI1 on the cell proliferation, migration and invasion in HCC. (A and B) BMI1 knockdown in Hep3B and HepG2 cells. (C) Colonies number of Hep3B and HepG2 transfected with siBMI1. (D and E) MTT assay detected the proliferation of Hep3B (D) and HepG2 (E) after BMI1 knockdown. (F) Cell migration analysis of Hep3B and HepG2 transfected with siNC and siBMI1. (G) Cell invasion analysis of Hep3B and HepG2 transfected with siNC and siBMI1. *P<0.05.

Abbreviation: HCC, hepatocellular carcinoma.

(Figure 6G) ability were inhibited by BMI1 silencing in Hep3B and HepG2 cells.

SNHG3 knockdown restrained the tumor progression in vivo

To evaluate whether SNHG3 is involved in HCC progression in vivo, a HepG2 cell nude mouse xenograft model was used. The total nude mice were separated into two groups, one group was stably transfected with siNC and the other with siSNHG3. The tumor size was measured every 7 days until the sacrifice on day 35. Tumor growth was significantly inhibited by SNHG3 knockdown from day 21 (Figure 7A). On day 35, we measured the weight, and SNHG3 silencing obviously inhibited the tumor growth (Figure 7B). Meanwhile, we checked the SNHG3, miR-139-5p, and BMI1 expression in the tumor. qRT-PCR demonstrated that SNHG3 (Figure 7C) were downregulated and miR-139-5p (Figure 7D) was upregulated in the tumor. In addition, immunohistochemical

assay revealed that BMI1 protein level was also decreased (Figure 7E). These data suggested that downregulation of SNHG3 could inhibit HCC development in vivo.

Discussion

Over the past few decades, researchers claimed that non-coding transcripts are transcriptional noise.²⁵ However, in recent years, whole genome and transcriptome sequencing suggested that non-coding RNAs (ncRNAs), including long non-coding RNAs (lncRNAs), may be involved in various cell processes,²⁶ for example, epigenetics, alternative splicing, RNA decay, and translation.^{27–29} Given that massive lncRNAs have been discovered, the detailed function of these lncRNAs needs to be elucidated. Emerging evidence demonstrated that dysregulated lncRNAs were correlated with various cancers,³⁰ for example, anti-NOS2A,³¹ BC200,³² HULC,³³ H19,³⁴ MALAT1,³⁵ HOTAIR,³⁶ etc. Previous reports demonstrated that SNHG3 play an important role in various cancer development. SNHG3 enhanced

