STAT3-induced upregulation of long noncoding RNA HNF1A-AS1 promotes the progression of oral squamous cell carcinoma via activating Notch signaling pathway

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ABSTRACT

Long non-coding RNAs (lncRNAs) are a group of biomarkers which can regulate the biological processes of various human cancers. LncRNA HNF1A-AS1 has been reported in human cancers for its oncogenic role. This study focused on the biological function and molecular mechanism of HNF1A-AS1 in oral squamous cell carcinoma (OSCC). The high expression of HNF1A-AS1 was examined in OSCC tissues and cell lines. Kaplan Meier method revealed that high expression of HNF1A-AS1 predicted poor prognosis for patients with OSCC. Results of loss-of-function assays demonstrated that silenced HNF1A-AS1 inhibited the proliferation, migration and epithelial-mesenchymal transition (EMT) of OSCC cells. Mechanically, HNF1A-AS1 was positively regulated by the transcription factor STAT3. Recently, Notch signaling pathway has been reported in human malignancies. In this study, we analyzed the correlation between HNF1A-AS1 and Notch signaling pathway. It was uncovered that the expression of Notch1 and Hes1 (the core factors of Notch signaling pathway) was negatively regulated by HNF1A-AS1 knockdown. Rescue assays further demonstrated the positive regulatory effects of HNF1A-AS1 on Notch signaling pathway in OSCC. In conclusion, upregulation of HNF1A-AS1 induced by transcription factor STAT3 promotes OSCC progression by activating Notch signaling pathway.

Introduction

Oral squamous cell carcinoma (OSCC) is widely recognized as the commonest oral malignancy. According to the previous studies, the incidence of OSCC ranks eighth in the world range.\(^1,2\) There are about 540,000 new cases of OSCC are identified every year. However, only half of these patients can survive for 5 years after the surgery. Despite advances have been made in surgery and medicine fields, the mortality of patients with OSCC is still high.\(^3\) More seriously, in some regions, such as the Eastern Europe, the mortality of OSCC is unceasingly increasing in the past two decades. Therefore, OSCC has gradually become a major concern for social problem and human health.\(^4\) The initiation and progression of OSCC are very complex. Although more and more molecular targets have been proved to be growth factors in biological processes of OSCC, the role of lncRNAs in OSCC still need to be further explored.\(^5\)

Over the several years, micro (mi) RNAs and IncRNAs have been proved to be able to serve as important factors in both physiology and pathological processes.\(^6–9\) So far, many researchers have reported the special functions of miRNAs in OSCC.\(^10–14\) In addition, a handful of studies proved the regulatory role of IncRNAs in OSCC.\(^15–17\) LncRNA HNF1A-AS1 (HNF1A antisense RNA 1, C12 or f27) has been reported in several types of human cancers, such as bladder cancer,\(^18\) NSCLC,\(^19\) and colorectal carcinoma.\(^20\) Nevertheless, the specific biological functions of it in OSCC is still marked. At first, we examined the expression of HNF1A-AS1 in OSCC tissues and cell lines. The prognostic value of HNF1A-AS1 for patients with OSCC was identified with Kaplan Meier method. The oncogenic role of HNF1A-AS1 in OSCC was identified by conducting loss-of-function assays.

We further detected the mechanism involved in HNF1A-AS1-mediated biological functions in OSCC. Based on previous reports, transcription factors can contribute to IncRNAs’ transcription, thus enhancing the expression of IncRNAs.\(^21–24\) In this study, we found that STAT3 is a transcription activator of HNF1A-AS1. Mechanism experiments further demonstrated the activation of STAT3 on HNF1A-AS1 transcription. Previous reports showed that upregulation of IncRNAs can promote tumorigenesis and tumor progression by activating Notch signaling pathway.\(^25–27\) In this study, HNF1A-AS1 could positively regulate the activity of Notch signaling pathway in OSCC. Rescue assays further proved the interaction between HNF1A-AS1 and NOTCH signaling pathway. In summary, this study revealed that STAT3-induced upregulation of HNF1A-AS1 promoted OSCC progression by activating NOTCH signaling pathway.

