Long Non-coding RNA SNHG12, a New Therapeutic Target, Regulates miR-199a-5p/Klotho to Promote the Growth and Metastasis of Intrahepatic Cholangiocarcinoma Cells

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Background: Small nucleolar RNA host gene 12 (SNHG12) is a newly identified long non-coding RNA (lncRNA) whose involvements have been explored in several cancers. Our study aimed to explore the functions of SNHG12 on intrahepatic cholangiocarcinoma (ICC) progression and its interaction with miR-199a-5p and Klotho.

Methods: RT-PCR was performed to examine the expressions of SNHG12, miR-199a-5p and Klotho in ICC cells. Cell counting kit-8 (CCK-8), colony formation assays and transwell assays were applied to analyze the proliferation, migration and invasion of ICC cells. Luciferase assays, RIP assays and RNA pull-down assays were carried out to demonstrate the direct binding relationships among SNHG12, miR-199a-5p and Klotho. The xenograft nude models were applied to test the effects of SNHG12 on ICC tumor growth.

Results: The expression of SNHG12 and Klotho was distinctly increased in ICC cells, while miR-199a-5p expressions were decreased. Functionally, the silence of SNHG12 inhibited the proliferation and metastasis of ICC cells, while miR-199a-5p overexpression exhibited an opposite result. Mechanistically, Knockdown of SNHG12 significantly suppressed the expressions of miR-199a-5p by sponging it, and then increased Klotho expression. The final in vivo experiments suggested that the silence of SNHG12 distinctly inhibited tumor growth.

Conclusion: Our findings indicated that SNHG12 inhibited cell proliferation and metastasis process of ICC cells through modulating the miR-199a-5p/Klotho axis and it is expected to become a potential therapeutic target for ICC.

Keywords: lncRNA SNHG12, intrahepatic cholangiocarcinoma, miR-199a-5p, Klotho, metastasis
INTRODUCTION

Intrahepatic cholangiocarcinoma (ICC) is a heterogeneous group of malignancies that occur at any location along the biliary tree (1, 2). In the last 20 years, it has been reported that the incidence and mortality of ICC is increasing in China (3). So far, one of the most effective treatments remains surgical removal for ICC patients (4). However, for these patients with advanced stages, operative treatments provide limited chances (5). It is necessary to develop novel adjuvant therapies for the improved prognosis of ICC patients. However, the poor understanding of the mechanisms underlying the development and progression of ICC restricts the developments of novel approaches. Thus, it is necessary to establish molecular modulators influencing developments and progression of ICC, which will promote the developments of efficient therapies.

Long non-coding RNA (LncRNA) is defined as an endogenous RNA with molecules > 200 nt in length (6). Unlike other non-coding RNAs including miRNAs, the potential involvements of LncRNAs in human illnesses remains largely uncovered. More and more studies have indicated that LncRNAs were involved in the life activity via modulating the expression of various genes through complex mechanisms (7, 8). Their regulatory effects on chromatin remodeling, gene transcription and intercellular signaling were also reported in several studies (9, 10). In addition, in tumor biology, the vital roles of LncRNAs were also demonstrated by a growing volume of literature (11). The dysregulation of LncRNAs has been confirmed to display tumor-promotive or suppressive roles in various types of tumors (12, 13). Nevertheless, to date, the functional roles of LncRNAs in the progression of ICC are unclear.

Small nucleolar RNA host gene 12 (SNHG12), located at the p35.3 region on chromosome 1, is a newly identified tumor-related lncRNA (14). In recent years, its dysregulated expression was frequently reported in several types of tumors (15–17). In some specific tumors, the oncogenic roles of SNHG12 were also reported in vitro and in vivo experiments, such as renal cell carcinoma and cervical cancer (18, 19). However, the expressions and functions of SNHG12 in ICC were rarely reported.

MATERIALS AND METHODS

Cell Lines and Transfection

Control cell line (BEC) and ICC cell lines (CCLP1, TFK-1, HuCCT1 and RBE) was obtained from the Institute of Cell Research, Chinese Academy of Sciences (Shanghai, China). All cells were cultured in DMEM medium (Procell, Yita Biology, Daxing, Beijing, China). The culture media were all supplemented with 10% fetal bovine serum (MedChemExpress, Technology, Wuhan, Hubei, China). The culture media were all supplemented with 10% fetal bovine serum (MedChemExpress, Pudong, Shanghai, China) and 50 μ/ml of penicillin and 50 mg/ml of streptomycin (Invitrogen, Shenneng, Guangdong, China). In a humidified incubator with 5% CO2 at 37°C, all cells were maintained.