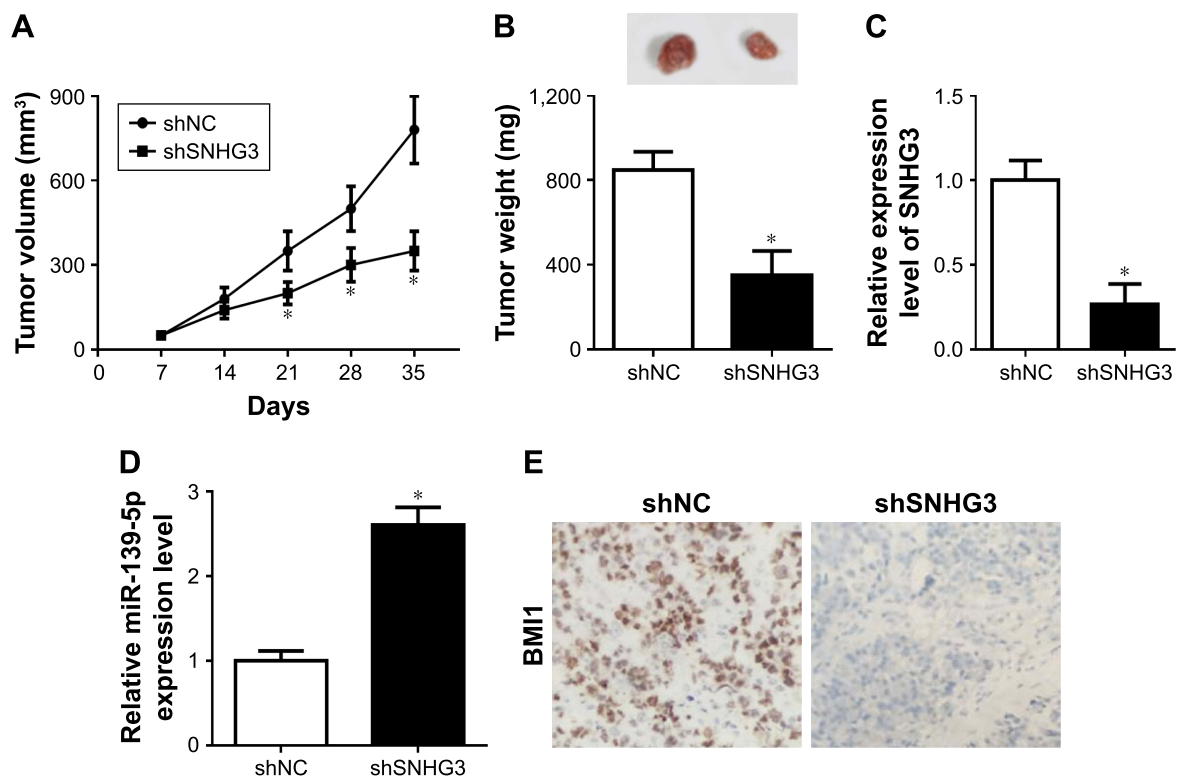


Figure 7 Down-regulation of SNHG3 repressed tumor progression in vivo. (A) Tumor volume was measured every 7 days after HepG2 stably transfected with siNC or siSNHG3 xenograft. (B) Tumor weight was measured on day 35. (C–F) SNHG3 (C), miR-139-5p (D), BMI1 mRNA (E), and BMI1 protein, magnification 200 \times . * $P < 0.05$. **Abbreviation:** SNHG3, small nucleolar RNA host gene 3.

the malignant process of colorectal cancer.³⁷ SNHG3 promoted proliferation and invasion in breast cancer³⁸ and lung adenocarcinoma.³⁹ In HCC, it was reported that SNHG3 was significantly upregulated and highly correlated with tumor size and relapse.⁴⁰ In our present work, we found that SNHG3 was upregulated in HCC tissues compared with normal tissues, which was consistent with previous data. Dissimilarly, we knocked down the expression of SNHG3 using RNAi and found that the proliferation, migration, and invasion were inhibited in HCC cell lines in vitro. Meanwhile, we used nude mice and xenograft assay to verify the inhibitory function of SNHG3 in vivo. So we hypothesized that SNHG3 was involved in the progression of HCC. However, another study has demonstrated SNHG3 silencing in HCCLM3, which is a highly metastatic HCC cell line, and overexpression in PLC/PRF/5, which is one lowly metastatic HCC cell line which didn't affect the cell proliferation.⁴¹ As shown in our supplemental material, we knocked down the expression of SNHG3 in HCCLM3, and the cell proliferation was inhibited (Figures S1–S4). We proposed that maybe the culture condition, knockdown efficiency, etc., were different in our two references, which led to distinct results. Besides, in other cancers

such as breast cancer and lung cancer,^{38,39} SNHG3 knockdown inhibited cell proliferation.

Many studies revealed that miR-139-5p is involved in the cancer development, such as breast cancer,⁴² colorectal cancer, myeloid leukemia,⁴⁴ etc. We found that miR-139-5p was downregulated in HCC. Through the bioinformatics database, we predicted that miR-139-5p is a potential target of SNHG3. Overexpression of miR-139-5p could impair the proliferation, migration, and invasion of Hep3B and HepG2 cells. Next, we performed bioinformatics analysis and luciferase reporter assay to show that BMI1 is a downstream mRNA target of miR-139-5p, which is consistent with another study.⁴⁵ However, Luo et al showed the relationship between miR-139-5p and BMI1 in the bladder cancer stem cell.⁴⁵ In their study, they suggested that BMI1 was a target and regulated by miR-139-5p, but didn't explain BMI1 biological function in bladder cancer. However, in our study, we silenced BMI1 in HCC cells and found cell proliferation, migration, and invasion were attenuated, which was similar to miR-139-5p overexpression or SNHG3 knockdown.

BMI1 is a member of the polycomb group of proteins. Many studies have shown that BMI1 is upregulated in many

cancers, including breast cancer,²² pancreatic cancer,⁴⁶ and HCC.⁴⁷ In our study, we found that BMI1 was upregulated in HCC tissues and cell lines. When knocking down the expression of BMI1, the proliferation, migration, and invasion of cells were inhibited.

