LncRNAs H19 and HULC, activated by oxidative stress, promote cell migration and invasion in cholangiocarcinoma through a ceRNA manner

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Abstract

Background: Long non-coding RNAs (lncRNAs) are known to play important roles in different cell contexts, including cancers. However, little is known about lncRNAs in cholangiocarcinoma (CCA), a cholangiocyte malignancy with poor prognosis, associated with chronic inflammation and damage to the biliary epithelium. The aim of the study is to identify if any lncRNA might associate with inflammation or oxidative stress in CCA and regulate the disease progression.

Methods: In this study, RNA-seq datasets were used to identify aberrantly expressed lncRNAs. Small interfering RNA and overexpressed plasmids were used to modulate the expression of lncRNAs, and luciferase target assay RNA immunoprecipitation (RIP) was performed to explore the mechanism of miRNA-lncRNA sponging.

Results: We firstly analyzed five available RNA-seq datasets to investigate aberrantly expressed lncRNAs which might associate with inflammation or oxidative stress. We identified that two lncRNAs, H19 and HULC, were differentially expressed among all the samples under the treatment of hypoxic or inflammatory factors, and they were shown to be stimulated by short-term oxidative stress responses to H2O2 and glucose oxidase in CCA cell lines. Further studies revealed that these two lncRNAs promoted cholangiocyte migration and invasion via the inflammation pathway. H19 and HULC functioned as competing endogenous RNAs (ceRNAs) by sponging let-7a/let-7b and miR-372/miR-373, respectively, which activate pivotal inflammation cytokine IL-6 and chemokine receptor CXCR4.

Conclusions: Our study revealed that H19 and HULC, up-regulated by oxidative stress, regulate CCA cell migration and invasion by targeting IL-6 and CXCR4 via ceRNA patterns of sponging let-7a/let-7b and miR-372/miR-373, respectively. The results suggest that these lncRNAs might be the chief culprits of CCA pathogenesis and progression. The study provides new insight into the mechanism linking lncRNA function with CCA and may serve as novel targets for the development of new countermeasures of CCA.

Keywords: Inflammation response, Oxidative stress, ceRNA, Cholangiocarcinoma, Migration and invasion

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Background
CCA is broadly described as malignancies originating from a malignant transformation of cholangiocytes and epithelial cells lining the intra-hepatic and extra-hepatic biliary ducts. CCA is difficult to diagnose and cure, partly due to the unknown molecular mechanisms underlying the development of CCA. In the past two decades, several studies have described molecular details regarding the malignant transformation of cholangiocytes and the environment of chronic inflammation in bile ducts with subsequent cholangiocyte damage, which is most commonly believed to contribute to CCA pathogenesis [1, 2]. A variety of inflammatory cytokines, including IL-6, IL-8, and TGFβ [2, 3], were found to contribute to the apoptosis, senescence, migration, invasion, and cell cycle regulation of malignant cholangiocytes [4]. Oxidative stress, potentially caused by Clonorchis sinensis, Opisthorchis viverrini, hepatitis B virus (HBV) infection, or inflammation [5–8], is also implicated in CCA development. The increased reactive oxygen species (ROS) produced by oxidative stress can damage lipids, proteins, and DNA and subsequently alter different cellular pathways and influence gene expression, which may induce tumor promotion and progression [9]. Persistent oxidative stress might also induce inflammatory responses, implying a complex association between oxidative stress and inflammation in CCA [7].

Previous studies have reported that malignant transformation of cholangiocytes was associated with chronic inflammation in the biliary epithelium; however, detailed molecular mechanisms of CCA promotion and progression are still unclear.

Recently, promising evidence has shown that non-coding RNAs serve as ideal diagnostic biomarkers and participate in oxidative stress and disease or cancer-related inflammation [10–14]. Small non-coding RNAs, especially microRNAs (miRNAs), are widely reported to play pivotal roles in disease pathogenesis, including the pathogenesis of CCA, and their post-transcriptional regulatory mechanisms [15, 16]. Several miRNAs, such as miR-370 [17], miR-373 [18, 19], and miR-21 [20, 21], are dysregulated and play vital roles in CCA. However, none of these miRNAs have been indicated to regulate inflammation or oxidative stress.

