Knockdown of Urothelial Carcinoma-Associated 1 Suppressed Cell Growth and Migration Through Regulating miR-301a and CXCR4 in Osteosarcoma MHCC97 Cells

Genglong Zhu,* Xialei Liu,* Yonghui Su,† Fangen Kong,‡ Xiaopeng Hong,* and Zhidong Lin†

*Department of Hepatobiliary Surgery, The Fifth Affiliated Hospital of Sun Yat-Sen University, Zhuhai, P.R. China
†Department of General Surgery, The Fifth Affiliated Hospital of Sun Yat-Sen University, Zhuhai, P.R. China
‡Department of Neurosurgery, The Fifth Affiliated Hospital of Sun Yat-Sen University, Zhuhai, P.R. China

Liver cancer is one of the most common malignancies in the world and a leading cause of cancer-related mortality. Accumulating evidence has highlighted the critical role of long noncoding RNAs (lncRNAs) in various cancers. The present study aimed to explore the role of lncRNA urothelial carcinoma-associated 1 (UCA1) in cell growth and migration in MHCC97 cells and its underlying mechanism. First, we assessed the expression of UCA1 in MHCC97 and three other cell lines by RT-qPCR. Then the expression of UCA1, miR-301a, and CXCR4 in MHCC97 cells was altered by transient transfection. The effects of UCA1 and miR-301 on cell viability, migration, invasion, and apoptosis were assessed. The results revealed that UCA1 expression was relatively higher in MHCC97 cells than in MG63, hFOB1.19, and OS-732 cells. Knockdown of UCA1 reduced cell viability, inhibited migration and invasion, and promoted cell apoptosis. However, the effect of UCA1 knockdown on cell growth and migration was blocked by miR-301a overexpression, whose expression was regulated by UCA1. We also found that miR-301a positively regulated CXCR4 expression. CXCR4 inhibition reversed the effect of miR-301a overexpression on cell growth and migration. Moreover, miR-301a activated the Wnt/β-catenin and NF-κB pathways via regulating CXCR4. The present study demonstrated that UCA1 inhibition exerted an antigrowth and antimigration role in MHCC97 cells through regulating miR-301a and CXCR4 expression.

Key words: Urothelial carcinoma-associated 1 (UCA1); MicroRNA-301a; CXCR4; Cell apoptosis; Migration and invasion; Liver cancer

INTRODUCTION

Liver cancer is one of the most common malignancies in the world and is a leading cause of cancer-related mortality. Hepatocellular carcinoma (HCC) represents a major global health problem, since it accounts for 70%–90% of all primary liver cancers worldwide. Epidemiologic evidence demonstrates that even though HCC primarily occurs in Asia and Africa, the medical and economic burden of HCC will still soar drastically in Western populations during the next decade. Thus, exploration of the underlying mechanisms of liver carcinogenesis is urgently needed, as well as the development of novel diagnostic biomarkers and therapeutic strategies for patients with HCC.

Recently, increasing evidence has highlighted the role of a group of long non-protein-coding RNAs (lncRNAs) in carcinogenesis, and it has been suggested that lncRNAs could be considered as biomarkers in various cancers. lncRNAs, a class of non-protein-coding RNAs longer than 200 bp in length, have been reported to play a crucial role in the process of cancer, including cell proliferation, cell apoptosis, metastasis, and differentiation. Several lncRNAs have been reported to be involved with the development and progress of HCC. For instance, lncRNA LINC00152 has been demonstrated to be upregulated in HCC tissues and HCC cell lines, and promoted cell proliferation in HCC by targeting EpCAM via the mTOR signaling pathway. Another lncRNA, highly upregulated in liver cancer (HULC), enhances epithelial–mesenchymal transition (EMT) to promote tumorigenesis and metastasis of HCC via regulating the microRNA 200a-3p (miR-200a-3p)/ZEB1 signaling pathway. Human urothelial carcinoma-associated 1 (UCA1) gene is located in chromosome 19p13.12, which has three exons and encodes two
Several studies have revealed that UCA1 is highly expressed in various cancers, such as bladder cancer, breast cancer, and colorectal cancer, and it promotes tumor growth and invasion. However, the role of UCA1 in HCC and its underlying molecular mechanisms are not fully elucidated.

