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Long Noncoding RNA CCAL Promotes Papillary Thyroid Cancer Progression by Activation of NOTCH1 Pathway

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Long noncoding RNA CCAL has been reported to promote tumor progression in various human cancers, including hepatocellular carcinoma, osteosarcoma, and colorectal cancer. However, the role of CCAL in papillary thyroid cancer remains largely unknown. In the present study, we found that the expression of CCAL was upregulated in papillary thyroid tumor tissues compared to adjacent normal tissues. Moreover, the expression of CCAL was positively related with papillary thyroid cancer severity and TNM stage and predicted poor prognosis. Besides, we found that knockdown of CCAL significantly inhibited papillary thyroid cancer cell proliferation, migration, and invasion in vitro and reduced tumor growth and metastasis in vivo. We found that knockdown of CCAL dramatically decreased the expression of NOTCH1 and suppressed the activation of the NOTCH1 signaling pathway. Furthermore, overexpression of NOTCH1 rescued the proliferation, migration, and invasion in papillary thyroid cancer cells. Taken together, our data indicated that CCAL promoted papillary thyroid cancer development and progression by activation of the NOTCH1 pathway, which provided a new insight on the design of therapeutic targets.

Key words: Papillary thyroid cancer; Colorectal cancer-associated lncRNA (CCAL); Proliferation; Migration; NOTCH1

INTRODUCTION

Thyroid cancer is one of the most malignant and prevalent endocrine tumors around the world¹. In recent years, the incidence of thyroid cancer has been rapidly increasing². Nearly 80% of thyroid tumors are papillary thyroid carcinomas (PTCs)³. For the treatment of PTC, surgical resection combined with radioiodine and levothyroxine treatment is the main method⁴. Although most patients with PTC displayed a relatively good prognosis after surgical treatment, there are still many PTC-induced deaths every year worldwide. Until now, there have been no effective diagnostic biomarkers and therapeutic targets for PTC. Therefore, it is quite necessary to discover the molecular mechanisms in PTC for the development of novel therapies.

Long noncoding RNAs (lncRNAs) are a class of transcripts with a length longer than 200 nucleotides⁵. Many reports show that lncRNAs have no protein-coding potential⁶. Increasing evidence demonstrates that lncRNAs exerted very important functions in various biological processes, including cell development, immune regulation,

and especially tumorigenesis⁷⁻⁹. More and more reports indicate that dysregulation of lncRNAs was closely related to tumor development and progression in various cancers^{10,11}. lncRNAs could regulate cellular proliferation, apoptosis, migration, and invasion in human cancers^{12,13}. For instance, overexpression of lncRNA small nucleolar RNA host gene 6 (SNHG6) is related to poor prognosis in gastric cancer and enhances the proliferation and epithelial-mesenchymal transition (EMT) of tumor cells¹⁴. lncRNA maternally expressed gene 3 (MEG3) could promote cisplatin-induced apoptosis in human glioma cells¹⁵. Some lncRNAs, such as metastasis-associated lung adenocarcinoma transcript 1 (MALAT1), HIT000218960, and homeobox C transcript antisense intergenic RNA (HOTAIR), are reported to be involved in the progression of thyroid cancer¹⁶⁻¹⁸. However, most lncRNAs are waiting for identification, and their roles need to be investigated in thyroid cancer.

Previous studies indicate that colorectal cancer-associated lncRNA (CCAL) serves as an oncogene in hepatocellular carcinoma (HCC), osteosarcoma (OS),

and colorectal cancer (CRC)^{19–21}. However, whether CCAL plays a role in PTC remains to be explored. In our study, we found that CCAL was upregulated in PTC tissues compared with adjacent normal tissues. Furthermore, we found that overexpression of CCAL predicted PTC malignance, high TNM stage, and poor prognosis. Knockdown of CCAL significantly inhibited PTC cell proliferation, migration, and invasion *in vitro* and *in vivo*. To identify the mechanism of action, we found that knockdown of CCAL remarkably suppressed the expression of NOTCH1 and activation of NOTCH1 signaling pathway. Overexpression of NOTCH1 rescued the proliferation, migration, and invasion of PTC cells. Our study demonstrated that CCAL promoted PTC progression by activation of the NOTCH1 pathway.