In addition to miRNAs, a set of long non-coding RNAs (lncRNAs), that are more than 200 nucleotides in length, have also been reported to play pivotal roles in a variety of diseases. Some lncRNAs, including ANRIL [22], HOTAIR [23], and H19 [24], influence many processes in various cancers, including chromatin remodeling, X chromosome inactivation, transcriptional regulation, molecular trafficking [25], inflammation responses, and oxidative stress [10–12, 14]. These observations suggest potential pathophysiological contributions from lncRNAs in CCA and imply that some lncRNAs might be involved in the inflammation response pathways stimulated by infection. Recently, the mechanisms and functions of several lncRNAs, such as IncRNA-HEIH [26], HULC [27], and HOTAIR [28], were uncovered in hepatic carcinoma, the most common hepatic malignancy, and these findings led to the construct of regulation networks and extended our knowledge regarding tumorigenesis in hepatic carcinoma. Similarly, dysregulated lncRNAs in CCA may play an important role in CCA promotion and progression through involvement in CCA key pathways, such as inflammation or oxidative stress. Currently, no lncRNAs have been reported to be associated with CCA progression.

In this study, we sought to determine if any lncRNA is stimulated by oxidative stress and trigger inflammation in CCA. We identified a set of lncRNAs that participated in inflammation and oxidative stress response. Two dysregulated lncRNAs H19 and HULC were shown to promote cell migration and invasion through ceRNA manners leading to the activation of inflammation cytokine IL-6 and chemokine receptor CXCR4. Our findings may provide potential biomarkers for CCA progression and potential therapeutic targets for the disease.

Methods
Cell lines
QBC939 human cholangiocarcinoma cells were obtained from Shuguang Wang (The Third Military Medical University, China). SK-cha-1 human cholangiocarcinoma cells were kindly provided by Dr. Chundong Yu (Xiamen University, China). RBE human cholangiocarcinoma cells and HEK293T human embryonic kidney cells were purchased from the Type Culture Collection of the Chinese Academy of Sciences (Shanghai, China). All CCA cell lines were cultured in RPMI-1640 medium with 10% fetal calf serum. HEK-293 T cells were cultured in DMEM medium with 10% fetal calf serum. All cells were maintained in a 37 °C humidified incubator with 5% CO₂.

qRT-PCR analysis
Cells were collected in EP tubes. Then, total RNA was extracted from tissue or cells using TRIzol reagent (Invitrogen, Carlsbad, CA, USA) following the manufacturer’s instructions. RNA was reverse-transcribed into cDNA with the ReverTra Ace® qPCR RT Kit (TOYOBO, Japan) or PrimeScript® RT reagent Kit with gDNA Eraser (Takara, Japan). Real-time PCR for lncRNA was performed using the SYBR Premix ExTaq real-time PCR kit (Takara, Japan) according to the manufacturer’s instructions with GAPDH as normalization control. The expression level for each lncRNA and mRNA was determined using the 2^−ΔΔCt method. The specificity and
reliability of the PCR were confirmed by sequencing the PCR product fragments. All primers are shown in Additional file 1: Table S1.

**Oxidative stress treatment in vitro**

CCA cells were treated with 1 μM H₂O₂ for 1 h and 1 μM glucose oxidase for 24 h for short-term and long-term oxide incubation in vitro, respectively.

**Vector construction**

Full-length H19-pcDNA 3.1 vectors were purchased from Integrated Biotech Solutions (Shanghai, China). The pcDNA 3.1 vector was used for IncRNA and mRNA overexpression, and the psiCHECK-2 vector was used for the luciferase target assay. Primers and oligonucleotides for full-length HULC, IL-6, and CXCR4 amplification and 3’-UTR segments of IL-6 and CXCR4 construction are listed in Additional file 1: Table S1.