The present study aimed to explore the role of UCA1 in the growth and metastasis of HCC cell lines and its underlying mechanism. We found that the expression of UCA1 was relatively higher in MHCC97 cells, and inhibition of UCA1 reduced cell viability, inhibited cell migration and invasion, and promoted cell apoptosis in MHCC97 cells. Moreover, we found that UCA1 inhibition might exert its antigrowth and antimigration role via regulating miR-301a and C-X-C chemokine receptor type 4 (CXCR4).

MATERIALS AND METHODS

Cell Culture and Treatment

Human liver cancer cell line MHCC97 and human osteosarcoma cell lines (MG63, hFOB1.19, and OS-732) were all obtained from the American Type Culture Collection (ATCC; Rockville, MD, USA). Cells were maintained in Dulbecco's modified Eagle’s medium (DMEM; HyClone, Logan, UT, USA) supplemented with 10% heat-inactivated fetal bovine serum (FBS; Gibco, Gaithersburg, MD, USA), 100 U/ml penicillin, and 100 μg/ml streptomycin (Life Technologies Corporation, Carlsbad, CA, USA) and were incubated at 37°C in an atmosphere of 95% air and 5% CO₂.

Transfection and Generation of Stably Transfected Cell Lines

Short hairpin RNA (shRNA) directed against human lncRNA UCA1 and CXCR4 was ligated into the U6/GFP/Neo plasmids (GenePharma, Shanghai, P.R. China) and were referred to as sh-UCA1 and sh-CXCR4. The plasmid PGPU6/GFP/neo-shControl (GenePharma) encoded with a nonsense sequence was used as the negative control (NC) of sh-UCA1 and sh-CXCR4 and were referred to as shNC and sh-NC, respectively. Meanwhile, the full-length CXCR4 sequence directed against CXCR4 was subcloned into the pEX-2 plasmid (GenePharma) and was referred to as pEX-CXCR4. Empty pEX-2 plasmid acted as an NC of pEX-CXCR4 and was referred to as pEX. The Lipofectamine 3000 reagent (Life Technologies Corporation) was used for the cell transfection following the manufacturer’s instructions. miR-301a inhibitor, miR-301a mimic, and their corresponding NCs (inhibitor control and mimic control) were synthesized by GenePharma and were transfected into MHCC97 cells with Lipofectamine 3000 reagent (Life Technologies Corporation).

Quantitative Reverse Transcription PCR (RT-qPCR)

Total RNA was extracted from cells using TRIzol reagent (Life Technologies Corporation) under the manufacturer's instructions. The One-Step SYBR® PrimeScript® PLUS RT-RNA PCR Kit (TaKaRa Biototechnology, Dalian, P.R. China) was used for the RT-qPCR analysis to detect the expression levels of UCA1. The TaqMan MicroRNA Reverse Transcription Kit and TaqMan Universal Master Mix II with the TaqMan MicroRNA Assay of miR-301a and U6 (Applied Biosystems, Foster City, CA, USA) were used for evaluating the expression levels of miR-301a. Expression level of CXCR4 was detected using the RNA PCR Kit (AMV) Ver.3.0 (TaKaRa Biototechnology) for transcription and SYBR® Premix Ex Taq™ II (TaKaRa Biototechnology) for qPCR. GAPDH and U6 were used for the normalization of mRNA and IncRNA or miRNA, respectively. The results were presented as fold changes relative to U6 or GAPDH and were calculated using the 2^−ΔΔCt method.

Cell Viability Assay

Cell viability was determined by trypan blue exclusion assay. Briefly, 1×10⁵ cells were seeded in 60-mm dishes and cultured for 48 h. Then cells were rinsed with phosphate-buffered saline (PBS), and the number of viable cells was determined by trypan blue exclusion (Beyotime Biotechnology, Shanghai, P.R. China) as previously described.

Migration and Invasion Assay

Cell migration was determined using a modified two-chamber migration assay with a pore size of 8 μm. For the migration assay, 1×10⁵ cells in 0.2 ml of serum-free medium were plated on the upper compartment of a 24-well Transwell culture chamber (Corning, Lowell, MA, USA), and the lower chamber was filled with 0.6 ml...
of medium containing 10% FBS. For the invasion assay, 1.5×10⁵ cells were plated on the upper chamber precoated with 20 μg of Matrigel (BD Biosciences, Bedford, MA, USA). After incubation for 48 h, the migrated and invaded cells in the lower chamber were fixed with 100% methanol and stained with 0.1% crystal violet (Sigma-Aldrich, St. Louis, MO, USA). The cells that did not migrate or invade through the pores were removed by cotton swabs. The migrated and invaded cells were counted in five random fields and expressed as the average number of cells per field. These experiments were done in triplicate and performed a minimum of three times.