Our present study revealed that SNHG3 can play an oncogenic role in the HCC development. We also observed that SNHG3 was upregulated in HCC tissues and cells, while miR-139-5p was downregulated. In addition, either SNHG3 knockdown or miR-139-5p overexpression could inhibit the proliferation, migration, and invasion in HCC cells. Bioinformatics analysis suggested that miR-139-5p was a target of SNHG3, and the expression of miR-139-5p was regulated by SNHG3. However, the accurate mechanism of regulatory function of SNHG3 in miR-139-5p expression is still needed to explore in the future study. Next, we predicted that BMI1 was a potential target mRNA of miR-139-5p, and BMI1 was upregulated in HCC tissues. The proliferation, migration, and invasion were impaired by BMI1 silencing. But, we cannot exclude that miR-139-5p may target other mRNAs and regulate HCC progression through unexplored pathways. Hence, exploring other potential mRNA targets will facilitate our understanding of miR-139-5p in HCC. Subsequently, we did the xenograft assay using HepG2 cells stably transfected with siSNHG3, and found that the tumor was significantly inhibited. In conclusion, it was indicated that lncSNHG3/miR-139-5p/BMI1 axis was involved in HCC proliferation, migration, and invasion.

Ethical statement

The ethical approval was obtained from the Research Ethics Committee of the First Affiliated Hospital of Sun-Yat-sen University (Guangzhou, China). Thirty-one specimens of cancerous tissue and 31 specimens of paracancerous tissue were collected from the HCC patients and all patients signed the informed consent form, and this study was performed in accordance with the ethical standards detailed in the Declaration of Helsinki. Animal experiments were conducted according to the National Institutes of Health Guidelines for the Care and Use of Laboratory Animals, and approved by the Institutional Animal Care and Use Committee of the First Affiliated Hospital of Sun-Yat-sen University.

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Author contributions

All authors contributed to data analysis, drafting and revising the article, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

Disclosure

The authors report no conflicts of interest in this work.