CCLP1 and TFK-1 cells were transfected with sh-SNHG12-1, sh-SNHG12-2 and sh-NC, miR-199a-5p mimics and miR-NC, miR-199a-5p inhibitors and NC inhibitors, pcDNA3.1/SNHG12 and pcDNA3.1, or pcDNA3.1/Klotho and pcDNA3.1 (all, Genepharma, Shanghai, China). Lipofectamine 2000 (Invitrogen, Shenneng, Guangdong, China) was used for the transfection.

RNA Isolation and Quantitative Real-Time PCR

According to the manufacturers’ instructions, TRIzol (Invitrogen, Yita Biology, Daxing, Beijing, China) was applied to extract total RNAs from ICC cells. Using the Reverse Transcriptase (Transgene, Beijing, China), the complementary DNA (cDNA) preparation was carried out based on the standard procedures. Then, RT-PCR assays were conducted on ABI 7600 using SYBR Premix ExTaq II kit (Takara, Haidian, Beijing, China). After RT-PCR data were corrected, the 2−ΔΔCt methods were used to calculate them. GAPDH acted as the endogenous control to normalize the data. Table 1 showed the primer sequences. The PCR was performed in triplicate.

Cell Viability Assays

Cellular viabilities were monitored using Cell Counting Kit-8 assays (LaiFuSai Technology, Nanjing, Jiangsu, China). The TFK-1 and CCLP1 cells after the transfection were grown in 96-well plates. Based on the manufacturer’s protocol, cellular viabilities were assessed every 24 h. The densities of every group were examined at 450 nm.

For colony formation assays, TFK-1 and CCLP1 cells were trypsinized after the transfection. Approximately 600 cells were plated in each well of 6-well plates and maintained for 14 days to form colonies. Subsequently, the colonies were fixed with methanol (Jizhihua Company, Pudong, Shanghai, China) and stained with 0.1% crystal violet (Baoman Biology, Yangpu, Shanghai, China). For each assay, three independent experiments were carried out.

Cell Invasion and Migration

TFK-1 and CCLP1 cells in migration and invasion ability were examined applying the transwell assays. For migration assays, 5 × 104 cells were seeded into the upper chamber of a transwell insert (pore size, 8 μm) in RPMI-1640 medium. The lower chamber was covered using the above medium containing 8% FBS. For invasion assays, on the lower surface of the chamber, the matrix

| Genes         | Sequences (5′-3′)                       |
|--------------|----------------------------------------|
| SNHG12: forward | TCTGTTGATGAGGACTTCCT                  |
| SNHG12: reverse | ACCTCCCTACGTACACACT                   |
| miR-199a-5p: forward | GCCAAGCGCAGTTGCGAC                    |
| miR-199a-5p: reverse | GTCCAGGTTGCGAGTTTCC                   |
| Klotho: forward | TGAAGGAGGACACCAGTGGGTGTT             |
| Klotho: reverse | CATGGATGCTTGGGCTCTAA                   |
| GAPDH: forward | AAAACGGATCACCAGGAGAGGA                  |
| GAPDH: reverse | AAGGAAATGAGAGCAGCAG                      |
| U6: forward | TCGCGGTGCTCTGGCTGCGA                   |
| U6: reverse | CCAATGCAGGGTCGAGGT                      |

Table 1: The primers used in this study for RT-PCR.
gel was covered. In addition, the rest of experiment steps were in line with the migrated assays. After incubation for 24 h, the TFK-1 and CCLP1 cells which have finished migration and invasion were fixed with crystal violet. Finally, under a microscope, the cells were captured and counted.

Subcellular Fraction
According to the protocol of PARIS™ Kit (Invitrogen, Haidian, Beijing, China), the cytoplasmic and nuclear fractions of TFK-1 and CCLP1 cells were isolated. RT-PCR was applied to examine the isolated RNAs.