**Apoptosis Assay**

The apoptotic cell rates were determined using an Annexin-V-Phycoerythrin (PE) Apoptosis Detection Kit (Beyotime Biotechnology). Briefly, the cells were harvested and washed with PBS three times. Then cells were resuspended with 500 μl of 1× binding buffer and were stained with 5 μl of annexin V-PE for 15 min in the dark at 37°C. Cell apoptosis was measured using a FACScan (Beckman Coulter, Fullerton, CA, USA), and the data were analyzed utilizing FlowJo software (Tree Star Inc., Ashland, OR, USA).

**Figure 2.** Knockdown of UCA1 inhibited cell growth and migration in MHCC97 cells. MHCC97 cells were transfected with shNC, sh-UCA1#1, and sh-UCA1#2. (A) The efficiency of transfection was quantified by RT-qPCR analysis. (B) Cell viability was measured by trypan blue exclusion assay. (C) Cell migration and (D) invasion were detected by Transwell migration assay. (E) Apoptotic cells and (F) the levels of apoptosis-related proteins were measured by flow cytometry and Western blot, respectively. *p<0.05, **p<0.01, ***p<0.001.
Western Blot Assay

The proteins used for Western blot were extracted using radioimmunoprecipitation assay (RIPA) lysis buffer (Beyotime Biotechnology), supplemented with protease inhibitors (Roche, Indianapolis, IN, USA). The BCA™ Protein Assay Kit (Pierce, Appleton, WI, USA) was used to quantify the concentration of proteins. Then 30 μg of protein was loaded and separated with a Bio-Rad Bis-Tris Gel system based on the manufacturer’s instructions. All the blots were transferred onto polyvinylidene difluoride (PVDF) membranes (Millipore, Billerica, MA, USA) and were blocked with 5% bovine serum albumin (BSA; Roche) for 2 h at room temperature. Subsequently, the blots were incubated with primary antibodies against Bcl-2 (#4223), Bax (#5023), caspase 3 (#9662), caspase 9 (#9502), GAPDH (#2118), p-p65 (#3033), p65 (#8242), IκBα (#4812), p-IκBα (#2859) (Cell Signaling Technology, Beverly, MA, USA), Wnt3a (ab28472), Wnt5a (ab72583), CXCR4 (ab197203), and β-catenin (ab6302) (Abcam, Cambridge, UK) overnight at 4°C. Then the blots were rinsed with TBST three times, followed by incubation with secondary antibody labeled with horseradish peroxidase for 1 h at room temperature. The membranes carrying blots and antibodies were placed in the ChemiDoc™ XRS system (Bio-Rad, Hercules, CA, USA), and 200 μl of Immobilon Western Chemiluminescent HRP Substrate (Millipore) was added to cover the membrane surface. The signals were captured and analyzed using Image Lab™ software (Bio-Rad).

Statistical Analysis

All experiments were repeated three times. The results of multiple experiments are presented as the mean ± standard deviation (SD). Statistical analyses were performed using GraphPad Prism 6 software (GraphPad Software, San Diego, CA, USA). The p values were calculated using a one-way analysis of variance (ANOVA). A value of p < 0.05 was considered to indicate a statistically significant result.

RESULTS

UCA1 Was Highly Expressed in MHCC97 Cells

We first detected the expression of UCA1 in the human liver cancer cell line MHCC97 and three other cell lines. As shown in Figure 1, the expression of UCA1 was highest in MHCC97 cells compared with the other three cell lines, human osteosarcoma MG63 and OS-732, and human osteoblast cell line hFOB1.19.