MATERIALS AND METHODS

Patient Samples

Fifty-two PTC tissue specimens were obtained from the Shanghai University of Traditional Chinese Medicine (Shanghai, P.R. China). No patients received radiation therapy or chemotherapy prior to surgery. Tissue samples were classified according to the World Health Organization criteria and stored in liquid nitrogen or at -80°C . This study was approved by the Institutional Review Board of Shanghai University of Traditional Chinese Medicine. Written informed consent was obtained from all participating patients.

Cell Lines and Cell Culture

Human thyroid cancer epithelium cell lines (BCPAP, FTC133, 8505C, TPC-1, and HTH-83) and a normal human thyroid epithelium cell line (TEC) were commercially purchased from the American Type Culture Collection (ATCC, Manassas, VA, USA). Cells were cultured in RPMI-1640 medium (Sigma-Aldrich; Merck KGaA, Darmstadt, Germany) supplemented with 10% fetal bovine serum (Thermo Fisher Scientific, Inc., Waltham, MA, USA) at 37°C in a humidified atmosphere with 5% CO_2 .

Reverse Transcription Quantitative Polymerase Chain Reaction (RT-qPCR)

Total RNA was extracted from tissue samples and cells using TRIzol[®] reagent (Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer's protocol. Total RNA was reverse transcribed into cDNA using the PrimeScript[™] RT reagent kit with gDNA Eraser (Takara Biotechnology Co., Ltd., Dalian, P.R. China). RT reaction mixtures contained 1 μg of total RNA, 1 μl of RT Enzyme Mix 1, 1 μl of RT Primer Mix, 4 μl of 5 \times PrimeScript Buffer 2, and 4 μl of ddH₂O. The temperature protocol was 37°C for 15 min and 85°C for 5 s. qPCR was performed on cDNA using the QuantiFast SYBR-Green PCR kit

(Qiagen GmbH, Hilden, Germany) on a StepOnePlus[™] Real-Time PCR System (Applied Biosystems; Thermo Fisher Scientific, Inc.). PCR mixtures contained 10 μl of 2 \times PCR Master Mix, 4 μl of sense and antisense primers, 2 μl of cDNA, and 4 μl of ddH₂O. The specificity of each PCR was confirmed using a melting curve analysis. PCR primers were synthesized by Invitrogen, Thermo Fisher Scientific, Inc. The results were normalized to the expression of GAPDH. All samples were assessed in triplicate.

siRNA-Mediated Interference

Small interfering RNAs (siRNAs) against CCAL consisting of three target-specific siRNAs were designed and synthesized by Shanghai GenePharma Co., Ltd. (Shanghai, P.R. China). BCPAP and TPC-1 cells at a density of 3×10^5 cells/ml were seeded into six-well plates and transfected with siRNAs using Lipofectamine[®] 2000 (Invitrogen; Thermo Fisher Scientific, Inc.) as the transfection reagent, according to the manufacturer's protocol. After 48 h of transfection, cells were harvested for RT-qPCR to verify the silencing of CCAL expression.

In Vivo Tumor Growth and Metastasis Assays

For tumor growth assay, 6-week-old male mice with severe combined immune deficiency (SCID; Institute of Zoology, Chinese Academy of Sciences, Beijing, P.R. China) received subcutaneous injections of 1×10^6 cells infected with siCCAL or control ($n=6$ mice/group). At 1, 3, 5, and 7 weeks after inoculation, tumor volume was monitored and calculated. All mice were sacrificed by euthanasia at 7 weeks postinoculation, and the tumors were removed. For tumor metastasis assay, 6-week-old male SCID mice were injected with 1×10^6 cells infected with siCCAL or control through the tail vein ($n=6$ mice/group). The mice were sacrificed by euthanasia at 8 weeks postinoculation. Anatomized mice were examined for metastasis in the lung. Animal experiments were performed in accordance with relevant guidelines and regulations of the Institutional Animal Care and Use Committees at Shanghai University of Traditional Chinese Medicine, and protocols were approved by the Institutional Animal Care and Use Committees at Shanghai University of Traditional Chinese Medicine.

Western Blot

Protein was extracted by cell lysis using RIPA (Beyotime, P.R. China) containing protease inhibitors (Beyotime). Protein lysates were separated in 10% SDS-PAGE and transferred onto PVDF membrane (Millipore, USA). The membranes were blocked by 5% milk in TBST buffer and incubated with primary antibodies overnight at 4°C . PVDF membranes were washed in TBST and incubated with horseradish peroxidase-conjugated secondary antibodies (ProteinTech Group, USA). Proteins were

visualized using ECL Western blotting substrate (Pierce, Rockford, IL, USA).