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Supplemental materials

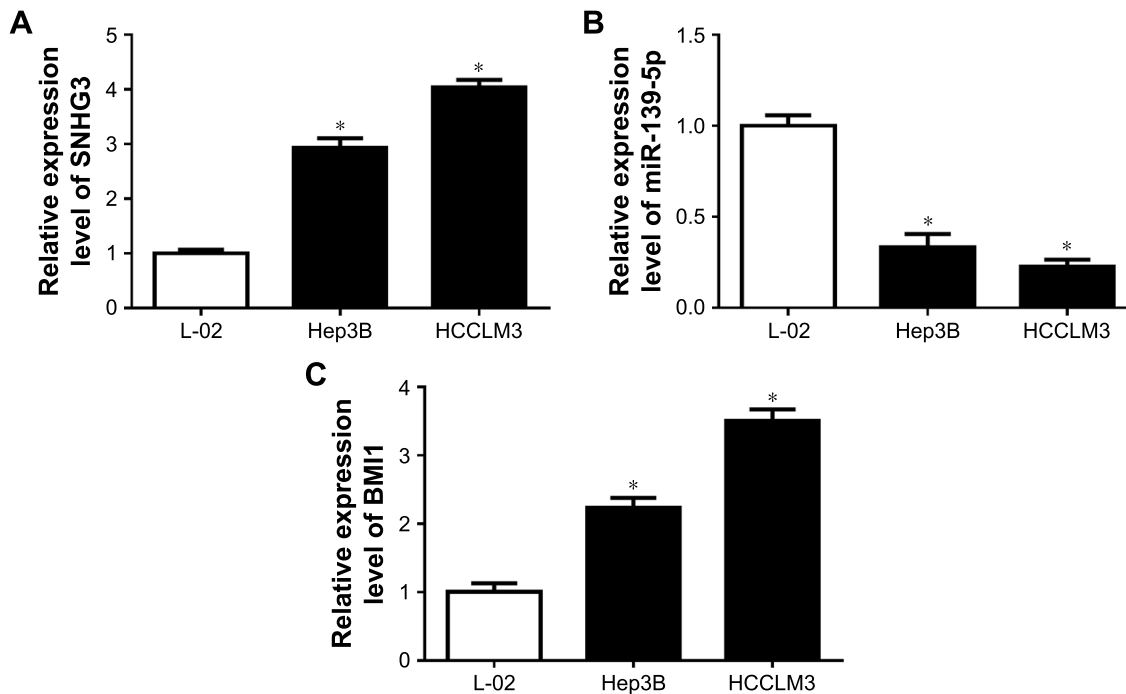


Figure S1 The expression of SNHG3, miR-139-5p and BMI1 in normal and HCC cells. **(A)** SNHG3 expression in L-01, Hep3B and HCCLM3 cells. **(B)** miR-139-5p expression in L-01, in Hep3B and HCCLM3 cells. **(C)** BMI1 expression in L-01, Hep3B and BMI1 cells. * $P < 0.05$.

Abbreviation: HCC, hepatocellular carcinoma.

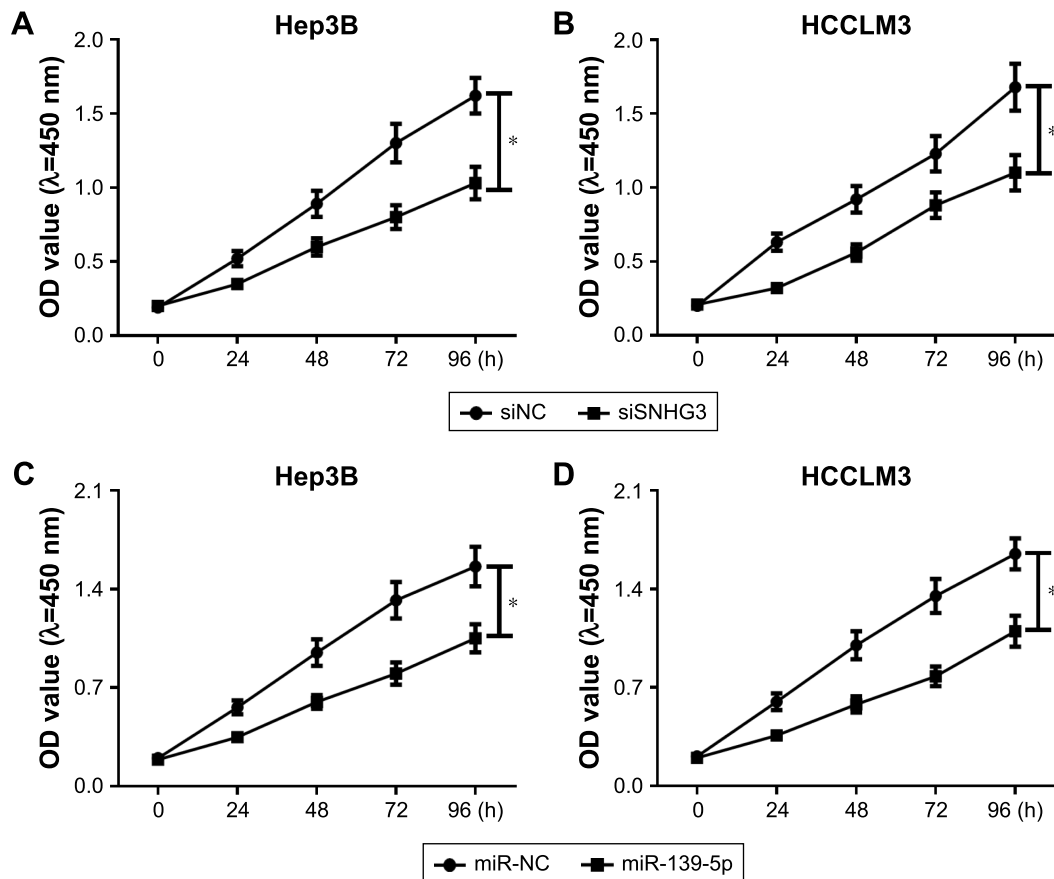


Figure S2 (Continued)

