High Expression of IncRNA AFAP1-AS1 Promotes the Progression of Colon Cancer and Predicts Poor Prognosis

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Abstract

Long non-coding RNAs (lncRNAs) are dysregulated in various cancers. However, the clinical relevance and functional roles of AFAP1-AS1 in colon cancer (CC) have not been clarified. We analyzed the lncRNA expression patterns in Gene Expression Omnibus (GEO) datasets and the Cancer Genome Atlas (TCGA) RNA-seq datasets, and found that the expression level of AFAP1-AS1 was significantly elevated in CC tissues. High levels of AFAP1-AS1 were associated with poor disease-free survival and overall survival in CC patients. In vitro experiments demonstrated that AFAP1-AS1 knockdown significantly inhibited the cell invasive and migration capability in CC cell line HT-29. AFAP1-AS1 knockdown also increased the expression of E-cadherin and ZO-1 while inhibited the expression of Vimentin, MMP9, ZEB1 and β-catenin, suggesting that AFAP1-AS1 is involved in the epithelial-mesenchymal transition (EMT) process of CC. Further studies confirmed that AFAP1-AS1 knockdown also affected the actin-cytokeratin signaling pathway. Thus, AFAP1-AS1 might be a potential novel diagnostic marker and therapeutic target for CC.

Key words: long non-coding RNA (lncRNA), AFAP1 antisense RNA 1 (AFAP1-AS1), colon cancer (CC), metastasis, prognosis

Introduction

Colon cancer (CC) is a common digestive system’s cancer that ranks fourth in cancer-associated mortality in the world and has increasing morbidity each year [1, 2]. Although encouraging progress for diagnosing and treating CC has been achieved, the underlying mechanisms of tumor development have yet to be fully elucidated.

Long non-coding RNAs (lncRNAs) are a kind of non-coding RNA longer than 200 nucleotides and lack significant open reading frames (ORFs) [3, 4]. Previous studies show that lncRNAs function in many biological processes, including cell differentiation, proliferation, and apoptosis [5, 6]. Recently, studies have shown that dysregulated lncRNAs also play important roles in human cancers [7-13]. Several lncRNAs are upregulated in CC tissues and cell lines relative to normal samples and therefore serve as oncogenes. BCAR4 promotes tumor growth and
metastasis in CC by activating the Wnt/β-catenin signaling pathway [14]. HOTAIR is associated with the epithelial-mesenchymal transition (EMT) in CC and functions as a prognostic marker [15]. HNF1A-AS1 greatly enhances cell metastasis [16]. However, some lncRNAs, such as HOXB-AS3 [17] and CASC7 [18], are down regulated in CC, indicating they may act as tumor suppressers and potential targets for clinical diagnosis, prognosis and therapy.

In this study, we found a candidate CC biomarker, AFAP1-AS1, by analyzing lncRNA gene expression data in the TCGA RNA-Seq and GEO datasets. AFAP1-AS1 was significantly elevated in CC tissues relative to normal tissues. We aimed to determine the expression level and prognostic significance of AFAP1-AS1 in CC as well as investigate the underlying molecular mechanisms.

**Materials and methods**

**Data mining and analysis**

The lncRNA expression data and clinical data of CC tissues and normal tissues were obtained from the Gene Expression Omnibus (GEO) profile database (http://www.ncbi.nlm.nih.gov/geo/, accession number GSE39582 [19] and GSE37364 [20]). The TCGA data of lncRNA levels (RNA-Seq dataset) in CC patients were obtained from the Cancer RNA-Seq Nexus database [21] (CRN, http://syslab4.nchu.edu.tw/index.jsp) and the clinical prognosis data were obtained from the GEPIA database [22] (Gene Expression Profiling Interactive Analysis, http://gepia.cancer-pku.cn/).

**Cell culture**

The human CC cell lines SW480, SW620, HCT116 and HT-29 were maintained in our laboratory. And cells were cultured in RPMI 1640 medium (GIBCO, USA) supplemented with 10% fetal bovine serum (FBS, GIBCO, USA), 100 U/ml penicillin and 100 ug/ml streptomycin (Invitrogen, USA) at 37°C in a humidified incubator air with 5% CO2.