**Cell transfection and luciferase target assay**

We plated 1.6 × 10⁴ RBE cells or 5 × 10⁴ HEK-293 T cells in 48-well plates. Then, 500 ng psiCHECK-2-derived reporter vectors (Promega, Madison, WI, USA), 500 ng pcDNA3.1 vectors overexpressing IncRNAs (Invitrogen Corporation, Carlsbad, CA, USA), and 30 nM miRNA mimics (GenePharma, Shanghai, China) were co-transfected with Lipofectamine® LTX (Invitrogen Corporation, Carlsbad, CA, USA) and Lipofectamine® 3000 (Invitrogen Corporation, Carlsbad, CA, USA). Finally, 48 h after transfection, cells were harvested for the dual luciferase reporter assay (Promega, Madison, WI, USA). Each experiment was repeated at least three times.

**Protein extraction, western blotting, and antibodies**

As previously described, 72 h after transfection, the RBE cells were harvested, and total protein was extracted from the cells using RIPA lysis buffer (Beyotime Biotechnology, Shanghai, China) with 1× complete ULTRA (Roche, Nutley, USA). The proteins were detected with antibodies against CXCR4 (ab2074, Abcam, USA), IL-6 (Roche, Nutley, USA). The protein levels were normalized to β-tubulin (32–2600, Invitrogen, USA). The protein levels were normalized to GAPDH or β-tubulin.

**RNA immunoprecipitation (RIP)**

In the assay, the enrichment of miRNAs binding with AGO2, a key component of the microRNA-containing RISC complex, were used as the positive controls, and U6 as the negative. The AGO2 (FB261-S2, Abnova, Taiwan) monoclonal primary antibody was rip grade, and the procedure of rip experiment was followed to the manual of Magan rip Kit (17–700, Millipore, USA).

**Scratch wound healing assay**

Transfected RBE cells were cultured in 24-well plates for 24 h in standard conditions until 80–100 % confluency was reached. Linear wound tracks were generated with sterile, 10-μl pipettes and maintained under standard conditions. The scratched cells were rinsed twice with PBS to remove non-adherent cells, and fresh RPMI-1640 medium (with no fetal calf serum) was added. Photographs of the centers of the gaps were taken using a 10× phase-contrast microscope (Zeiss). Cell migration at 0 and 24 h after scratching was evaluated by determining the wound distance at two random wound gap locations. Three independent scratch wound experiments were used for calculations.

**Migration and invasion assays**

Migration and invasion assays were conducted using Transwell chambers (8 μm, Corning Costar Co., Cambridge, USA) according to the manufacturer’s instructions. For the migration assays, 8 × 10⁴ RBE cells transfected after 24 h were plated in the top chambers lined with a non-coated membrane. For the invasion assay, the chamber inserts were coated with a 1:9 deliquation of Cultrex® Basement Membrane Extract (Trevigen, USA). Then, 1.2 × 10⁵ cells were plated in the top chamber, while 600 μl RPMI-1640 with 10 % fetal bovine serum was added to the lower chamber as a chemotactrant. After a 24-h incubation at 37 °C, cells located in the lower chamber were fixed with 100 % methanol and stained with 0.1 % crystal violet. Cells were counted in ten fields for triplicate membranes at 10× magnification using a microscope (Zeiss). Five random sights in each sample were selected to analyze cell count, and the mean of triplicate experiments was calculated.

