Long noncoding RNA H19 is a critical oncogenic driver and contributes to epithelial-mesenchymal transition in papillary thyroid carcinoma

Wei-Quan Liang¹,*
De Zeng²,*
Chun-Fa Chen¹
Shu-Ming Sun¹
Xiao-Feng Lu¹
Chun-yan Peng³
Hao-Yu Lin¹

¹Department of Thyroid and Breast Surgery, The First Affiliated Hospital of Shantou University Medical College, Shantou, Guangdong 515000, People’s Republic of China.
²Department of Medical Oncology, Cancer Hospital of Shantou University Medical College, Shantou, Guangdong 515031, People’s Republic of China.
³Department of Clinical Laboratory, Taihei Hospital of Hubei University of Medicine, Hubei 442008, People’s Republic of China.

*These authors contributed equally to this work.

Background: Growing evidence has indicated that the long noncoding RNA H19 (IncRNA H19), frequently deregulated in almost all tumor types tested, acted as a pivotal contributor to both cancer initiation and progression. However, the role of IncRNA H19 in human papillary thyroid carcinoma (PTC) remains controversial. The aim of the study was to investigate the expression and potential function of IncRNA H19 in human PTC.

Patients and methods: The IncRNA H19 level was determined by quantitative real-time (RT)-PCR analyses in 58 PTC tissue samples and their paired paracancerous tissue samples. RNA interference, RT-PCR analysis, and Western blot assay were used to determine the impact of IncRNA H19 on epithelial-mesenchymal transition (EMT) markers in human PTC cells. The migratory and invasive capacities of PTC cells were determined by wound-healing and transwell migration and invasion assays.

Results: IncRNA H19 expression was 2.417-fold higher in PTC tissues than their paired paracancerous tissue (95% CI: 1.898–2.935, P<0.0001). Higher level of IncRNA H19 was correlated to elevated expression of Vimentin, ZEB2, Twist, and Snail2. Inhibition of IncRNA H19 resulted in upregulation of E-cadherin and downregulation of Vimentin both at mRNA and protein levels. Conversely, enforced expression of the exogenous IncRNA H19 led to E-cadherin mRNA and protein downregulation and relative upregulation of Vimentin. Moreover, wound-healing and transwell migration and invasion assays showed that IncRNA H19 could promote the migratory and invasive abilities of PTC cells.

Conclusion: The level of IncRNA H19 was significantly higher in PTC tissues than paired paracancerous tissue or normal tissues. Overexpression of IncRNA H19 was correlated with higher tumor burden of PTC. It also contributes to EMT process, as well as promotes migration and invasion of PTC cells.

Keywords: long noncoding RNA H19 (IncRNA H19), papillary thyroid carcinoma, epithelial-mesenchymal transition.

Introduction

Growing concern over the sharp increase in the prevalence of thyroid cancer in past decades, particularly papillary thyroid carcinoma (PTC), has prompted investigators to explore the molecular underpinnings of the disease, leading to the identification of BRAFV600E and RAS mutations, as valid therapeutic targets.¹–³ Recently, the Encyclopedia of DNA Elements project, which was designed to discover and characterize novel RNAs transcripts, has revealed that a large amount of primary or processed protein-coding and noncoding transcripts, including thousands of long noncoding...
RNAs (lncRNAs), are predominantly implicated in the human genome. Notably, increasing studies have demonstrated that deregulated expression of lncRNAs was significantly involved in the pathogenesis of multiple types of cancer.

The lncRNAs are a new class of nonprotein transcripts with over 200 bases in length. Emerging evidence has indicated that the IncRNA H19, frequently deregulated in almost all tumor types tested, acted as a critical contributor to both cancer initiation and progression. Li et al reported that overexpression of IncRNA H19 enhances carcinogenesis and metastasis of gastric cancer. Guan et al reported that overexpression of IncRNA H19/miR-675 promotes tumorigenesis in head and neck squamous cell carcinoma. A study by Shi et al demonstrated that IncRNA H19 predicts poor prognosis in melanoma cells. Moreover, preclinical and clinical studies have highlighted the therapeutic potential against IncRNA H19. For instance, Zhao et al showed that downregulation of IncRNA H19 inhibits migration and invasion of human osteosarcoma through the NF-kB pathway.