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Materials and methods

Clinical samples

62 pairs of human OSCC tissues and their adjacent normal tissues were collected from patients who were diagnosed with OSCC in The Affiliated Stomatological Hospital of Nanchang University from October, 2012 to December, 2017. All patients participated in this study had signed the ethics consents before all experiments were conducted. In addition, this study had acquired the approval of the ethic committee of The Affiliated Stomatological Hospital of Nanchang University.

Cell culture and treatment

Five OSCC cell lines (CAL-27, HN5, SCC-15, SCC-9 and Tca8113) and the Keratinocytes of the oral cavity (NHOK) were all commercially obtained from American Type Culture Collection (ATCC, Manassas, VA). Cells were maintained in Dulbecco’s Modified Eagle’s Medium (DMEM, Invitrogen, Grand Island, NY, USA) which had been mixed with a complex (10% FBS + 1% L-glutamine + penicillin at a density of 25 units/ml + streptomycin at a concentration of 25 g/ml).

To activate Notch signaling, Tca8113 cells were incubated with recombinant Jagged-1/Fc (R&D Systems, Minneapolis, USA). To knock down HNF1A-AS1 or STAT3, the short hairpin RNAs (shRNAs) specially targeting HNF1A-AS1 (sh-HNF1A-AS1#1, sh-HNF1A-AS1#2, sh-HNF1A-AS1#3) or STAT3 (sh-STAT3) and negative control shRNA (sh-NC) were designed and constructed by GenePharma (Shanghai, China). Similarly, the small interfering RNA (siRNA) against JAK1 and JAK2 (si-JAK1, si-JAK2) and their negative control siRNAs (si-NC) were synthesized by GenePharma. For STAT3 overexpression, pcDNA3.1 vector containing the whole sequence of STAT3 and empty vector (pcDNA-NC) were constructed and obtained from GenePharma. All transfections were finished by using Lipofectamine 2000 (Invitrogen, USA).

RNA extraction and qrt-pcr

Based on the user guide, the Trizol reagent (Invitrogen, Carlsbad, CA, USA) was utilized for extraction of total RNA from the tissues or transfected cells. Next, with the help of the ABI PRISM 7000 Fluorescent Quantitative PCR System (Applied Biosystems, Foster City, CA, USA), quantitative real-time PCR was carried out. The expression of HNF1A-AS1 was normalized to GAPDH.

MTT assay

Briefly, SCC-15 and Tca8113 cells were cultured in 24-well plates at a concentration of 3000 cells per well. Next, we replaced media with MTT regent. The blue crystals were dissolved by DMSO. Finally, the absorbance was measured with a spectrophotometer at 570 nm.

CCK-8 assay

Based on the recommendation of manufacturers, CCK-8 assay was carried out in two OSCC cell lines. Briefly, Cells transfected with sh-NC or sh-HNF1A-AS1 were seeded into 96-well plates. 10 μL CCK-8 solutions were added into each well at different time points at 37°C. At last, the absorbance was measured with a spectrophotometer at 450 nm.

Colony formation assay

Additionally, colony formation assay was further conducted to detect cell proliferation. Before single-cell suspensions, trypsin was used for digestion of transfected OSCC cells. Then, 6-well plates were utilized for seeding cells (500 cells/well). Next, we waited for about two weeks for necessary incubation. Cells were then fixed with methanol for half an hour at room temperature. Afterwards, we stained them with 0.5% crystal violet for about 60 minutes. Finally, we used a Nikon camera (Nikon, Japan) to photograph.

Flow cytometry

SCC-15 and Tca8113 cells were harvested and collected for further experiments at 48 hours after transfection. Based on the instructions for users, cells were stained with PI with the help of the Cycle TESTTM PLUS DNA Reagent Kit (BD Biosciences). The result was analyzed with a flow cytometer (FACScan®, BD Biosciences). Finally, the percentages of the cells in different phases (G0-G1, S, and G2-M) were counted for comparison.