ChIP Assay

ChIP assay was performed according to a previous study²². In brief, cells were treated with 1% formaldehyde for 10 min for cross-linking, crashed with SDS lysis buffer, and followed by ultrasonication. Standard ChIP assay was performed using H3K4me3 antibody. Enriched DNA fragments were eluted and analyzed by qRT-PCR.

Cellular Proliferation Assays

Cells were seeded into 96-well plates at a density of 1×10^3 cells/well. After 24, 48, and 72 h of incubation at 37°C, cellular viability was evaluated using a cell counting kit-8 (CCK-8) assay (Dojindo Molecular Technologies, Inc., Kumamoto, Japan) according to the manufacturer's protocol. The absorbance was measured at a wavelength of 450 nm using a Multiskan™ GO Microplate Spectrophotometer (Thermo Fisher Scientific, Inc.).

A colony formation assay was also performed. Cells were seeded into six-well plates and cultured at 37°C with 5% CO₂ for 14 days. Colonies were fixed with methanol at room temperature for 20 min and stained with 0.1% crystal violet (Sigma-Aldrich; Merck KGaA). The total number of visible colonies was determined under an optical microscope (Olympus Corporation, Tokyo, Japan). All experiments were repeated three times.

Cellular Migration and Invasion Assays

Cellular migration was assessed using 6.5-mm Transwell inserts with 8.0- μ m-pore polycarbonate membranes (Costar; Corning Incorporated, Corning, NY, USA). A cell migration or invasion (coated with Matrigel) assay was performed using the Transwell inserts. Briefly, 2×10^5 transfected and nontransfected cells were suspended and seeded into the upper chambers of the inserts. Medium supplemented with 10% FBS was added into the lower chambers as a chemoattractant. Following 24 h of incubation at 37°C, cells on the upper surface of the membrane were removed; cells that had migrated to the lower membrane were fixed with 100% methanol at room temperature for 20 min and stained with crystal violet. Cells were observed using an optical microscope (Olympus Corporation). Cells were counted in five random fields from each well, and the average number of migrated or invaded cells was calculated. The assays were performed in triplicate.

Statistical Analysis

The statistical significance of the differences between groups was assessed using Student's *t*-test for pairwise comparisons or one-way analysis of variance followed by Fisher's least significant difference post hoc test for

multiple comparisons. A value of $p < 0.05$ was considered to indicate a statistically significant difference. Data are expressed as the mean \pm standard deviation of three independent experiments. Statistical analysis was performed using the SPSS software version 20.0 (IBM Corp., Armonk, NY, USA).

RESULTS

CCAL Was Highly Expressed in PTC Tissues

To determine the function of CCAL in PTC, we examined the expression of CCAL in PTC tissues ($n=52$) and normal tissues ($n=52$) by RT-qPCR. Results indicated that CCAL was upregulated in PTC tissues compared with adjacent normal tissues ($p < 0.001$) (Fig. 1A). Moreover, as shown in Figure 1B and C, upregulated CCAL expression was significantly correlated with multifocality ($p < 0.01$) and TNM stage ($p < 0.05$). Then we checked the expression level of CCAL in PTC cell lines (BCPAP, FTC133, 8505C, HTH-83, and TPC-1). Compared to the normal human thyroid epithelium cell line (TEC), CCAL was highly expressed in PTC cell lines as shown by RT-qPCR (Fig. 1D). We then divided the PTC tissues into two subgroups based on CCAL expression. We performed Kaplan–Meier analysis to determine the correlation between CCAL expression and clinical outcome in PTC patients. Results indicated that patients with a higher CCAL expression have worse overall survival ($p=0.029$) (Fig. 1E).

CCAL Knockdown Suppressed Cell Proliferation

To further explore the effect of CCAL on PTC cells, we silenced CCAL in BCPAP and TPC-1 cells (Fig. 2A). To understand the impact of CCAL on the proliferation of thyroid carcinoma cell, CCK-8 assay was conducted after transfection. As shown in Figure 2B and C, the proliferation rate of BCPAP and TPC-1 cells was suppressed by transfecting with siCCAL at 48 and 72 h ($p < 0.05$). As shown in Figure 2D, the colony formation assay showed a significant reduction in colony numbers consistent with the CCK-8 assay. To further evaluate the effect of CCAL expression on cell proliferation, the cell cycle distribution was explored in BCPAP and TPC-1 cells transfected with siCCAL by flow cytometry, and the results showed that the cell cycle progression was stagnated at the G₁–G₀ phase compared with the control cells transfected with empty vector (Fig. 2E). Taken together, CCAL knockdown could inhibit PTC cell proliferation by blocking the cell cycle.