Despite enormous breakthroughs in the understanding of the role of IncRNA H19 in multiple types of tumor progression, the exact role and function of IncRNA H19 in human PTC remain under-investigated. A major mechanism through which BRAF mutation might contribute to primary PTC progression is through induction of EMT. Specifically, the Snail/E-cadherin axis has been demonstrated as a pathway dysregulated by BRAF, leading to EMT. The possibility that IncRNA H19 can regulate EMT program is an attractive option to explore. In this study, we identified that the IncRNA H19 was significantly increased in PTC tissues and associated with classic EMT markers in PTCs, suggesting that IncRNA H19 may be required for human PTC tumorigenesis and development.

Patients and methods

Patients and clinical tissue samples

Written informed consents were received from all patients before enrollment, and this study was approved by the Ethics Committee of Shantou University Medical Hospital. Fifty-eight PTC tissue samples and their paired paracancerous tissue samples were obtained from patients (age range: 27–72 years; 16 males and 42 females), who were first diagnosed between June 2016 and December 2017 at the First Affiliated Hospital of Shantou University Medical Hospital for quantitative real-time (qRT)-PCR analyses and paraffin-embedded pathological investigation. All medical histories of the patients were well-documented according to the eighth edition of the American Joint Committee on Cancer TNM system.

Cell lines and cell culture

The human PTC cell line B-CPAP was purchased from the Chinese Academy of Sciences (Beijing, China). This cell line was established from the tumor tissue of a woman aged 72 years with metastatic PTC in 1992. Cells were cultured in RPMI-1640 medium, containing 10% FBS, 100 U/mL penicillin, and 100 mg/mL streptomycin (all from Thermo Fisher Scientific, Waltham, MA, USA) in a 5% CO₂ incubator at 37°C.

RNA purification and RT-PCR analysis

All obtained tissues during surgery were immediately stored in liquid nitrogen prior to use. Total RNA (1 µg) was isolated from tissues or cells using TRIzol (Thermo Fisher Scientific) following the manufacturer’s instructions and stored at −80°C. Reverse transcription was performed using a PrimeScript™ RT reagent kit (Takara Bio Inc., Shiga Prefecture, Japan) according to the manufacturer’s instructions. To detect mRNA expression, qRT-PCR was performed with an SYBR Select Master Mix (Thermo Fisher Scientific) and a CFX96 RT-PCR Detection System (Bio-Rad Laboratories Inc., Hercules, CA, USA). To normalize the amount of mRNA in each sample, GAPDH was used. Primer sequences used in RT-PCR are listed in Table S1.

Small interfering RNAs (siRNAs) and transfection

The siRNAs were purchased from Suzhou GenePharma Co., Ltd. (Suzhou, China). The siRNAs are shown in Table S2. Cells were transfected using Lipofectamine 3000 (Life Technologies, Carlsbad, CA, USA). A total of 1×10⁶ B-CPAP cells were inoculated into each well of a 6-well culture plate. The next day, 75 pmol of siRNA was combined with 3.75 µL of Lipofectamine 3000 (Life Technologies) by strictly following the manufacturer’s instructions.

Western blot assay

Cells were lysed in RIPA buffer (Cell Signaling Technology, Danvers, MA, USA) with protease inhibitors. The extracted protein sample was measured using the BCA protein assay and was stored at −80°C until required for use. The 30 µg protein sample was separated by 8% SDS-PAGE and transferred onto a PVDF membrane (EMD Millipore, Billerica, MA, USA). After transferring, the membrane was blocked in...
5% nonfat milk and then incubated with primary antibodies at 4°C overnight. The antibodies are shown in Table S3. The next day, the membrane was incubated with secondary antibodies and subjected to electrochemiluminescence detection (Applygen Technologies, Inc., Beijing, China).