**siRNA transfection**

The sequences of the two siRNAs that targeted AFAP1-AS1 were siRNA1, 5'-GGGCTTCAATTTACAAGCAT-3’ and siRNA2, 5'-CTCTATCGCTCAACA CGTT-3’. Sequences of non-target negative controls (NC) were obtained from Life Technologies. The cells were cultured in 6-well plates over night. And then these cells were transfected with either 50 nM of siRNA1 or siRNA2 or NC using Lipofectamine RNAiMAX Reagent (Invitrogen, Breda, The Netherlands) in accordance with the manufacturer’s instructions.

**qRT-PCR**

The total RNA was extracted using Trizol agent according to the manufacturer’s instructions. SYBR PremixExTaq II kit (Takara, Dalian, China) was used to carry out the reactions. The CFX96 Real-Time PCR Detection System (Bio-Rad, Hercules, CA, USA) was used to measure the relative expression levels of each gene. The gene-specific primers used were as follows: AFAP1-AS1, 5’-AACCTGTTAGGAGGAGAAGA-3’ and 5’-CACACAGGGAAATGAGGAGG-3’; E-cadherin, 5’-GAGAACCGATCCAGCATA-3’ and 5’-AACAGACCCCTCTAGAGACACG-3’; Vimentin, 5’-TGCCCAACCGAAGACAGATG-3’ and 5’-GCCTTCAATTTACAAGCAT-3’; MMP-9, 5’-AGG GGTCTTGTTCCAACCTC-3’ and 5’-AGCATTTACC GTCCAGGATGATG-3’; ZEB1, 5’-GATATGAGATGAC CGATCGTCAG-3’ and 5’-CTGGTCTCTTACGAGT GCC-3’; β-actin, 5’-TCACCAACTGGGACGACATG-3’ and 5’-GTCACCGAGTCATCCAGAT-3’. β-actin was used as an endogenous control.

**Wound healing assay**

We performed wound healing assays to assess the migration capacity of tumor cells as described previously [23-26]. Briefly, after transfection, the cells were seeded into 6-well plates (5×10^5 cells/well) and incubated for 24 h. When the cell cultures reached 95% confluency, a 10 μl tip was used to create a scratch in the cell monolayer. Images were captured at different time points after scratching (0 h, 24 h and 48 h) by microscopy to measure gap closure after wounding.

**Transwell invasion assays**

Transwell Cell Culture Inserts (BD Biosciences, New Jersey, USA) were used to measure the invasive capacity [27-29]. In brief, HT-29 cells (1×10^5 cells in 100 μl of 2% FBS medium) were seeded on the top chamber and then placed into 24-well plates. The bottom well contained 800 μl of growth medium with 15% FBS. After the cells were incubated for 36 h at 37°C, the cells on the upper surface were removed, while the cells that invaded through the filter pores were fixed in 4% paraformaldehyde for 30 min and stained with gentian violet. Invaded cells were imaged using a microscope, and the cell number was counted in 5 randomly selected high-power fields.

**Western blotting**

The cells were harvested after 48h of treatment, and proteins were extracted in three decontamination lysis steps. Protein was quantified with a BCA protein assay kit, separated by 10% SDS-PAGE gels and then transferred onto PVDF membranes (Millipore, Billerica, MA, USA). We then followed standard
procedures using the following antibodies: E-cadherin, β-catenin, Vimentin, MMP9, ZEB1 and ZO-1 (Cell Signaling Technology, Danvers, MA, USA); MPRIP, CTTN, RHOA, RHOC, RAB1B and β-actin (Proteintech, Wuhan, China). Immunoreactive bands were visualized by electrochemiluminescence. Visualization of β-actin expression was used as a loading control.

**Statistical analysis**

All the numerical data are expressed as the mean ± standard deviation (SD). Student’s t-tests were used to analyze the data. Curve fitting analyses were performed with GraphPad Prism Software (GraphPad Software, San Diego, CA, USA), and the results of the analyses were considered significant if \( P < 0.05 \).