Knockdown of CCAL Inhibited Cell Migration and Invasion

A previous study showed that CCAL could promote the metastasis of OS¹⁹. To further determine whether CCAL also regulates the metastasis in PTC, we performed

Transwell assays with CCAL-depleted or control BCPAP and TPC-1 cells. As shown, CCAL knockdown significantly inhibited the numbers of migrated and invaded BCPAP and TPC-1 cells (Fig. 3A and B). EMT is a hallmark of metastatic neoplasms. As shown in Figure 3C, knockdown of CCAL upregulated the epithelial cell marker E-cadherin and downregulated vimentin and Snail, which are characteristic of mesenchymal cells. These data suggested that CCAL promoted PTC cell migration and invasion.

CCAL Knockdown Inhibited Tumor Growth and Metastasis In Vivo

In the next step, to evaluate the effect of CCAL on PTC cells in vivo, we subcutaneously injected BCPAP cells into SCID mice. Tumor volumes were measured at weeks 1, 3, 5, and 7 postinjection, and the mice were sacrificed at 7 weeks. The volume and weight of the tumors derived from CCAL-depleted BCPAP cells were lower than those of the control group (Fig. 4A–C). Furthermore, the rate of lung metastasis was also lower in xenograft tumors expressing siCCAL than control group (Fig. 4D). Then Western blot was used to assess the proliferation and metastasis of formed tumor tissues. As shown in Figure 4E, knockdown of CCAL reduced the levels of

cyclin D1, Snail, and Twist, which suggested that CCAL knockdown inhibited tumor proliferation and metastasis in vivo.

Knockdown of CCAL Inhibited the Activation of NOTCH1 Pathway in PTC Cells

Emerging evidence has shown that NOTCH1 signaling is important for tumor growth and metastasis in PTC^{22–24}. In our study, we also found that knockdown of CCAL significantly inhibited the expression of NOTCH1, HEY1, and HEY2 in BCPAP and TPC-1 cells, which suggested that CCAL knockdown led to inactivation of NOTCH1 signaling (Fig. 5A and B). To determine how CCAL regulates NOTCH1 expression, we performed ChIP assay with CCAL-silenced or control BCPAP cells. We found that CCAL knockdown significantly inhibited the enrichment of histone H3 lysine 4 trimethylation (H3K4me3), an active histone modification, on NOTCH1 promoter (Fig. 5C), suggesting that CCAL regulates NOTCH1 transcription accessibility. To further determine whether CCAL-mediated regulation on PTC cell proliferation and metastasis relied on the NOTCH1 signaling pathway, we restored the activation of NOTCH1 pathway by overexpressing NOTCH1 in CCAL-silenced BCPAP and TPC-1 cells (Fig. 5D). As shown by the CCK-8 and