**Results**

**H19 and HULC involved in inflammation-associated pathways triggered by oxidative stress in CCA cell lines**

In human bile duct cells, inflammatory responses were usually triggered by oxidative stress, which was induced by HBV, HCV, or excretory-secretory proteins released by *C. sinensis* or *O. riverine* [6, 8]. We speculated that the IncRNAs stimulated by oxidative stress may trigger inflammation in CCA. To identify whether any IncRNA related with inflammation is regulated by oxidative stress, we firstly reanalyzed several GEO data, GSE76743 [29], GSE57539 [30], GSE67106 [31], GSE55146 [32], and GSE70544 [33], and identified 40 IncRNAs responding to oxidative stress or inflammatory response (Fig. 1a). Among these, XIST, CDKN2B-AS1, CRNDE, TUG1, SOX2-OT, etc. have been reported playing crucial roles in inflammation pathway of cancer progression. Importantly, two IncRNAs, HULC and H19, showed higher expressed among all of the five selected datasets,
suggesting that they may involve in inflammation pathway that is regulated by oxidative stress. To validate the expression patterns of HULC and H19 under oxidative stress, we then used three human CCA cell lines, QBC939, SK-cha-1, and RBE, for short-term or long-term oxide incubation using H2O2 and glucose as described in previous studies [34] and measured the expression levels of HULC and H19 by qPCR. The results showed that both lncRNAs were significantly up-regulated in three cell lines (Fig. 1b), suggesting a potential function in acute oxidative stress responses. Similarly, HULC and H19 were also significantly up-regulated in at least two cell lines after long-term oxidative stress induced by glucose oxidase (Fig. 1c), and these two lncRNAs might be more likely to contribute to the pathogenesis of CCA through chronic inflammation caused by long-term oxidative stress. It is worth noting that HULC and H19 were up-regulated after both short- and long-term oxidative stress, implying their pivotal roles in inflammation promotion and CCA pathogenesis. Additionally, we have also investigated the heme oxygenase-1 (HO-1) gene expression, which were reported to have the cytoprotective effect under various pathophysiological stimulating conditions, such as oxidative stress [35]. As a result, the expression of HO-1 had increased significantly under the oxidative stress condition (Additional file 1: Figure S1), suggesting that the HO-1 may also protect cholangiocarcinoma from oxidative stress [36, 37].

**IncrRNAs H19 and HULC regulate cholangiocyte migration and invasion**

Combined with previous observations, we speculated that H19 and HULC are stimulated by oxidative stress
and regulate inflammation pathways involved in CCA pathogenesis. To validate this speculation, we investigated the function of H19 and HULC in CCA cells. Several inflammation pathways are involved in CCA by modifying migration and invasion; therefore, we used scratch wound healing assays to further demonstrate the function of H19 and HULC in migration potency. After 24 h in culture, CCA cells with forced expression of H19 or HULC (Additional file 1: Figure S2A) dramatically promoted wound closure compared to the control (Fig. 2a), suggesting that H19 and HULC have migration regulating roles. We also performed migration assays and invasion assays to validate the migration and invasion abilities of H19 and HULC. Consistence with the scratch wound healing assays, H19 and HULC significantly expedited migration (2.0-fold for H19 and 3.3-fold for HULC, Fig. 2b) and invasion (1.77-fold for H19 and 1.64-fold for HULC, Fig. 2c), which further strengthened the migration and invasion roles of H19 and HULC in CCA. Furthermore, we also investigated the migration and invasion abilities of RBE cells when knocking down H19 or HULC. The results showed that the migration (Additional file 1: Figure S3A) and invasion (Additional file 1: Figure S3B) abilities of RBE cells were decreased when H19 or HULC was knocked down (Additional file 1: Figure S2B). These data imply that H19 and HULC may regulate the inflammation factors regulating migration and invasion after stimulating oxidative stress.

**H19 and HULC sponge a set of miRNAs and the inflammation factors**

We further investigated the potential target genes of H19 and HULC and the underlying regulatory mechanisms. Previous studies showed that both H19 and HULC are similar to ceRNAs [38] in human embryonic kidney cells and human embryonal carcinoma cells by sponging let-7 and miR-372, respectively [27, 39]. To determine whether H19 and HULC have similar roles in CCA, we focused on the potential functions of the let-7 and miR-372 families in CCA cells (Additional file 1: Figure S4A). We used bio-informational predicting and found the let-7 might target IL-6, one of the most important inflammatory factors in the pathogenesis, growth and migration of CCA [40], and miR-372 and miR-373 might target CXCR4, a chemokine receptor involved in CCA progression and metastasis [41] (Fig. 3a). Subsequently, a

![Fig. 2](image-url) Enforced H19 and HULC expression increased the ability of migration and invasion in CCA cells. a Scratch wound healing assays of CCA cells. b Migration assays of CCA cells. c Invasion assays of CCA cells
Fig. 3 (See legend on next page.)
In conclusion, our results suggest that two inflammation-associated lncRNAs in CCA, H19 and HULC, target inflammatory genes, including IL-6 and CXCR4, using ceRNA methods and primarily regulated cell migration and invasion (Fig. 5). These lncRNAs might serve as key regulators in CCA.