RNA-BINDING PROTEIN IMMUNOPRECIPITATION ASSAY
Following the manufacturer’s directory, the Magna RIP RNA-Binding Protein Immunoprecipitation Kit (Millipore, Nanjing, Jiangsu, China) was applied to perform the RIP assays. Briefly, TFK-1 and CCLP1 cells at 75–85% confluency gathered and lysed using RIP lysis buffer. 150 μl of the extracts of TFK-1 and CCLP1 cells were then incubated with RIP buffer which contained magnetic beads conjugated to negative control normal mouse IgG or human anti-Ago2 antibody. Then, to digest the protein, Proteinase K was used to incubate the collected samples for 24 h. After collecting the immunoprecipitated RNAs, RT-PCR assays were applied to examine the purified RNAs.

RNA Pull-Down Assays
The bio-miR-199a-5p or bio-NC probes were dissolved in washing and binding buffer, followed by the incubation using Dynabeads M-290 Streptavidin (Solaibao Biology, Haidian, Beijing, China) at 25°C for 15 min. After generating probe-coated beads, the probe-coated beads were applied to incubate cell lysates of TFK-1 and CCLP1. Then, the RNA complexes were collected for further examination of the relative enrichment of RNAs by the use of RT-PCR.

Luciferase Reporter Assays
StarBase and TargetScan were used to predict the putative binding sites between SNHG12 and miRNA-199a-5p, miRNA-199a-5p and Klotho. The sequences of wild type SNHG12 (SNHG12-WT), mutant type SNHG12 (SNHG12-MuT), wild type 3′-UTR of Klotho (WT-Klotho), and mutant type 3′-UTR of Klotho (MUT-Klotho) containing the putative binding site with miR-199a-5p were amplified and cloned into the pmiRGLo vector (Promega). Then, by the use of Lipofectamine 2000 (Invitrogen, Shenzhen, Guangdong, China), luciferase reporter vectors were co-transfected with miR-NC or miR-199a-5p mimics into TFK-1 and CCLP1 cells. A dual-luciferase reporter assay system examined the luciferase activity.

Animal Experiments
All BALB/c nude mice aged 6–7 weeks and weighing 20–22 g were used for in vivo assays. The animal studies were performed after we received the approval from Zhejiang Provincial People’s Hospital. About 5.0 × 10⁶ CCLP1 cells in the log phase were inoculated into the right flank of every nude mouse by subcutaneous injection. Sh-NC and sh-SNHG12-1 groups include six mice. The size of tumor was determined every 4 days. After 28 days, all mice were sacrificed. After their tumors were excised, the tumor weight was examined. Moreover, the tumor volumes were calculated using the formula: length × width² × 0.5.

Statistical Analysis
All statistical analyses were performed using SPSS 17.0 (SPSS, Chicago, USA). The significance of difference between groups was analyzed by two-tailed Student’s t-test one-way analysis or two-way analysis of variance. A two-sided p-value of <0.05 was considered to be statistically significant.

RESULTS
The Distinct Upregulation of SNHG12 in ICC Cells and Its Oncogenic Roles
To identify the functional lncRNA in ICC progression, we searched TCGA datasets and focused on SNHG12 which was distinctly overexpressed in ICC specimens (n = 36) compared with non-tumor specimens (n = 9) (Figure 1A). Then, RT-PCR assays also showed that SNHG12 expression was distinctly increased in four ICC cell lines compared with BEC cells (Figure 1B). Then, to explore the function of SNHG12 in the ability of ICC cells, we down-regulated SNHG12 expression in CCLP1 and TFK-1 cells using sh-SHG12-1 and sh-SHG12-2, which was confirmed by RT-PCR (Figure 1C). CCK8 and clone formation assays revealed that silence of SNHG12 observably suppressed the viability of CCLP1 and TFK-1 cells (Figures 1D,E). Transwell experiments indicated that knockdown of SNHG12 distinctly suppressed the metastasis ability of CCLP1 and TFK-1 cells (Figures 1F,G). On the contrary, we observed that overexpression of SNHG12 promoted the proliferation, migration and invasion of CCLP1 and TFK-1 cells (Figures 2A–E).