**Results**

**LncRNA AFAP1-AS1 is upregulated in CC**

To explore the clinical relevance of AFAP1-AS1, we first investigated its expression in two publicly available CC datasets (GSE39582 and GSE37364). AFAP1-AS1 was significantly upregulated in CC tissues compared to normal tissues (Figure 1A &1B). We next validated AFAP1-AS1 expression in an independent CC RNA-Seq dataset (TCGA-COAD). The results revealed that CC tissues have higher AFAP1-AS1 expression relative to normal tissues (Figure 1C).

**Association between AFAP1-AS1 expression and CC patient survival**

Based on the data obtained from GEPIA, we analyzed the prognostic value of AFAP1-AS1 in CC in the TCGA-COAD RNA-Seq dataset. The patients were classified into two groups according to the AFAP1-AS1 expression level: the low AFAP1-AS1-expressing group (n = 105), in which the expression of AFAP1-AS1 was lower than the median value, and the high AFAP1-AS1-expressing group (n = 135), in which the expression of AFAP1-AS1 was higher than the median value. Compared with the patients in the low expression group, patients expressing high levels of AFAP1-AS1 had significantly shorter overall survival (OS) and disease-free survival (DFS) (Figure 2A & 2B).

**Knockdown of AFAP1-AS1 negatively regulates CC cell migration and invasion**

We measured the expression level of AFAP1-AS1 in 4 CC cell lines (SW480, SW620, HCT116, and HT-29) by qRT-PCR. Since HT-29 had the highest AFAP1-AS1 expression level (Figure 3A), we selected this cell line for further study. Two siRNAs targeting AFAP1-AS1 were transfected into HT-29 cells. The second siRNA (siRNA2) had a greater knockdown effect as measured by qRT-PCR (Fig. 3B). To elucidate the role of AFAP1-AS1 in CC, siRNA2 or a negative control (NC) siRNA was transfected into HT-29 cells. Migration studies revealed that the gap distance in the siRNA2-treated group was much longer than the NC group, suggesting that the knockdown of AFAP1-AS1 inhibited cell migration ability (Figure 3C & 3D). Transwell matrigel invasion assay was used to assess the effects of AFAP1-AS1 knockdown on cell invasion. As expected, the number of invading cells in the AFAP1-AS1 knockdown group was much lower than the number in the NC group (Figure 3E &3F).

**Silencing AFAP1-AS1 influences the expression of EMT-related genes**

To explore how AFAP1-AS1 regulates CC cell migration and invasion, we measured the expression level of EMT-related genes, E-cadherin, vimentin, MMP9, ZEB1, β-catenin and ZO-1 in CC cells after AFAP1-AS1 silencing. As shown by our qRT-PCR results, silencing AFAP1-AS1 upregulated the expression of E-cadherin while inhibiting the expression of vimentin, MMP9 and ZEB1 (Figure 4A). We also measured protein levels of EMT marker proteins by western blotting. Similarly, silencing AFAP1-AS1 upregulated E-cadherin and ZO-1 protein while downregulating β-catenin, vimentin, MMP9 and ZEB1 (Figure 4B).

![Figure 1. AFAP1-AS1 is elevated in CC tissues relative to normal tissues. The AFAP1-AS1 expression data in CC tissues were obtained from the GEO datasets in accession GSE39582 (A) and GSE37364 (B). Analysis revealed that CC tissues have a higher AFAP1-AS1 expression level than normal tissues. (C) Comparison of AFAP1-AS1 expression levels in CC tissues and normal tissues from the TCGA-COAD RNA-Seq dataset within the Cancer RNA-Seq Nexus database.](http://www.jcancer.org)
Knockdown of AFAP1-AS1 influences the actin-cytokeratin signaling pathway

Previous studies have demonstrated that down regulation of AFAP1-AS1 suppressed tumor migration and invasion by regulating EMT-related genes. Future studies should investigate which signaling pathways are involved in tumor progression. In a previous study, we showed that RHOA, RAC2, RAB10 and RAB11A were significantly upregulated, while RHOC, RAB11B and LASP1 were down regulated after silencing AFAP1-AS1 in nasopharyngeal carcinoma cells [30]. We therefore measured these molecules in CC cells. Western blotting analysis showed that CTNN, RHOA and RAB1B were elevated, while MPRIP was lower after silencing AFAP1-AS1 (Figure 4C).