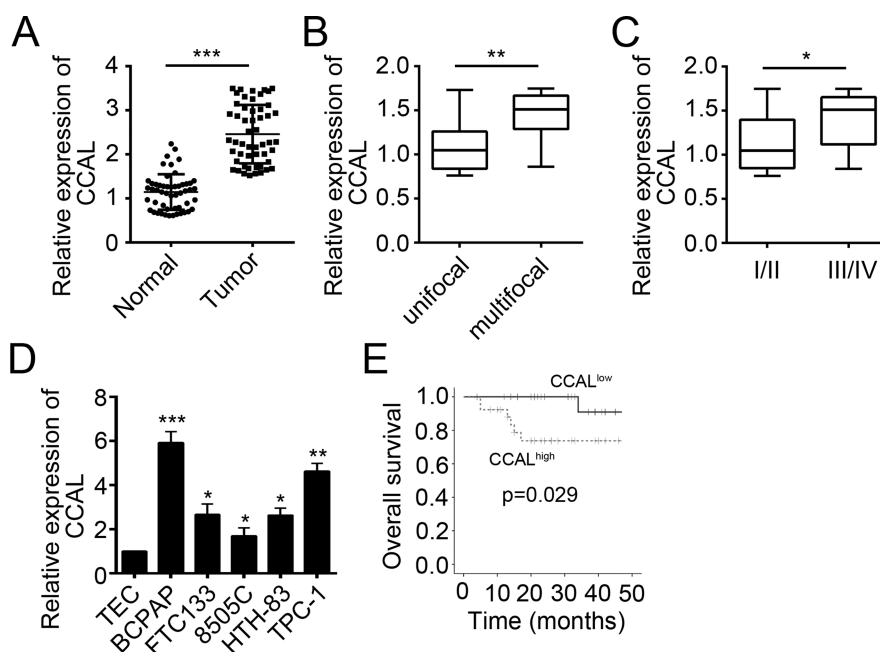


Figure 1. Colon cancer-associated long noncoding RNA (CCAL) was highly expressed in papillary thyroid carcinoma (PTC) tissues. (A) Comparing the different expression of CCAL between PTC tissues ($n=52$) and normal thyroid tissues ($n=16$) by reverse transcription quantitative polymerase chain reaction (RT-qPCR). (B) CCAL expression was dramatically upregulated in patients with multifocal PTC. (C) CCAL expression was dramatically upregulated in patients with advanced TNM stage. (D) CCAL expression between different types of thyroid carcinoma cell lines and a normal human thyroid epithelium cell line (TEC) by RT-qPCR. (E) Kaplan–Meier survival analysis based on CCAL expression in PTC tissues. * $p<0.05$, ** $p<0.01$, and *** $p<0.001$ by two-tailed Student’s t -test. All data presented are shown as means \pm SD collected from three independent experiments.

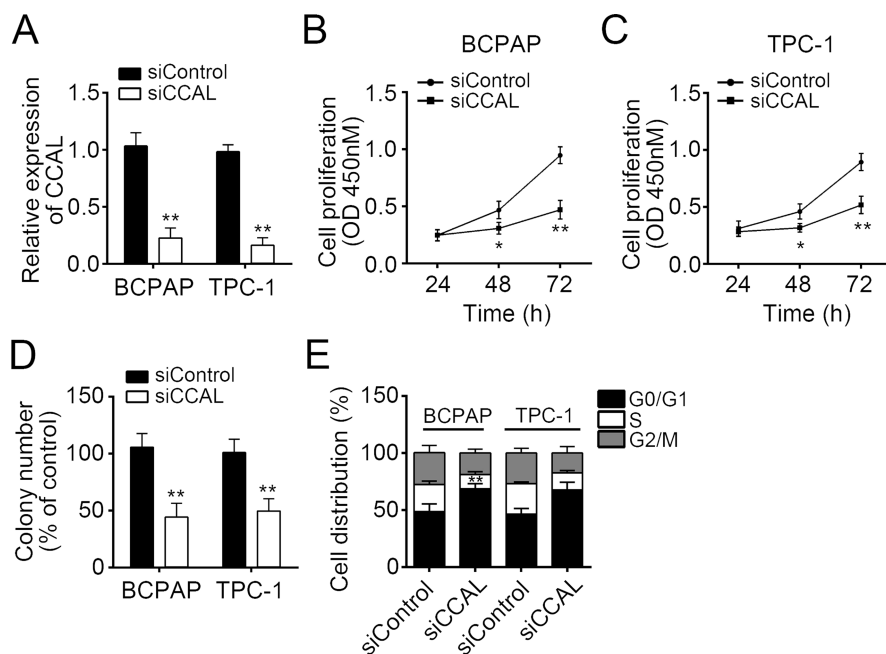


Figure 2. CCAL knockdown suppressed cell proliferation. (A) RT-qPCR was used to determine the knockdown efficiency of CCAL in BCPAP and TPC-1 cells. (B–D) Cell counting kit-8 (CCK-8) and colony formation assays were used for the detection of cell proliferation potential in BCPAP and TPC-1 cells. (E) Cell cycle distribution of BCPAP and TPC-1 cells transfected with siCCAL relative to cells transfected with control was measured by propidium iodide staining using flow cytometry. * $p < 0.05$ and ** $p < 0.01$ by two-tailed Student's *t*-test. All data presented are shown as means \pm SD collected from three independent experiments.