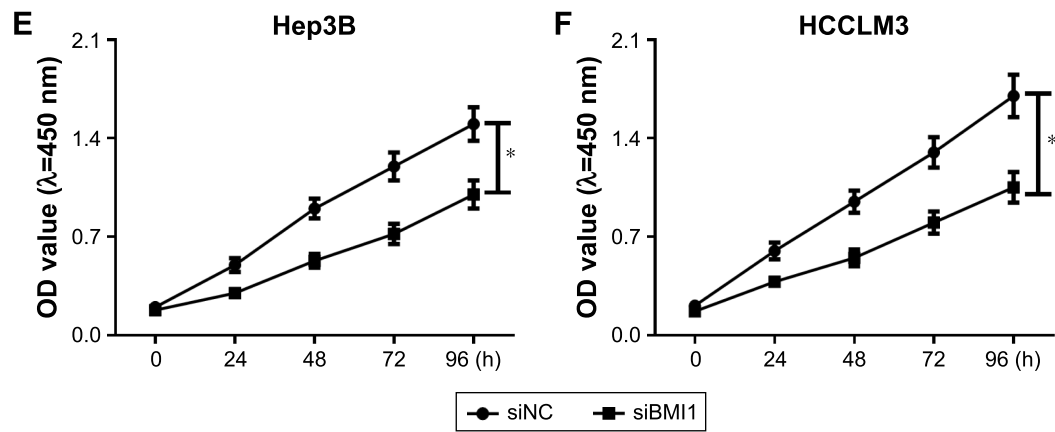


Figure S2 Effect of SNHG3, miR-139-5p and BMI1 on cell proliferation in HCC. MTT assay detected the proliferation of Hep3B (A) and HCCLM3 (B) after SNHG3 knockdown. MTT assay measured the proliferation of Hep3B (C) and HCCLM3 (D) after miR-139-5p overexpression. MTT assay evaluated the proliferation of Hep3B (E) and HCCLM3 (F) after BMI1 downregulation. *P<0.05.

Abbreviations: HCC, hepatocellular carcinoma; MTT, 3-(4,5-dimethyl-2-thiazolyl)-2,5-diphenyl-2-H-tetrazolium bromide.