Discussion

Emerging evidence indicates that lncRNAs play important roles in multiple physiological and pathological processes, including tumor development [31-37]. AFAP1-AS1, also known as LOC84740, is located on chromosome 4p16.1 and acts as an antisense lncRNA regulator of the AFAP1 gene [30]. AFAP1-AS1 was originally identified in esophageal adenocarcinoma [38], is frequently upregulated in CC and serves a vital role in a variety of tumor types, including pancreatic cancer, hepatocellular carcinoma and lung cancer [39-42]. In liver cancer, AFAP1-AS1 is highly expressed and promotes cell growth by activating the RHOA/RAC2 pathway [43].

Previous studies confirmed via qRT-PCR that AFAP1-AS1 was frequently highly expressed in colorectal cancer (CRC) tissues and that AFAP1-AS1 is a potential diagnostic biomarker for CRC patients [44]. In the present study, we collected data from different platforms to determine the expression tendency of AFAP1-AS1 and its prognostic value in CC. Our analysis is consistent with that of previous studies; AFAP1-AS1 was elevated in CC tissues. Our findings also confirmed that AFAP1-AS1 expression was highly correlated with poor prognosis. These findings suggest that AFAP1-AS1 may act as an oncogene in CC. We found that HT-29 cells had a higher expression level of AFAP1-AS1. Therefore, HT-29 cells were used to further explore the role of AFAP1-AS1 in CC. We found AFAP1-AS1 knockdown significantly suppressed cell migration and invasion ability. We also determined that AFAP1-AS1 acts as a tumor promoter. EMT was characterized by cancer cell migration and invasion. E-cadherin suppression is the first step of EMT initiation, resulting in reduced cell adhesion. Our data suggest that AFAP1-AS1 knockdown causes the upregulation of E-cadherin while inhibiting Vimentin, MMP9 and ZEB1 mRNA expression. Furthermore, AFAP1-AS1 knockdown increases E-cadherin and ZO-1 protein levels while decreasing β-catenin, Vimentin, MMP9 and ZEB1. These findings provide evidence for the anti-metastasis action of AFAP1-AS1 silencing.

Cell migration and invasion are important aspects of cancer progression, and studies have highlighted the key signaling pathways that mediate tumor cell migration and invasion through regulation of the cytoskeleton assembly [45-47]. In a previous study, we confirmed that some molecules associated with the actin-cytokeratin signaling pathway were
regulated by AFAP1-AS1 in nasopharyngeal carcinoma [30]. We also measured the molecules key to the actin-cytokeratin signaling pathway in the present study and found that CTTN, RHOA and RAB1B were elevated while MPRIP decreased after silencing AFAP1-AS1. These results suggest that AFAP1-AS1 knockdown might affect tumor cell invasion and migration through regulation of the actin-cytokeratin signaling pathway.

In conclusion, we found and confirmed that AFAP1-AS1 is upregulated in CC. AFAP1-AS1 knockdown significantly suppressed CC progression by regulating the actin-cytokeratin signaling pathway and EMT-related genes, indicating that AFAP1-AS1 may be a novel therapeutic target for CC.
Figure 4. AFAP1-AS1 knockdown influences EMT-associated gene expression and the actin-cytokeratin signaling pathway. (A) qRT-PCR-measured mRNA expression of E-cadherin, vimentin, MMP9 and ZEB1 in siRNA2-transfected HT-29 cells. (B) The protein expression of E-cadherin, vimentin, MMP9, ZEB1, β-catenin and ZO-1 were detected by Western blotting. (C) Western blotting analysis showed that CTTN, RHOA and RAB1B were elevated while MPRIP decreased after silencing AFAP1-AS1.

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Competing Interests
The authors have declared that no competing interest exists.

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