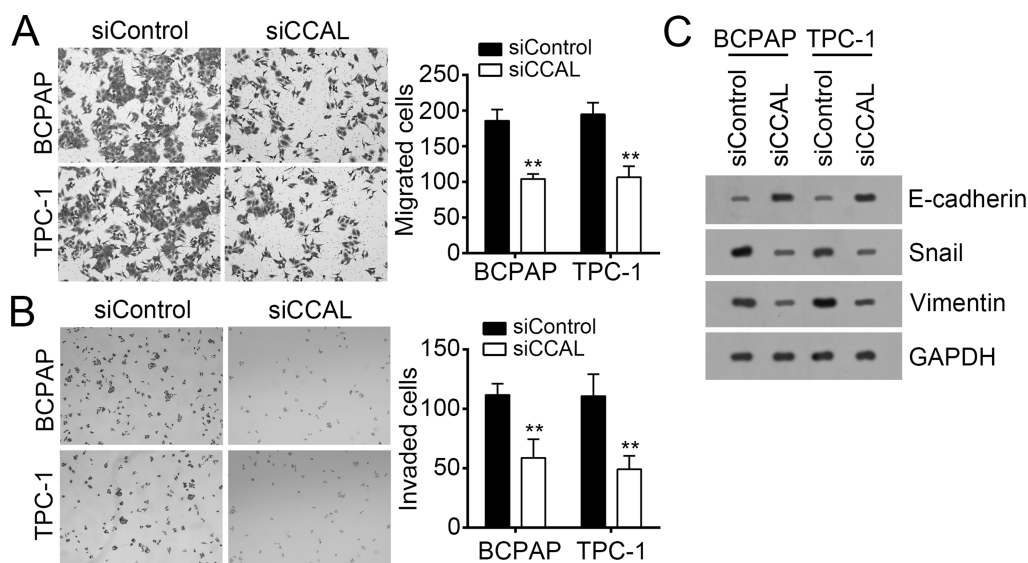


Figure 3. Knockdown of CCAL inhibited cell migration and invasion. (A) Transwell migration and (B) invasion assays were used to assess the migration and invasion abilities of BCPAP and TPC-1 cells. (C) The protein levels of Snail, vimentin, and E-cadherin were checked by Western blot in BCPAP and TPC-1 cells. GAPDH acted as loading control. ** $p < 0.01$ by two-tailed Student's *t*-test. All data presented are shown as means \pm SD collected from three independent experiments.

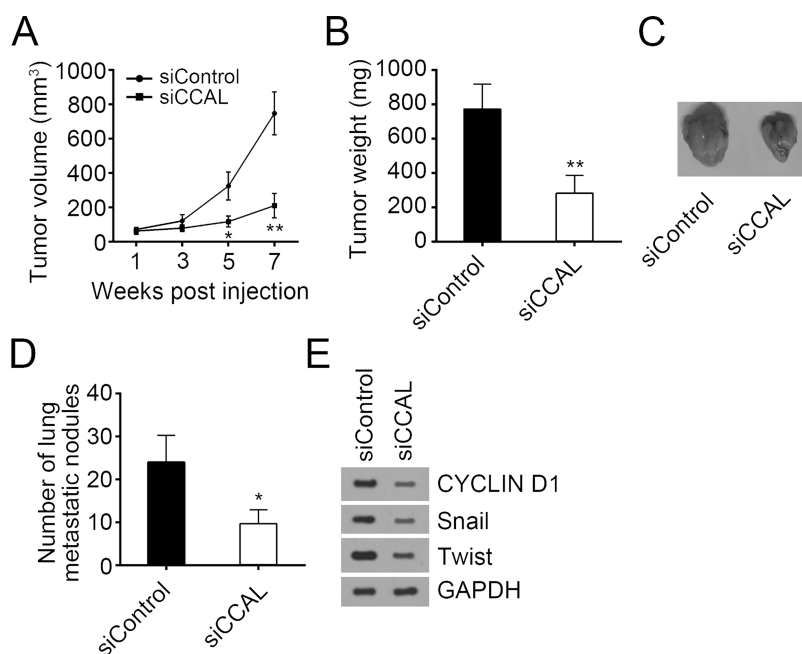


Figure 4. CCAL knockdown inhibited tumor growth and metastasis in vivo. (A–C) BCPAP cells were injected into nude mice subcutaneously. Then tumor volumes (A) and tumor weight (B and C) were measured at indicative time points. (D) The numbers of metastatic foci in the lungs of mice from various groups at 8 weeks after tail vein injection were counted. (E) The protein levels of cyclin D1, Twist, and Snail were measured by Western blot. * $p < 0.05$ and ** $p < 0.01$ by two-tailed Student's *t*-test. All data presented are shown as means \pm SD collected from three independent experiments.