For apoptosis analysis, we treated OSCC cells with fluorescein isothiocyanate (FITC), Annexin V and propidium iodide (PI) avoid light at normal temperature in accordance with the recommendations of manufacturer. Subsequently, apoptotic analysis was finished by using FACScan®.

Cell migration assay

To measure the migratory ability of OSCC cells, cells were repeatedly suspended in serum-free medium (100 μL). The treated cells were then plated in the upper chamber of each insert (8-μm pore size, Corning, USA) without the membrane for the transwell migration assay. To incubate cells, the lower chamber was filled with DMEM medium which containing 10% FBS. After 12 hours, migrated cells were fixed in the lower surface of the insert. Then stained them with 0.1% crystal violet. Finally, the number of migrated cells was counted under a light microscope.

Immunofluorescence

Before the immunofluorescence assay, cells grown on glass slices. Next, fixation and permeation of cells were separately finished in 4 % formaldehyde for ten minutes and with 0.3 % Triton X-100. Next, the slices were blocked by the goat serum for a quarter of an hour at 37°C. Then incubation of anti-E-cadherin (1:80, Bioworld, MN, USA) and anti-N-cadherin (1:80, Bioworld, MN, USA) for all night at 4°C with. Samples were washed three times before the incubation of goat TRITC labeled secondary antibody (1:70, Bioworld, MN, USA) (37°C and 1 h). After stained by DAPI (Genview Inc,
Shanghai, China), the fluorescence was visualized with a microscope whose magnification was × 400.

**Western blotting analysis**

According to the recommendation of manufacturer, protein lysates were extracted from tissues or cells by a RIPA kit (Beyotime, Shanghai, China). And protein concentrations were examined by utilizing an enhanced BCA Protein Assay kit (Beyotime). Loading buffer was used to mix the samples which containing equal amounts of protein. Besides, those samples were separated by SDS-PAGE. Next, they were transferred to the PVDF membranes (Millipore, Billerica, MA, USA). These membranes were blocked by a buffer containing 5% defatted milk, and followed by the treatment of primary antibodies: the rabbit anti-Notch1 (ab8925, 1: 2,000 dilution), anti-Hes1 (ab71559, 1: 2,000 dilution), anti-STAT3 (ab68153,1: 2,000 dilution), anti-p-STAT3 (ab76315,1: 2,000 dilution) and GAPDH (1: 3,000 dilution) at 4°C overnight (8 h), followed by treatment with secondary antibody Goat Anti-Mouse IgG (ab6785, 1:2,000) conjugated to horseradish peroxidase for 2 h at room temperature. All antibodies used in this study were acquired from Abcam (San Francisco, CA, USA). To visualize blot images, chemiluminescence (Beyotime) was employed. ImageQuant LAS 4000 System (GE Healthcare, Little Chalfont, UK) was used for further analysis.

**Chromatin immunoprecipitation (chip) assay**

Chromatin immunoprecipitation (ChIP) was performed using the SimpleChIP® Enzymatic Chromatin IP Kit (CST, USA) based on the standard methods provided by supplier. SCC-15 and Tca8113 cells were treated with 4% formaldehyde to covalently crosslink the proteins to DNA for 15 min and then collected for ultrasonic to shear cross-linked chromatin DNA into two fragments. Thereafter, the complex was immunoprecipitated with anti-STAT3 antibody (ab32500, Abcam) and bound-protein G magnetic beads (Life Technologies, CA, USA). Normal rabbit immunoglobulin G (IgG) was used as a negative control. The precipitated chromatin DNA was washed and purified to recover DNA which was analyzed by qRT-PCR.

**Luciferase reporter assay**

Bioinformatics analysis was used to predict the binding sites between transcription factor STAT3 and HNF1A-AS1 promoter. Three binding sites of STAT3 in HNF1A-AS1 promoter were co-transfected into SCC-15 and Tca8113 cells along with pcDNA-STAT3 (termed STAT3), sh-STAT3 and their corresponding negative control (NC and sh-NC) using Lipofectamine 2000 (Invitrogen, CA, USA). 48 h after transfection, cells were harvested. The luciferase activity was measured using Dual-Luciferase® Reporter Assay System (Promega, Madison, WI, USA). Results were normalized to Renilla luciferase activity.