MiR-199a-5p Suppressed the Proliferation and Metastasis of ICC Cells
It has been demonstrated that many cytoplasmic lncRNAs act as ceRNAs by competitively binding miRNAs. By the use of subcellular fractionation, our group showed that SNHG12 was expressed both in the cytoplasm and nucleus, and a larger proportion of SNHG12 was observed in the cytoplasm (Figure 3A). Using StarBase v2.0 software (screening conditions: pan-cancer number: 5, medium, clip data), several miRNAs were predicted to have a high probability of binding to SNHG12 (Figure 3B). Among the 5 miRNAs, miR-199a-5p has been demonstrated to serve as an anti-oncogene in some cancers (20, 21). Thus, we chose it for further study. RT-PCR assays indicated that miR-199a-5p expressions were distinctly decreased in ICC cells (Figure 3C). Further qRT-PCR showed that miRNA-199a-5p mimics effectively overexpressed miR-199a-5p in CCLP1 and TFK-1 cells compared to miR-NC group (Figure 3D). Gain-of-function experiments indicated that overexpression of miR-199a-5p distinctly suppressed the proliferation, migration and invasion of CCLP1 and TFK-1 cells (Figures 3E–H).
FIGURE 1 | Increased SNHG12 expression and its oncogenic roles in ICC cells. (A) Higher levels of SNHG12 were observed in ICC specimens than normal specimens by amazing TCGA datasets. (B) RT-PCR determined the expressions of SNHG12 in ICC cells and BEC cells. (C) SNHG12 expression was examined in TFK-1 and CCLP1 cells transfected with sh-NC, sh-SNHG12-1 or sh-SNHG12-2. (D) Cell proliferation analysis. TFK-1 and CCLP1 cells were detected for 24, 48 and 72 h after transfection. (E) Proliferation of TFK-1 and CCLP1 cells transfected with sh-SNHG12-1 or sh-SNHG12-2 as determined by colony formation assays. (F,G) Transwell assays were used to detect cellular invasion and migration in SNHG12-knockdown TFK-1 and CCLP1 cells. *p < 0.05, **p < 0.01.
SNHG12 Promotes ICC progression via Sponging miR-199a-5p

To explore whether a ceRNA mechanism between SNHG12 and miR-199a-5p existed, we conducted RIP assays and found that SNHG12 and miRNA-199a-5p coexisted in CCLP1 and TFK-1 cells (Figure 4A). StarBase v2.0 showed miR-199a-5p contained a putative binding site with SNHG12 (Figure 4B). Luciferase reporter assays indicated that miR-199a-5p mimics distinctly decreased the luciferase activity of SNHG12-WT, while the relative luciferase activity of SNHG12-MUT remained unchanged (Figure 4C). Functional assays revealed that miR-199a-5p inhibition reversed the suppressor effects of SNHG12 knockdown on the proliferation, migration and invasion of CCLP1 and TFK-1 cells (Figures 4D–G).

Increased Expression of Klotho in ICC via a ceRNA Mechanism With SNHG12

To demonstrate the ceRNA network, our group was required to discover the target genes of miR-199a-5p. Using starbase 2.0, Klotho may be a potential one. We examined the expressions of Klotho in ICC cells using RT-PCR and observed Klotho was distinctly increased (Figure 5A). Then we explore whether the dysregulation of SNHG12 may exhibit an effect on the expression of Klotho. The results of RT-PCR suggested that knockdown of SNHG12 suppressed Klotho, while overexpression of SNHG12 exhibited a reverse effect (Figure 5B). Moreover, RIP assays demonstrated that SNHG12, miR-199a-5p and Klotho coexisted in CCLP1 and TFK-1 cells (Figure 5C). Pull-down assays also demonstrated that biotinylated miR-199a-5p could pull down SNHG12 and Klotho in CCLP1 and