Transwell assays, overexpression of NOTCH1 rescued the proliferation and invasion of CCAL-silenced BCPAP and TPC-1 cells (Fig. 5E–G). Taken together, CCAL promoted the proliferation, migration, and invasion of PTC cells through activating the NOTCH1 signaling pathway.

DISCUSSION

PTC patients have died mainly as a result of insufficient specific diagnostic biomarkers and therapeutic strategies²⁵. Defining the molecular mechanism of PTC progression and identifying novel therapeutic targets for PTC intervention are of great importance. More and more evidence shows that lncRNAs have broad biological functions, especially in cell growth, differentiation, and tumorigenesis²⁶. lncRNAs may be a good candidate for tumor diagnosis and prognosis. For example, lncRNA chromosome 17 open reading frame 91 (C17orf91) is a potential prognostic marker and serves as an oncogene in ovarian cancer²⁷. A recently identified lncRNA named sex-determining region Y box 21 antisense 1 (SOX21-AS1) indicates poor prognosis in lung adenocarcinoma²⁸. In this study, we found that CCAL was highly expressed in PTC tissues and aimed to determine its function in PTC.

Dysregulation of lncRNAs has been related to diverse human cancers including HCC, gastric cancer, and colon cancer^{11,29,30}. Previous reports showed that a lncRNA was highly expressed in HCC tissues and associated with

HCC metastasis and TNM stage¹⁹. They found that CCAL promoted HCC cellular invasion and proliferation and inhibited cell apoptosis by activating the Wnt/ β -catenin pathway. Another report indicated that CCAL was significantly overexpressed in OS tissues compared with adjacent normal tissues²⁰. Increased expression of CCAL was also correlated with advanced TNM stage and metastasis in OS. Additionally, Ma and colleagues demonstrated that CCAL was a key regulator of CRC progression. CCAL could activate Wnt/ β -catenin signaling via suppression of activator protein 2 α (AP-2 α) expression in CRC²¹. Until now, the function of CCAL is largely unknown. We found that the expression of CCAL was significantly upregulated in PTC tissues. Increased expression of CCAL was positively related with advanced TNM stage and predicted poor prognosis in PTC. By CCK-8 and Transwell assays, we found that knockdown of CCAL in BCPAP and TPC-1 cells remarkably suppressed cellular proliferation, migration, and invasion in vitro. Moreover, xenograft experiments indicated that knockdown of CCAL markedly inhibited tumor growth and lung metastasis in vivo. Thus, our data suggested that CCAL might be a good biomarker for PTC patients' prognosis and be a potential therapeutic target.

Previous studies widely proved that NOTCH signaling is very important for tumor progression in various human cancers^{23,24,31–33}. For instance, miR-199a-3p inhibits