Knockdown of UCA1 Inhibited Cell Growth and Migration in MHCC97 Cells

We then investigated the role of UCA1 in cell viability, migration, invasion, and apoptosis in MHCC97 cells. The expression of UCA1 was inhibited by transfection with sh-UCA1. The efficiency of transfection was identified by RT-qPCR. As shown in Figure 2A, the level of UCA1 was significantly reduced in MHCC97 cells transfected with sh-UCA1#1 and sh-UCA1#2, and the transfection efficiency of sh-UCA1#2 was higher than that of sh-UCA1#1 (p<0.05 or p<0.001). Thus, we selected sh-UCA1#2 for the suppression of UCA1 in the following experiments. Cell viability was significantly reduced in MHCC97 cells after transfection with sh-UCA1#2 (p<0.01) (Fig. 2B). Meanwhile, knockdown of UCA1 reduced cell migration and invasion (Fig. 2C and D) (p<0.05). We also found that knockdown of UCA1 promoted cell apoptosis in MHCC97 cells, as evidenced by increasing the percentage of apoptotic cells (p<0.001) (Fig. 2E), as well as inducing the expression of the proapoptosis factors (Bax, cleaved caspase 3, and cleaved caspase 9) and reducing the Bcl-2 expression (Fig. 2F). Overall, these results revealed that knockdown of UCA1 suppressed cell growth and migration in MHCC97 cells.

miR-301a Overexpression Blocked the Effect of UCA1 on Cell Growth and Migration in MHCC97 Cells

The RT-qPCR analysis showed that the expression of miR-301a was remarkably decreased in sh-UCA1-transfected cells compared with the corresponding controls (p<0.001) (Fig. 3). Then we further investigated the role of miR-301a in the modulation of UCA1 knockdown in cell growth and migration. As shown in Figure 4A, miR-301a overexpression inhibited the cell viability reduction caused by sh-UCA1 transfection (p<0.05). Consistently, miR-301a overexpression increased cell migration and invasion even though the cells were transfected with sh-UCA1 (p<0.05 or p<0.01) (Fig. 4B and C). The results of the flow cytometry and Western blot showed that miR-301a overexpression reversed...
the promoting effect of sh-UCA1 on cell apoptosis, in that the apoptotic cell rate was reduced and the expression of proapoptotic proteins was inhibited, while anti-apoptotic protein expression was elevated ($p<0.05$ or $p<0.01$) (Fig. 4D and E). These results suggested that the modulation of UCA1 inhibition on cell growth and migration was associated with the downregulation of miR-301a.

The Expression of CXCR4 Was Positively Regulated by miR-301a

As shown in Figure 5A, the mRNA expression of CXCR4 was upregulated by miR-301a mimic, while it was downregulated by a miR-301a inhibitor ($p<0.05$). Similar results were observed in protein levels by Western blot analysis (Fig. 5B). These results showed that
miR-301a positively regulated the expression of CXCR4 in MHCC97 cells.

CXCR4 Was Involved in the Regulatory Effect of miR-301a on Cell Growth and Migration

Since we found that CXCR4 expression was regulated by miR-301a, we further explored whether CXCR4 participated in the modulation of miR-301a on cell growth and migration of MHCC97 cells. MHCC97 cells were transfected with miR-301a or cotransfected with sh-CXCR4. As shown in Figure 6A and B, the mRNA and protein expressions of CXCR4 were upregulated by miR-301a mimic (p<0.01). However, this modulation was reversed by sh-CXCR4 transfection (p<0.05), which further confirmed the regulatory relationship between CXCR4 and miR-301a. Interestingly, we found that the progrowth and promigration effect of miR-301a mimic was reversed by sh-CXCR4, as demonstrated by declining cell viability (p<0.05) (Fig. 6C) and inhibition of cell migration and invasion (p<0.05) (Fig. 6D and E), reducing apoptotic cell rate (p<0.01 or p<0.001) (Fig. 6F) as well as regulating the protein expression of apoptosis-related core factors (Fig. 6G). From the above, it indicated that miR-301a overexpression enhanced cell growth and migration through upregulation of CXCR4 expression.

miR-301a Enhanced the Activation of the Wnt/β-Catenin and NF-κB Signaling Pathway Through Upregulating CXCR4

Results in Figure 7A showed that miR-301a inhibitor suppressed the expressions of Wnt3a, Wnt5a, and β-catenin, while miR-301a mimic acted as an opposite regulatory effect. However, the inhibitive effect of the miR-301a inhibitor was blocked by CXCR4 overexpression, as pEX-CXCR4 increased the expression of the three proteins related with the Wnt/β-catenin signaling pathway. Similar regulation was found in the NF-κB signaling pathway in that the pathway was activated by miR-301a mimic while suppressed by miR-301a inhibitor. Results in Figure 7B revealed that miR-301a inhibitor suppressed the phosphorylation of IκBα and p65, and miR-301a mimic accelerated the expression of p-IκBα and p-p65. Furthermore, overexpression of CXCR4 blocked the suppressive influence on the NF-κB signaling pathway. In general, these results suggested that miR-301a enhanced the activation of the Wnt/β-catenin and NF-κB signaling pathways through upregulating CXCR4.