**Wound healing assay**

Thirty minutes before the application of injury lines, B-CPAP cells were treated with Mitomycin C (25 mg/mL) (Beyotime Institute of Biotechnology, Jiangsu, China). Injury line was made with a 2-mm wide tip separated on cells plated in culture dishes at 90% confluency. After washing with PBS, cells could migrate in complete medium and photographs were taken (×40) after 48 hours. An average of five random widths of each injury line was measured for quantitation.

**Transwell invasion assays**

Cell culture inserts (8 M pore size; BD, Franklin Lakes, NJ, USA) and Matrigel invasion chambers (BD) were used to perform migration and invasion assays, respectively. Transfected cells were serum-starved for 24 hours and then 3×10^4 B-CPAP cells in serum-free medium were seeded into the upper chamber. Complete medium was added to the bottom chamber. Cells were stained with 0.1% crystal violet for migration assays after 36 and 48 hours for invasion assays. Each assay was performed in triplicate. The number of cells from five fields in each well were counted by two investigators (HYL and DZ).

**Clinical databases**

The ONCOMINE datasets (http://www.oncomine.org/) were used to analyze the expression of *lncRNA H19* in various cancers against normal tissue as well as the correlated expression genes in thyroid cancer.20 The correlations between the expression levels of *lncRNA H19* and *Vimentin*, *ZEB2*, *Twist*, or *Snail2* in thyroid cancers were determined through analysis in the cBioPortal database (http://www.cbioportal.org/index.do).20 We analyzed thyroid carcinoma (TCGA, Provisional) and PTC (TCGA, Cell 2014) datasets. The expression levels of *lncRNA H19* in a series of cancers were analyzed by CCLE database (https://portals.broadinstitute.org/ccle/home), which is an online encyclopedia of a compilation of gene expression, chromosomal copy number, and massively parallel sequencing data from 947 human cancer cell lines.21

**Statistical analysis**

Statistical analysis was performed using the SPSS version 18.0. Data from three independent experiments were presented as the mean ± SD. Statistical analysis was performed using Mann–Whitney U test and Wilcoxon signed-rank test. The clinicopathological characteristics were analyzed by two-way ANOVA. For clinical data from ONCOMINE and cBioPortal datasets, Student’s t-test was used, and two times of fold change with a *P*-value of <0.01 was defined as clinically significant. Correlation coefficients between RNA levels were obtained through Spearman’s rank correlation analysis as was described earlier.22 A value of *P*<0.05 was considered as a statistically significant.

**Ethical statement**

The study was approved by the ethics committee of the First Hospital of Shantou University Medical College. Patient consent was written informed consent, in compliance with the Declaration of Helsinki.

**Results**

The expression level of *lncRNA H19* was significantly increased in PTC tissues

*lncRNA H19* has been identified as an oncogenic gene in multiple cancer types and its elevated expression was tightly linked to tumorigenesis and cancer progression. ONCOMINE analysis revealed that *lncRNA H19* expression was significantly higher in cancer than normal samples across a wide variety of datasets in different cancer types. A total of 220 different types of studies about *lncRNA H19* were collected in the ONCOMINE database (Figure S1) in which 37 studies showed the increase of *lncRNA H19* expression and 22 studies showed a decrease. In addition, CCLE analysis demonstrated that the expression level of *lncRNA H19* in thyroid cancer cells ranked the eleventh highest in a variety of cancer cell lines tested (Figure S2). Thus, we performed the qRT-PCR assay to detected PTC tissue samples and corresponding adjacent nontumorous tissues samples from 58 patients who were first diagnosed in our hospital. Our data showed that *lncRNA H19* expression was 2.417-fold higher (95% CI: 1.898–2.935) in PTC tissues than their paired paracancerous tissues (Figure 1A, *P*<0.0001). These results implied that *lncRNA H19* may be distinctively expressed higher in thyroid cancer, suggesting it might play unique roles in the development of thyroid cancer.