**Discussion**

The importance of several dysregulated lncRNAs has been implicated in developmental regulation and disease pathogenesis, and the function of many dysregulated lncRNAs have been reported in various cancers [28, 43–45]. However, no studies on lncRNAs associated with CCA migration and invasion have been reported. In this study, we found that H19 and HULC targeted IL-6 and CXCR4, respectively, by sponging miRNAs in CCA cells. The study provides new insight into the mechanism linking lncRNA function with CCA and may serve as novel targets for the development of new countermeasures of CCA.

A number of studies have revealed that IL-6 was released by autocrine and paracrine by malignant cells or immune cells and functioned in tumorigenesis [40, 46, 47]. Generally, IL-6 was secreted by noncancer stem cells in low-attachment culture conditions and enriched Oct4 gene expression by activating the IL-6R/JAK2/STAT3 signaling pathway in chronic inflammatory disease, such as cholangiocarcinoma [47, 48]. Our previously study also found treatment of CAA cells with IL-6 can active paracrine IL-6/STAT3 pathway in inflammation and CCA initiation [49]. IL-6 is a multifunctional inflammatory cytokine that plays a major role in the response of cholangiocytes to inflammation [40, 50]; and increased concentrations of IL-6 during inflammation...
Fig. 4 (See legend on next page.)
in the biliary tract stimulates several pathways, including the JAK-STAT pathway, the p38 MAPK pathway, the PI3-kinase pathway, and is involved in survival and growth of malignant cholangiocytes [40, 51]. CXCR4 is a chemokine receptor involved in several inflammatory processes and diseases, including CCA [52], and induces CCA cell migration and invasion via the ERK 1/2 and Akt pathways [53, 54]. However, a single IncRNA can regulate multiple target genes, and a powerful inflammation-related IncRNA might also target multiple inflammation genes, for example, H19 targets TGFβ1 in prostate cancer cells [55]; TGFβ1 is a key regulator in CCA inflammation [1], suggesting the pivotal role of inflammation regulation by H19. Therefore, we speculate that H19 and HULC may act as primary regulators of other downstream inflammation genes that initiate and sustain CCA.

The growing evidences have indicated that H19 involved in both proliferation and differentiation processes, together with epithelial to mesenchymal transition (EMT) and also mesenchymal to epithelial transition (MET), suggesting its contribution in tumor both initiation and progression [56]. H19 has an evolutionary conservation secondary structure, suggesting its structure-dependent functions, which include binding to enhancer of zeste homolog 2 (EZH2) [57], and interacting with the methyl-CpG-binding domain protein 1 (MBD1) and recruiting it to some of its targets that maintain the repressive H3K9me3 histone marks in their loci [58]. H19 has also been recently found to interact with P53 proteins, and led to its inactivation, indicating that H19

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**Fig. 4** The mechanism of H19 and HULC that regulated genes. **a** AGO2 rip-qPCR of miRNAs and IncRNAs in RBE cells. The enrichment of miRNAs binding with AGO2 were used as the positive controls and U6 as the negative. **b** AGO2 rip-qPCR of miRNAs and IncRNAs in QBC939 cells. Triplicate experiments were analyzed by mean ± SD, p value < 0.001, ***, <0.01, **, <0.05, *.