DISCUSSION

An increasing number of studies reveal that the aberrant expression of lncRNAs is highly involved with the progression and prognosis of cancers, serving a role as either an oncogene or a tumor suppressor gene18–21. Thus, it will be beneficial and meaningful to deeply clarify the biological and molecular mechanisms of lncRNAs in cancer. The expression of UCA1 has been reported to be highly expressed in various cancers, and it may serve the role of an oncogene13–16. In our present study, we found that lncRNA UCA1 was relatively expressed higher in the human liver cancer cell line (MHCC97 cells) compared with three human osteosarcoma or osteoblast cell lines (MG63, OS-732, and hFOB1.19). Downregulation of UCA1 decreased cell viability, inhibited cell migration, and induced cell apoptosis of MHCC97 cells. Our results were consistent with a previous study in which the expression of UCA1 has been reported to be highly expressed in various cancers, and it may serve the role of an oncogene13–16. In our present study, we found that IncRNA UCA1 was relatively expressed higher in the human liver cancer cell line (MHCC97 cells) compared with three human osteosarcoma or osteoblast cell lines (MG63, OS-732, and hFOB1.19). Downregulation of UCA1 decreased cell viability, inhibited cell migration, and induced cell apoptosis of MHCC97 cells. Our results were consistent with a previous study in which the expression of UCA1 is aberrantly upregulated in HCC tissues, and UCA1 depletion inhibited the growth and metastasis of HCC cell lines in vitro and in vivo22.

Even though the role of UCA1 in HCC has been reported previously in one study, it still remains significant to explore the underlying mechanism(s) for us
Figure 6. CXCR4 was involved in the regulatory effect of miR-301a on cell growth and migration. MHCC97 cells were transfected with sh-CXCR4 or sh-NC, or cotransfected with sh-CXCR4 and miR-301a mimic, or mimic control. The (A) mRNA and (B) protein expression levels of CXCR4 were tested by RT-qPCR analysis and Western blot analysis, respectively. (C) Cell viability, (D) cell migration, (E) invasion, and (F) apoptotic cells were assessed by trypan blue exclusion assay, Transwell migration assay, and flow cytometry assay, respectively. (G) The protein levels of apoptosis-related factors were measured by Western blot. *p<0.05, **p<0.01, ***p<0.001.

Figure 7. miR-301a enhanced the activation of the Wnt/β-catenin and NF-κB signaling pathways through upregulating CXCR4. MHCC97 cells were transfected with miR-301a inhibitor, miR-301a mimic, or their corresponding controls (mimic control and inhibitor control), or cotransfected with miR-301a inhibitor and pEX-CXCR4. The protein expression of (A) Wnt/β-catenin- and (B) NF-κB signaling pathway-related core factors was measured by Western blot.
to better understand the pathogenesis of HCC. A previous study highlighted that UCA1 contributed to the progression of HCC via directly downregulating miR-216b expression and promoting the activation of the FGFR1/ERK signaling pathway. However, our study revealed that knockdown of UCA1 exerted its antigrowth and antitmigration roles through downregulation of miR-301a and CXCR4, which is also involved in the inactivation of the Wnt/β-catenin and NF-κB signaling pathways. miR-301a, located in the human chromosome 17q22-17q23, has been previously reported to be overexpressed in many kinds of human cancers and shown to be an oncogene in gastric cancer, pancreatic cancer, liver cancer, and colorectal cancer. In HCC, miR-301a was significantly upregulated, and inhibition of miR-301a suppressed HepG2 cell growth, migration, and invasion, and induced cell apoptosis through targeting homeobox gene Gax and then modulating NF-κB expression. In the present study, we found that miR-301a was downregulated by UCA1 knockdown. Furthermore, miR-301a overexpression reversed the inhibitory effect of UCA1 knockdown on cell growth and migration, which is consistent with the previous study. However, the relationship between miR-301a and UCA1 and the underlying regulatory mechanisms need to be further explored in future studies.