**Elevated lncRNA H19 expression correlated with tumor burden of PTC**

To further investigate the correlation between *lncRNA H19* expression and clinicopathological features, qRT-PCR analy-
ses of lncRNA H19 were performed using samples from 58 PTC patients. When the 58 tumor tissues were stratified on clinical progression, we found that higher lncRNA H19 expression is remarkably associated with increased tumor size (Figure 1B, \( P = 0.0407 \)). In addition, lncRNA H19 expression levels were much higher in lymph node metastasis group than those in patients without lymph node metastasis (Figure 1C, \( P = 0.0305 \)). Moreover, high lncRNA H19 expression was correlated with PTC patients of stage III/IV than those in patients with stage I/II (Figure 1D, \( P = 0.0033 \)). Consistently, when 58 PTC patients were divided into two group, lncRNA H19\textsuperscript{high} (>median), and lncRNA H19\textsuperscript{low} (<median), our data also revealed that more patients in increased tumor size group (\( P = 0.036 \)), lymph node metastasis group (\( P = 0.002 \)), and higher stage group (\( P = 0.004 \)) correlated with lncRNA H19\textsuperscript{high} group (Table 1). However, the level of lncRNA H19 expression was not associated with age, gender, extrathyroidal extension status, pathological subtype, and whether combined with nodular goiter between these two groups (Table 1). It was, therefore, postulated that the high lncRNA H19 expression level was correlated with more malignant phenotypes with increased proliferation and potential of metastasis in PTC.

**Higher expression of lncRNA H19 correlated with elevated expression of mesenchymal phenotypes markers Vimentin, ZEB2, Twist, and Snail2**

To explore the inherent properties of lncRNA H19 and its potential mechanism in PTC, we used the ONCOMINE and cBioPortal databases to analyze the correlation of lncRNA H19 in connection with other featured biomarkers in PTC. Interestingly, in ONCOMINE co-expression analysis, it was found that lncRNA H19 expression was significantly correlated with the expression of ZEB2, a typical mesenchymal phenotypes marker (Figure 2, \( r = 0.700 \)). Moreover, gene correlation targeted analysis from TCGA database showed that in thyroid carcinoma, higher expression of lncRNA H19
we verified whether *IncRNA H19* was associated with typical EMT markers in PTC cell line B-CPAP. Suppression of *IncRNA H19* by siRNA caused a significant upregulation of epithelial marker *E-cadherin* expression at both the mRNA and protein levels, while that of *Vimentin*, a mesenchymal marker, decreased as compared to controls (Figure 4A and B). Two different siRNA sequences were used to silence *IncRNA H19* separately to exclude the possibility of nonspecific target. It was found that siH19-a and siH19-b effectively suppressed *IncRNA H19* and finally caused downregulated *Vimentin*, while upregulating *E-cadherin* both at mRNA and protein levels (Figure 4A and B). We chose siH19-b with better suppression effect on *IncRNA H19* for further experiments. Conversely, enforced expression of the exogenous *IncRNA H19* led to *E-cadherin* mRNA and protein downregulation and relative upregulation of *Vimentin* (Figure 4C and D). These results implied that *IncRNA H19* was able to mediate the process of EMT in PTC cells.