**Fig. 5** Schematic representations of pathways modulated by H19 and HULC in CCA cells. H19 and HULC are up-regulated by oxidative stress induced by viruses or liver flukes and sponging miRNAs, such as let-7a/let-7b and miR-372/miR-373. This increases the expression of inflammation-related genes, including IL-6 and CXCR4, resulting in abnormal inflammation responses and pathogenesis of CCA.
plays a role in tumorigenesis [59]. HULC was reported to promote the proliferation and regulated cell cycle of HCC through down-regulation of the tumor suppressor gene CDKN2C (p18) and involved in signaling pathways including ATM/ATR and p53 [60, 61]. In this study, we revealed the potential pathway of H19 that is targeted by let-7a/let-7b, causing the partial inactivation of IL-6 in its mediated inflammatory responses triggered by oxidative stress in cholangiocarcinoma. We also found that HULC was targeted by miR-372 and miR-373 and further activated the chemokine receptor CXCR4 in the progression of migration and invasion in cholangiocarcinoma cells. The results suggest a potential pathway of lncRNAs participated in the progression of CCA.

Conclusions
Our study revealed that H19 and HULC were up-regulated by oxidative stress and target IL-6 and CXCR4 via ceRNA patterns of sponging let-7a/let-7b and miR-372/miR-373, respectively. Furthermore, overexpressed H19 and HULC might promote cell migration and invasion, suggesting that these lncRNAs might be the chief culprits of CCA pathogenesis and progression (Fig. 5). Our results provide new insight into the mechanism linking lncRNA function with CCA and may serve as novel targets for the development of new countermeasures of CCA.

Additional file

Additional file 1: Figure S1. The expression of heme oxygenase-1 (HO-1) had increased significantly under the oxidative stress condition. HO-1 stimulated by short-term oxidative stress using H2O2 (A) and long-term oxidative stress using glucose oxidase (B) in QBC939, SK-cha-1, and RBE cells. Figure S2. qPCR validations of forced expressed H19 and HULC in CCA cells (A) and knockdown of IncRNAs, H19, and HULC in RBE cells by siRNAs. Figure S3. Knockdown of H19 and HULC expression decreased the ability of migration and invasion in CCA cells. Migration assays of CCA cells (A). Invasion assays of CCA cells (B). Figure S4. MiRNAs targeted inflammation-associated genes, IL-6 and CXCR4. A. Schematic representation of the interaction between miRNAs and H19 or HULC (A). Let-7a/let-7b and miR-372/miR-373 suppressed luciferase activity of vectors with 3’-UTR of IL-6 and CXCR4 in RBE cells (B). Figure S5. The histogram of miRNA down-regulation by transfecting the miRNA inhibitors into RBE cells under oxidative stress. The expression levels of IL-6 (A) and CXCR4 (B) increased significantly. Figure S6. Western Blot validations of forced expressed IL-6 and CXCR4 in CCA cells. CXCR4-1 and CXCR4-2 represent two alternative splicing of CXCR4. The right histogram of relative gray values represented the protein level. Figure S7. Forced IL-6 and CXCR4 expression increased the ability of invasion in CCA cells. Scratch wound healing assays of CCA cells (A). Migration assays of CCA cells (B). Invasion assays of CCA cells (C). Table S1. Primer sequence for qPCR and vector construction. (PDF 1109 kb)

Abbreviations
CCA: Cholangiocarcinoma; ceRNAs: Competing endogenous RNAs; EMT: Epithelial to mesenchymal transition; EZH2: Enhancer of zeste homolog 2; H3K27: Histone H3 Lys 27; HBV: Hepatitis B virus; HO-1: heme oxygenase-1; IncRNAs: Long non-coding RNAs; MBD1: Methyl-CpG-binding domain protein 1; MET: Mesenchymal to epithelial transition; miRNAs: microRNAs; NF-kB: Nuclear factor kappa B; RIP: RNA immunoprecipitation; ROS: Reactive oxygen species; STAT3: Signal transducer and activator of transcription 3

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Availability of data and materials
The dataset supporting the conclusions of this article is included within the article.

Authors’ contributions
WTW, HY, and WPP conceived and carried out the experiment and wrote the paper. BWH and ZHC performed the in vitro experiments. YQC conceived the experiment and co-wrote the paper. All authors read and approved the final manuscript.

Competing interests
The authors declare that they have no competing interests.

Consent for publication
Not applicable.

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Not applicable.
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