FIGURE 2 | Overexpression of SNHG12 promoted the proliferation and metastasis of TFK-1 and CCLP1 cells. (A) RT-PCR assay was used to detect SNHG12 level in SNHG12 overexpression plasmids transfected TFK-1 and CCLP1 cells. (B,C) Overexpression of SNHG12 distinctly promoted the proliferation of TFK-1 and CCLP1 cells. (D,E) Transwell assay indicated the migrated and invaded cell number in TFK-1 and CCLP1 cells. **p < 0.01.
miR-199a-5p overexpression promoted the proliferation and metastasis of TFK-1 and CCLP1 cells. (A) Relative SNHG12 levels in nuclear and cytosolic fractions of TFK-1 and CCLP1 cells. (B) Bioinformatics analysis to select relevant miRNAs. (C) The histogram of miR-199a-5p expressions in four ICC cells and BEC cells. (D) RT-PCR examined the expressions of miRNA-199a-5p in TFK-1 and CCLP1 cells transfected with miR-NC or miR-199a-5p mimics. (E) CCK-8 assays revealed that OD values of TFK-1 and CCLP1 cells were distinctly declined when transfected with miR-199a-5p mimics. (F) Cell clone number was distinctly decreased when transfected with miR-199a-5p mimics. (G,H) Transwell assays of migration and invasion of TFK-1 and CCLP1 cells after treatment with miR-199a-5p mimics or miR-NC. **p < 0.01.
TFK-1 cells (Figure 5D). Besides, luciferase reporter assays demonstrated the predicted binding sites between miR-199a-5p and Klotho (Figures 5E,F). In addition, we performed rescue experiments with overexpressed Klotho, finding that overexpression of Klotho reversed the distinct suppression of SNHG12 knockdown on the proliferation, migration and invasion of CCLP1 and TFK-1 cells (Figures 6A–D). Finally, Xenografts model was applied for the determination of oncogenic roles of SNHG12 in vivo. As presented in Figure 6E, the tumor growth speed was slower on nude mice after subcutaneous injection with sh-SNHG12-1 than sh-NC group. Moreover, our group showed that the tumor volume and weight were distinctly lessened in sh-SNHG12-1 group compared with sh-NC group (Figures 6F,G).

![Figure 4](link)

**FIGURE 4** Knockdown of miR-199a-5p reversed the distinct suppression of SNHG12 knockdown on the ICC progression. (A) RIP assay to confirm the coexistence of SNHG12 and miR-199a-5p in TFK-1 and CCLP1 cells. (B) Schematic outlining the predicted binding sites between miR-199a-5p and SNHG12. (C) miR-199a-5p mimics markedly reduced luciferase activity in SNHG12-wild not in SNHG12-Mut in TFK-1 and CCLP1 cells. (D–G) Rescue experiments were used to determine the efficacy of miR-199a-5p knockdown on SNHG12 down-regulation via CCK-8, colony formation assay and transwell assays. **p < 0.01.**
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FIGURE 5 | Klotho is a target gene of miR-199a-5p. (A) Four ICC cells exhibited an increased expression of Klotho compared with BEC cells using RT-PCR. (B) RT-PCR for the expression of Klotho in TFK-1 and CCLP1 cells transfected with sh-NC, sh-SNHG12-1, pcDNA3.1 or pcDNA3.1/SNHG12. (C) RIP assays to verify the coexistence of three molecules. (D) RNA pull-down assay to test enrichment of SNHG12 and Klotho pulled down by miR-199a-5p. (E) Schematic representation of the predicted target site for miR-199a-5p in SNHG12. (F) Luciferase reporter assay in TFK-1 and CCLP1 cells, co-transfected with the reporter plasmid and the indicated miRs. **p < 0.01.