**IncRNA H19** promotes migration and invasion abilities of PTC cells

To evaluate whether the expression of *IncRNA H19* regulates cellular function via inducing EMT, we performed gain-of-function and loss-of-function studies on B-CPAP cells. We first assessed the effect of *IncRNA H19* overexpression on B-CPAP cells invading through the Matrigel-coated membrane (Figure 5A). As displayed in Figure 5A, after culturing for 48 hours, the width of the scratch wound in siH19-b B-CPAP cells had reduced to 75% as compared to controls (Figure 5B), indicating that loss of *IncRNA H19* expression inhibits cell motility. In contrast, when enforcing expression of the exogenous *IncRNA H19* in B-CPAP cells, the scratch wound width had healed to 26% as compared to 0 hours, dwarfing the 44% gap healed by control cells. This result suggests that *IncRNA H19* overexpression was capable of promoting the motility of B-CPAP cells (Figure 5C). We further investigated whether *IncRNA H19* influenced the migration and invasion abilities of B-CPAP cells via transwell migration and invasion assays. As shown in Figure 5D, the number of migrated cells transfected with siH19 was significantly lower (~52%) than the number of migrated cells transfected with si-control (Figure 5F). Similarly, invasion assays showed that knocking down *IncRNA H19* via siRNAs resulted in fewer cells invading through the Matrigel-coated membrane (Figure 5E and G). Moreover, the ectopic expression of *IncRNA H19* in B-CPAP cells promoted cellular migration and invasion (Figure 5D–G). These results indicate that *IncRNA H19* enhances the migration and invasion abilities of PTC cells.

### Table 1 Clinicopathological characteristics of 58 PTC patients according to *H19* expression

<table>
<thead>
<tr>
<th>Clinicopathological features</th>
<th><em>H19</em>&lt;sup&gt;hi&lt;/sup&gt; (&lt;median, n=29)</th>
<th><em>H19</em>&lt;sup&gt;lo&lt;/sup&gt; (&lt;median, n=29)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>17 11</td>
<td>12 18</td>
<td>0.115</td>
</tr>
<tr>
<td>&lt;55 years</td>
<td>6 10</td>
<td>23 19</td>
<td>0.240</td>
</tr>
<tr>
<td>≥55 years</td>
<td>23 19</td>
<td>11 19</td>
<td>0.036†</td>
</tr>
<tr>
<td>Tumor size</td>
<td>≤2 cm 11</td>
<td>11 19</td>
<td>0.002†</td>
</tr>
<tr>
<td>&gt;2 cm</td>
<td>18 10</td>
<td>18 10</td>
<td>1.000</td>
</tr>
<tr>
<td>LN metastasis</td>
<td>Yes 20</td>
<td>No 9</td>
<td>0.004†</td>
</tr>
<tr>
<td>Distance metastasis</td>
<td>Yes 27</td>
<td>No 2</td>
<td>0.581</td>
</tr>
<tr>
<td>Clinical stage</td>
<td>I+II 11</td>
<td>III+IV 18</td>
<td>0.730</td>
</tr>
<tr>
<td>Extrathyroidal extension</td>
<td>Yes 18</td>
<td>No 11</td>
<td>0.788</td>
</tr>
<tr>
<td>Pathological subtype</td>
<td>Classical 25</td>
<td>Follicular 4</td>
<td></td>
</tr>
<tr>
<td>Nodular goiter</td>
<td>Yes 12</td>
<td>No 17</td>
<td></td>
</tr>
</tbody>
</table>

Notes: †P-value<0.05 was considered statistically significant.
Abbreviations: LN, lymph node; PTC, human papillary thyroid carcinoma.

*H19* correlated with elevated mesenchymal biomarker *Vimentin* (Figure 3A, r=0.22, P<0.001), *ZEB2* (Figure 3C, r=0.19, P<0.001), *Twist* (Figure 3E, r=0.56, P<0.001), and *Snail2* (Figure 3G, r=0.48, P<0.001). Meanwhile, correlation analysis in PTC showed that elevated *IncRNA H19* correlated with increased expression of *Vimentin* (Figure 3B, r=0.20, P<0.001), *ZEB2* (Figure 3D, r=0.23, P<0.001), *Twist* (Figure 3F, r=0.57, P<0.001), and *Snail2* (Figure 3H, r=0.50, P<0.001). These results showed that the expression of *IncRNA H19* in PTC positively associated with mesenchymal phenotypes marker, which imply that *IncRNA H19* might play a distinct role in the processing of EMT and lead to more aggressive clinicopathological features of PTC.