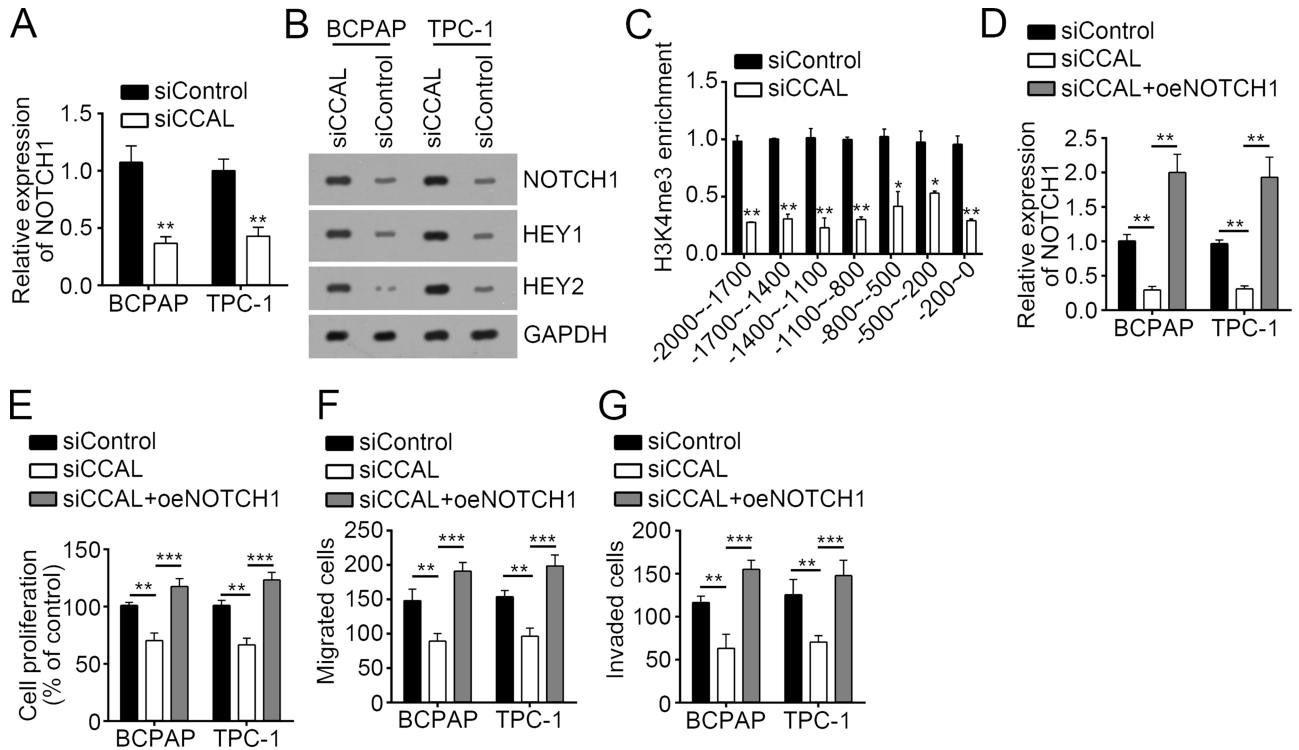


Figure 5. Knockdown of CCAL inhibited the activation of NOTCH1 pathway in PTC cells. Knockdown of CCAL significantly inhibited the (A) mRNA and (B) protein levels of NOTCH1 and the protein levels of its target genes (HEY1 and HEY2) in BCPAP and TPC-1 cells. (C) CCAL knockdown significantly inhibited the enrichment of histone active modification H3K4me3 on NOTCH1 promoter. ChIP assay was conducted with CCAL-silenced or control BCPAP cells. (D) RT-qPCR analysis indicated that NOTCH1 was restored in CCAL-depleted BCPAP and TPC-1 cells. (E) CCK-8 assays showed that overexpression of NOTCH1 rescued the proliferation of CCAL-depleted BCPAP and TPC-1 cells. Transwell assays indicated that overexpression of NOTCH1 rescued the (F) migration and (G) invasion of CCAL-depleted BCPAP and TPC-1 cells. * $p < 0.05$, ** $p < 0.01$ and *** $p < 0.001$ by two-tailed Student's *t*-test. All data presented are shown as means \pm SD collected from three independent experiments.

cell proliferation and induces apoptosis by suppressing Jagged1–Notch signaling in human HCC³¹. The NOTCH1 pathway promoted microRNA-151-5p expression and contributed to gastric cancer progression³². In PTC, researchers found that NOTCH1 signaling was activated and promoted the proliferation and cell cycle of PTC cells²⁴. Another study indicates that activation of NOTCH1 signaling inhibits Prospero homeobox 1 (PROX1) activity and contributes to the malignant behavior of thyroid cancer cells²³. A previous study also shows that the NOTCH1 receptor was upregulated and serves as a marker of lymph node metastases in PTC³³. In our study, we showed that CCAL promoted the activation of NOTCH1 signaling pathway. By Western blot, we found that CCAL knockdown significantly inhibited the protein levels of NOTCH1 and its target genes (HEY1 and HEY2). Moreover, we also found that restoration of NOTCH1 promoted the proliferation, migration, and invasion of PTC cells. Our data revealed the relationship between CCAL expression and NOTCH1 pathway activation in PTC.

In summary, our research, for the first time, demonstrated the function of CCAL in PTC and explained its

functional mechanism. We found that CCAL promoted the proliferation, migration, and invasion of PTC cells, at least in part, by activation of the NOTCH1 signaling pathway. Our findings highlight the importance of the CCAL/NOTCH1 axis in PTC progression.

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