DISCUSSION

In recent years, several studies have reported the potential effects of SNHG12 on several tumors. For instance, Liu et al. reported that SNHG12 was highly expressed in renal cell carcinoma and its overexpression through the transfection of pcDNA3.1/SNHG12 promoted the proliferation and invasion of renal cell carcinoma cells via increasing CDCA3 expression (18). In cervical cancer, SNHG12 was shown to exhibit a high level, and its knockdown was observed to modulate the radiosensitivity of cervical cancer via upregulating CDK1 expression through sponging miR-148a (19). The similar effects of SNHG12 acting as a tumor promotor were also observed in other tumors, such as esophageal squamous cell carcinoma cells and prostate cancer (22, 23). These
FIGURE 6 | The upregulation of Klotho reversed functional depletion caused by SNHG12 knockdown. (A) CCK-8 assays were applied to determine the proliferation of TFK-1 and CCLP1 cells cotransfected with sh-NC, sh-SNHG12-1 or sh-SNHG12-1 + pcDNA3.1/Klotho. (B) Colony formation assays were used to detect the proliferation of TFK-1 and CCLP1 cells cotransfected with sh-NC, sh-SNHG12-1 or sh-SNHG12-1 + pcDNA3.1/Klotho. (C–E) Transwell assays were applied to determine the migration and invasion of TFK-1 and CCLP1 cells cotransfected with sh-NC, sh-SNHG12-1 or sh-SNHG12-1 + pcDNA3.1/Klotho. (F,G) Tumors derived from mice in two different groups were presented. (F,G) Tumor volume and weight in the xenograft mice from the SNHG12-knockdown group and the control group. **p < 0.01.
findings highlighted the important effects of SNHG12 in tumor progression. In this study, we searched TCGA datasets, finding that SNHG12 was an overexpressed lncRNA in ICC, which was also confirmed in four cell lines using RT-PCR. Then, we firstly provided evidence that knockdown of SNHG12 distinctly suppressed the proliferative and metastatic ability of ICC cells, while overexpression of SNHG12 exhibited an opposite effect. Our findings filled in the gaps in the fields of the potential effects of SNHG12 on ICC progression.

Then, we explored the potential mechanisms involved in the oncogenic roles of SNHG12 on ICC progression. It has been demonstrated that many cytoplasmic lncRNAs act as competing endogenous RNAs (ceRNAs) via binding miRNAs (24). In ICC, several lncRNAs such as lncRNA MT1JP and lncRNA AGAP2-AS1 have been shown to exhibit their tumor-related effects via sponging miRNAs to modulate the expression of various genes involved in tumor progression (25, 26). Previously, SNHG12 was also reported to act as a ceRNA in several tumors, such as cervical cancer and oral squamous cell carcinoma (19, 27). These findings encouraged us to explore the ceRNA mechanisms about SNHG12. Using Bioinformatics analysis, we screened five miRNAs. Importantly, among the five candidates, miR-199a-5p has been shown to suppress the proliferation and metastasis of hepatocellular carcinoma cells (21). Thus miR-199a-5p was used for further experiments. We found overexpression of miR-199a-5p suppressed the proliferation and metastasis of CCLP1 and TFK-1 cells. In mechanistic studies, we observed the coexistence of miR-199a-5p and SNHG12 in ICC cells and the binding sites between them. More importantly, rescue experiments confirmed that knockdown of miR-199a-5p reversed the anti-oncogenic roles of SNHG12 knockdown. These findings suggested SNHG12 may exhibit its tumor promoter effects via sponging miR-199a-5p.

To explore the possible mechanisms of miRNAs, establishing their functional targets counts a great deal. Thus, we searched internet databases, finding that klotho may be a potential target of miR-199a-5p. Klotho gene, encoding a 130-kDa transmembrane protein, is located on chromosome 13q12 (28). It has been demonstrated that Klotho is frequently expressed in the distal tubule of the renal, and less frequently in some other human specimens (29). In recent years, many studies have reported Klotho as an oncogene in several tumors, including hepatocellular carcinoma (30, 31). In this study, we also provided evidence that Klotho expression was increased in ICC cells. We firstly provided evidence which demonstrated the location and binding situation of SNHG12, miR-199a-5p and Klotho, demonstrating the ceRNA mechanism. Rescue experiments also confirmed Klotho overexpression saved functional attenuation caused by down-regulation of SNHG12. Finally, in vivo assays also demonstrated SNHG12 served as a tumor promoter in ICC.

**CONCLUSION**

Our findings demonstrated that SNHG12 facilitated the growth and metastasis of ICC cells by sponging miR-199a-5p and regulating Klotho expression, and provided a potential marker and therapeutic target for ICC patients.

**DATA AVAILABILITY STATEMENT**

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

**ETHICS STATEMENT**

The animal study was reviewed and approved by Zhejiang Provincial People's Hospital Animal Care Committee.

**AUTHOR CONTRIBUTIONS**

QH and H-GY designed and supervised the study. H-GY, QH, T-pW, and S-aH performed most experiments and wrote the manuscript. C-zH and C-hJ helped to perform parts of the experiments. QH and T-pW analyzed the data and revised the article. All authors contributed to the article and approved the submitted version.

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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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