**IncRNA H19** regulates mRNA and protein expression of EMT markers *E-Cadherin* and *Vimentin*

Since the above data from the public database and clinical sample have shown that *H19* was correlated with EMT in PTC,
Discussion

Thyroid cancer is the most common endocrine malignancy across the world. Its prevalence has been steadily increasing in the past few decades, primarily due to a remarkable increase in the incidence of PTC. Multiple patterns of genetic alteration, including BRAF mutations, RAS mutations, and RET/PTC rearrangements, have been identified to be critically involved in the tumorigenesis and progression of PTCs. Targeting therapies toward these genetic alterations have led to a significant improvement in the therapeutic response and survival of patients with PTCs.

Except for a series of protein-coding genetic alteration, a variety of noncoding transcripts have also been found to play essential roles in the development of a number of solid tumors. However, up to now, the alteration in the noncoding transcript in PTCs remained under-investigated. It has been recognized in most studies that lncRNA H19 overexpresses and contributes to tumorigenesis and progression of multiple cancer types, including gastric cancer, glioma, and head and neck squamous cell carcinoma. Nevertheless, it was found that lncRNA H19 was under-expressed in intra-tumoral hepatocellular carcinoma (HCC) tissues, as compared to peritumoral tissues. Moreover, lncRNA H19 could suppress the progression and metastasis of HCC. Lv et al also reported that downregulation of lncRNA H19 and MiR-675 promotes migration and invasion of human HCC cells through AKT/GSK-3beta/Cdc25A signaling pathway.

Figure 2 H19 expression was significantly correlated with ZEB2 expression in thyroid carcinoma (shown in red frame, from ONCOMINE correlation analysis).
Figure 3 Elevated H19 expression correlated with higher expression of Vimentin, ZEB2, Twist, and Snail2 in thyroid carcinoma.

Notes: In the TCGA (the Cancer Genome Atlas) database, the Pearson’s chi-squared test was performed to analyze the correlation of H19 and some typical mesenchymal markers. (A and B) Gene correlation targeted analysis between H19 and Vimentin in samples of (A) thyroid carcinoma or (B) PTC. (C and D) Gene correlation targeted analysis between H19 and ZEB2 in samples of (C) thyroid carcinoma or (D) PTC. (E and F) Gene correlation targeted analysis between H19 and Twist in samples of (E) thyroid carcinoma or (F) PTC. (G and H) Gene correlation targeted analysis between H19 and Snail2 in samples of (G) thyroid carcinoma or (H) PTC.

Abbreviation: PTC, human papillary thyroid carcinoma.
IncRNA H19 has been recognized to act as an oncogenic driver in a number of cancer types, such as gastric, breast, and lung cancers. However, the specific function of IncRNA H19 in thyroid cancer remains controversial. For example, Wang et al reported that IncRNA H19 represses cell viability, and restrains migratory and invasive capabilities of thyroid cancer cells through downregulation of IRS-1. Lan et al demonstrated that inhibition of IncRNA H19 promotes the proliferation and motility of PTC. On the contrary, Liu et al reported that increased IncRNA H19 expression levels are associated with poor prognosis in thyroid cancer patients. Another study by Liu et al suggested that increased expression of IncRNA H19 promoted growth, migration, and invasion in thyroid cancer, whereas H19 knockdown impaired TC cell viability. Our analysis in the ONCOMINE database demonstrated that IncRNA H19 expression was markedly higher in thyroid cancer than normal tissue samples. Particularly, the qRT-PCR assay of 58 PTCs and corresponding adjacent nontumorous tissues from our institution also showed that IncRNA H19 expression was considerably higher in PTC tissues than paired paracancerous tissues. These findings consistently suggested that IncRNA H19 distinctively was expressed higher in PTCs than normal counterparts. Notably, we found that higher IncRNA H19 level was remarkably associated with larger tumor size, increased number of lymph node metastasis, as well as more advanced clinical stage, implying that higher IncRNA H19 level is correlated to higher tumor burden of PTCs.