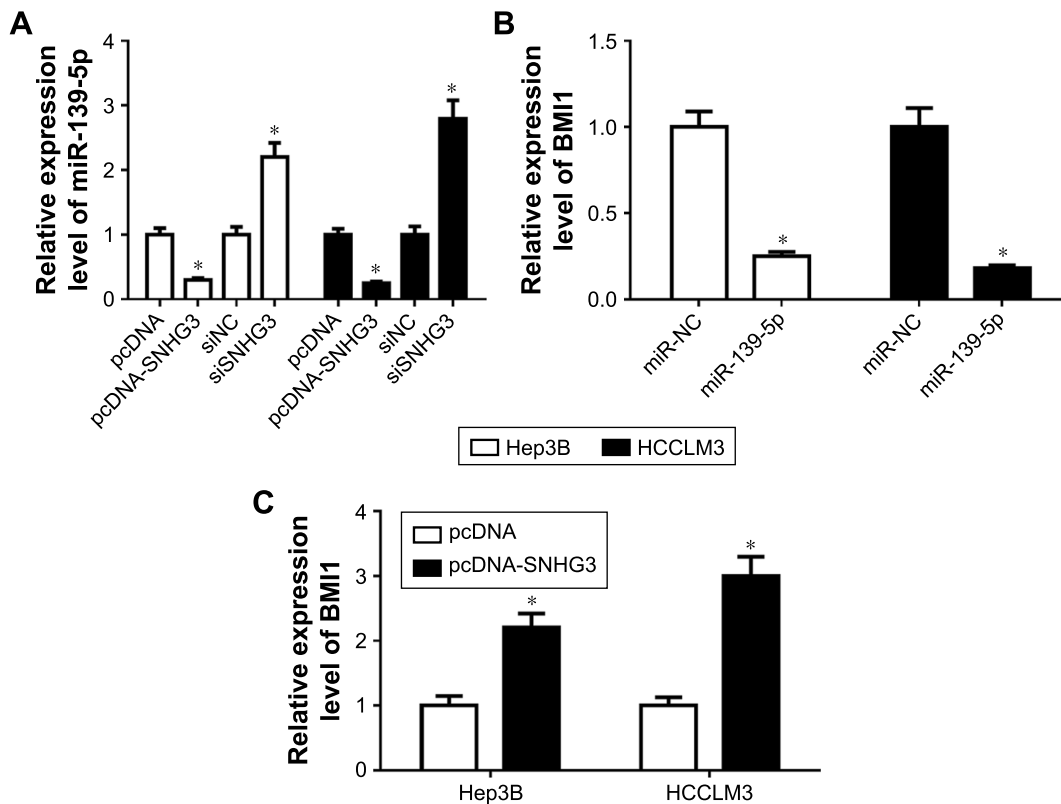


Figure S3 The expression of miR-139-5p and BMI1 in HCC cells. (A) miR-139-5p expression in Hep3B and HCCLM3 cells transfected with pcDNA, pcDNASNHG3, si-NC or siSNHG3. (B) BMI1 expression in Hep3B and HCCLM3 cells after miR-139-5p overexpression. (C) BMI1 expression in Hep3B and HCLM3 cells after SNHG3 overexpression. *P<0.05.

Abbreviation: HCC, hepatocellular carcinoma.

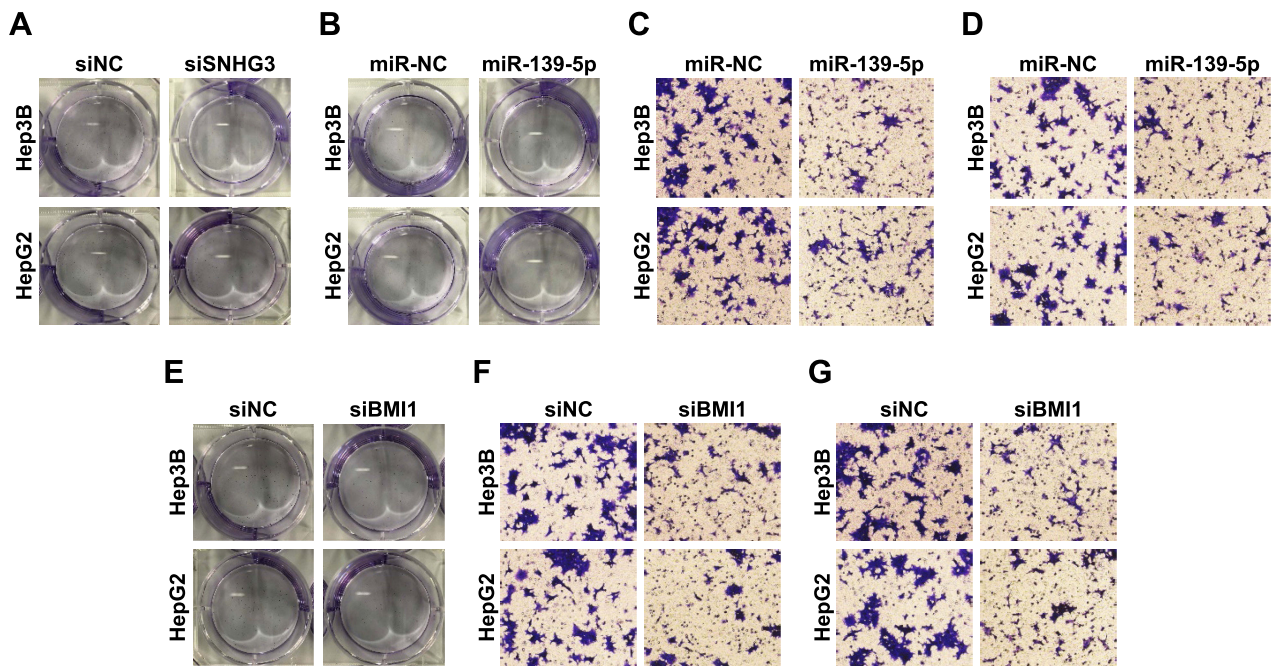


Figure S4 siNHG3 regulated cell migration and invasion by targeting miR-139-5p/BMI1 axis. **(A)** Colony formation in HepG2 and Hep3B cells transfected with siNC and siSNHG3. **(B)** Colony formation in HepG2 and Hep3B cells transfected with miR-NC and miR-139-5p. **(C, D)** Cell migration and invasion of HepG2 and Hep3B cells transfected with miR-NC and miR-139-5p. **(E)** Colony formation in HepG2 and Hep3B cells transfected with siNC and siBMI1. **(F, G)** Cell migration and invasion of HepG2 and Hep3B cells transfected with siNC and siBMI1. Magnification 200 \times .

Abbreviation: HCC, hepatocellular carcinoma.

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