CXCR4, the receptor for stromal cell-derived factor (SDF-1), plays an important role in angiogenesis and is associated with tumor progression. CXCR4 has been indicated as a critical factor for breast cancer metastasis through the interaction with SDF-1. Inhibition of CXCR4 by siRNA impairs the invasion of breast cancer cells in the Matrigel invasion assay and inhibits breast cancer metastasis in an animal model. Meanwhile, it has been reported that CXCR4 promotes the EMT process and progression of colorectal cancer. Furthermore, CXCR4 acts in colorectal cancer EMT and progression through the regulation of the Wnt/β-catenin signaling pathway. In the theoretical sense, miRNAs are involved in the mediation of gene expression via targeting their binding sites, usually in the 3′-UTR of mRNA, and this leads to posttranscriptional or translational repression. However, a recent study demonstrated that miRNA positively regulated the expression of downstream mRNA, due to its being complementary to the promoter sequences of specific genes. Qu et al. reported that miR-558 was upregulated and positively correlated with HPSE expression in neuroblastoma tissues and cell lines, as well as directly targeted the HPSE promoter to activate its transcription. Similar with this study, our results suggested that the expression of CXCR4 was positively regulated by miR-301a. This might be because miR-301a upregulated the expression of CXCR4 through directly targeting the CXCR4 promoter. However, studies are still needed to clarify this hypothesis and further explore the relationship between miR-301a and CXCR4 in the future. In addition, we found that CXCR4 inhibition blocked the effect of miR-301a overexpression on cell viability, migration, invasion, and apoptosis, which suggested that inhibition of CXCR4 might exert antigrowth and antitmetsasis effects on MHCC97 cells.

The Wnt/β-catenin signaling pathway plays critical roles in cell proliferation, differentiation, and adhesion. It has been demonstrated that CXCR4 promotes tumor progression by activating the canonical Wnt/β-catenin signaling pathway in pancreatic cancer and colorectal cancer. Furthermore, a previous study showed that upregulated miR-301a in breast cancer promoted tumor metastasis by targeting PTEN and activating Wnt/β-catenin signaling. Similarly, the NF-κB pathway has been reported to be regulated by CRCX4 and miR-301a. Thus, we hypothesized that miR-301a may regulate the activation of Wnt/β-catenin and NF-κB signaling in liver cancer cells through CXCR4. We found that miR-301a mimic enhanced the activation of the Wnt/β-catenin and NF-κB signaling pathways, while miR-301a inhibitor exerted the opposite effect. Moreover, CXCR4 inhibition reversed the regulatory effect of miR-301a on these signaling pathways. It suggested that miR-301a might activate the Wnt/β-catenin and NF-κB signaling pathways through the regulation of CXCR4.

In conclusion, the present study demonstrated that UCA1 was highly expressed in MHCC97 cells. Knockdown of UCA1 inhibited cell viability, migration, and invasion, and promoted cell apoptosis in MHCC97 cells through the regulation of miR-301a and CXCR4. Our study might help us better understand the progression of HCC and shed new light on the diagnosis and treatment for HCC.

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incRNAs, long non-coding RNAs, act as regulatory elements in the transcriptome and can influence various biological processes, including cell proliferation, apoptosis, and metastasis. They often function as competing endogenous RNAs (ceRNAs), sponging up miRNAs to relieve their repression of target genes.

In human ovarian cancer, IncRNA UCA1 has been identified as a significant biomarker. A study by Zhang et al. (2016) demonstrated that UCA1 expression is upregulated in ovarian cancer cells and contributes to tumor proliferation, cell cycle progression, and invasion. Additionally, UCA1 expression is correlated with poor prognosis in ovarian cancer patients.

In breast cancer, UCA1 has also been shown to play a critical role. Jia et al. (2014) reported that UCA1 promotes breast cancer cell proliferation and invasiveness through activation of the mTOR signaling pathway. Furthermore, UCA1 contributes to progression of hepatocellular carcinoma through inhibition of miR-216b and activation of FGFR1/ERK signaling pathway. Oncotarget 2015;6(10):7899–917.

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