Notably, emerging evidences in recent years indicated that IncRNA H19 acts as an important inducer of EMT.

**Figure 4** H19 regulates expression of EMT marker E-cadherin and Vimentin in PTC cells.

**Notes:** (A and B) Expression of E-cadherin and Vimentin in B-CPAP cells at protein and mRNA levels analyzed by (A) Western blot and (B) RT-PCR, respectively, when silencing H19 by siH19-a and siH19-b. (C and D) Expression of E-cadherin and Vimentin in B-CPAP cells at protein and mRNA levels analyzed by (C) Western blot and (D) RT-PCR, respectively, when overexpressing H19. *P<0.05 as compared to control groups.

**Abbreviations:** EMT, epithelial-mesenchymal transition; PTC, papillary thyroid carcinoma; RT-PCR, real-time PCR.
Figure 5 H19 knockdown in B-CPAP cells inhibits EMT while overexpression promotes EMT.

**Notes:** (A–C) Wound healing assay showed the knockdown of H19 inhibited the cellular motility of B-CPAP cells, while overexpression of H19 enhanced such motility. (A) Representative pictures and (B and C) quantitative data of wound recovery after 48 hours cell culture. Each sample contained three wells. (D–G) Knockdown of H19 decreased the cellular migration and invasion of B-CPAP cells, while overexpression of H19 enhanced such motility. (D) Representative pictures of migration and (E) invasion, (F) quantitative data of migration, or (G) invasion assays. All experiments were performed at least thrice and data were statistically analyzed by two-way ANOVA. *P<0.05 vs control. Error bars indicate SD.

**Abbreviation:** EMT, epithelial-mesenchymal transition.
Matouk et al have reported that the role of the oncofetal lncRNA H19 in tumor metastasis orchestrating the EMT-MET decision.42 Zhou et al reported that the lncRNA H19 mediates breast cancer cell plasticity during EMT and MET plasticity by differentially sponging miR-200b/c and let-7b.43 However, whether lncRNA H19 acts as an important inducer of EMT in papillary thyroid cancer remained unclear. Our analysis with ONCOMINE and cBioPortal databases, as well as TCGA datasets consistently demonstrated that lncRNA H19 was positively associated with key EMT factors. It was found that higher expression of lncRNA H19 was correlated to elevated expression of Vimentin, as well as critical EMT-associated transcription factors, including ZEB2, Twist, and Snail2.

Loss or decreased E-cadherin and increased Vimentin are universally regarded as the hallmark of the EMT process.44 In the present study, we first used the human PTC cell line B-CPAP to characterize the potential role of lncRNA H19 and found that lncRNA H19 was able to regulate mRNA and protein expression of EMT markers E-Cadherin and Vimentin, which suggested that lncRNA H19 was pivotal driver of EMT in PTC cells. In addition, our study also showed that lncRNA H19 promotes migration and invasion abilities of PTC cells. Shi et al reported that lncRNA H19 predicts poor prognosis in patients with melanoma and regulates cell growth, invasion, migration, and EMT in melanoma cells.12 Liao et al reported that downregulation of lncRNA H19 inhibits the migration and invasion of melanoma cells by inactivating the NF-κB and PI3K/Akt signaling pathways.45

Conclusion

In summary, the expression level of lncRNA H19 was significantly higher in PTC tissues than paired paracancerous tissue or normal tissues. Elevated lncRNA H19 expression was correlated with higher tumor burden of PTC. lncRNA H19 regulates mRNA and protein expression of classic EMT markers E-Cadherin and Vimentin and promotes migration and invasion of PTC cells. lncRNA H19 is a critical onco- genetic driver in PTC, and targeted therapy against lncRNA H19 might be a potential avenue, in conjunction with other conventional therapy, to increase the efficacy for patients with PTC.

Acknowledgments

This work was partly supported by the Natural Science Foundation of Guangdong Province, China (No. 2018A030313562), National Natural Science Foundation of China (81600358), and Youth Innovative Talent Project of Colleges and Universities in Guangdong Province, China (No. 2017KQNCX073).

Disclosure

The authors report no conflicts of interest in this work.

References

15. Knauf JA, Sartor MA, Medvedovic M, et al. Progression of BRAF-induced thyroid cancer is associated with epithelial-mesenchymal transition requiring concomitant MAP kinase and TGFβ signaling. Oncogene. 2011;30(28):3153–3162.


## Supplementary materials

### Table S1 Primers used in RT-PCR

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward primer</th>
<th>Reverse primer</th>
</tr>
</thead>
<tbody>
<tr>
<td>H19</td>
<td>5′-TCTGAGAGATTCAAAGCCTCCAC-3′</td>
<td>5′-GTCTCCACAACCTCAAACCAGTG-3′</td>
</tr>
<tr>
<td>E-cadherin</td>
<td>5′-TGCCCAGAAATGAAAAAGG-3′</td>
<td>5′-GTGTATGTGGCAATGCCTGTT-3′</td>
</tr>
<tr>
<td>Vimentin</td>
<td>5′-GAGAATTGTCCGTTAAGC-3′</td>
<td>5′-TCCAGAGCTTCCGTGAGT-3′</td>
</tr>
<tr>
<td>GAPDH</td>
<td>5′-TGGACTCCACGACGTACCTCAG-3′</td>
<td>5′-ACATGTTCAATATGATTCCA-3′</td>
</tr>
</tbody>
</table>

Abbreviation: RT-PCR, real-time PCR.

### Table S2 Oligonucleotide sequences for siRNA constructs

<table>
<thead>
<tr>
<th>SiRNAs</th>
<th>sense (5′–3′)</th>
</tr>
</thead>
<tbody>
<tr>
<td>siH19-a</td>
<td>5′-GCAAGCAGAGACAGUGUCC-3′</td>
</tr>
<tr>
<td>siH19-b</td>
<td>5′-CCAAGUAUAAGACACCAU-3′</td>
</tr>
<tr>
<td>siNC</td>
<td>5′-UUCUCCGAACGUGUGACGU-3′</td>
</tr>
</tbody>
</table>

### Table S3 Antibodies used in this study

<table>
<thead>
<tr>
<th>Antibody</th>
<th>Catalog number</th>
<th>Company</th>
<th>Concentration rate</th>
<th>Species</th>
</tr>
</thead>
<tbody>
<tr>
<td>E-cadherin</td>
<td>NCh-38</td>
<td>Dako Denmark A/S (Glostrup, Denmark)</td>
<td>1:3000</td>
<td>Rabbit</td>
</tr>
<tr>
<td>Vimentin</td>
<td>D21H3</td>
<td>Cell Signaling Technology (Danvers, MA, USA)</td>
<td>1:2000</td>
<td>Rabbit</td>
</tr>
<tr>
<td>GAPDH</td>
<td>sc-32333</td>
<td>Santa Cruz Biotechnology Inc. (Dallas, TX, USA)</td>
<td>1:3000</td>
<td>Mouse</td>
</tr>
</tbody>
</table>
**Figure S1** Expression pattern of H19 in different tumor types.

**Notes:** This figure shows the number of datasets with statistically significant RNA overexpression (red) or down-expression (blue) of the target gene (cancer vs. normal tissue). The P-value threshold is 0.01. The number in each cell represents the number of analyses that meet the threshold within those analysis and cancer types. The gene rank was analyzed by percentile of target gene in the top of all genes measured in each research. Cell color is determined by the best gene rank percentile for the analyses within the cell.
Figure S2. The RNA expression level of H19 in thyroid cancer cells ranked eleventh highest in a variety of cancer cell lines from Cancer Cell Line Encyclopedia analysis (shown in red frame).

Abbreviations: RMA, return material authorization; CML, chronic myelocytic leukemia; AML, acute myelocytic leukemia; NSC, non-small cell.