**Statistical analysis**

All data were shown as the mean ± SD from more than two independent experiments. Statistical analysis was performed using Prism5 (GraphPad, USA) software. Kaplan-Meier method was applied to generate the overall survival curves. The correlation between HNF1A-AS1 and STAT3 in OSCC tissues was determined by Spearman’s correlation analysis. We analyzed differences by utilizing Student’s t test or one-way ANOVA analysis. Data were considered statistically significant only when p < 0.05.

**Results**

**High expression of HNF1A-AS1 predicted unfavorable prognosis for patients with OSCC**

At first, we measured the expression pattern of HNF1A-AS1 in OSCC tissues and cells by performing qRT-PCR analysis. Compared with that in the normal tissues or the normal cells, HNF1A-AS1 was expressed much higher in OSCC tissues or cells (Figure 1(a-b)). According to the mean value of HNF1A-AS1 expression in 62 OSCC tissues, the tissue samples were divided into two groups (HNF1A-AS1 high expression group and low expression group). Next, the correlation between HNF1A-AS1 expression and clinicopathological features of OSCC patients was analyzed. We uncovered that the expression levels of HNF1A-AS1 was observably related with Nodal Invasion, T stage and Differentiation, but not with age, gender and other factors (Table 1). Proportional hazards method analysis revealed a fact that HNF1-AS1 is a crucial prognostic element for OSCC patients (Table 2). Kaplan Meier analysis determined the negative effect of HNF1A-AS1 expression on the overall survival rate of CCA patients (Figure 1(c)).
HNF1A-AS1 knockdown suppressed OSCC cell proliferation by inducing cell apoptosis and causing cell cycle arrest

According to the data in Figure 1(b), HNF1A-AS1 was expressed highest in SCC-15 and Tca8113 cells. Here, we silenced HNF1A-AS1 in SCC-15 and Tca8113 cells by transfecting with specific shRNAs (sh-HNF1A-AS1#1, sh-HNF1A-AS1#2, sh-HNF1A-AS1#3). As shown in Figure 2(a), the best transfection efficiency was obtained when cells were transfected with sh-HNF1A-AS1#2. Therefore, sh-HNF1A-AS1#2 was chosen to do next assays. Loss-of-function assays were then conducted in SCC-15 and Tca8113 cells. Results of MTT and CCK-8 assay showed the inhibitory effects of HNF1A-AS1 knockdown on cell proliferation (Figure 2(b,c)). Colony formation assays revealed the negative effect of HNF1A-AS1 knockdown on cell proliferation (Figure 2(d)). To further demonstrate the effect of HNF1A-AS1 on cell proliferation, we examined the expression of Ki-67 and PCNA. As presented in Figure 2(e), both Ki-67 and PCNA were downregulated in SCC-15 and Tca8113 cells transfected with sh-HNF1A-AS1#2. Flow cytometry analysis was used to identify whether HNF1A-AS1 affects cell proliferation by regulating cell cycle distribution. Cell cycle was arrested at G0/G1 phase after HNF1A-AS1 was silenced in SCC-15 and Tca8113 cells (Figure 2(f)). Whereas, cell apoptosis was accelerated by the silencing of HNF1A-AS1 (Figure 2(g)).

Knockdown of HNF1A-AS1 inhibited OSCC cell migration and epithelial-mesenchymal transition

Furthermore, we detected the potential role of HNF1A-AS1 in cell migration and EMT process. The inhibitory effect of sh-HNF1A-AS1 on cell migration was identified by transwell assay (Figure 3(a)). Subsequently, the protein levels of EMT markers were examined in OSCC cells transfected with sh-HNF1A-AS1. The result showed the increased levels of E-cadherin and β-catenin (epithelial markers) as well as the decreased levels of N-cadherin and Vimentin (mesenchymal markers) (Figure 3(b)). The results of immunofluorescence further demonstrated that silenced HNF1A-AS1 induced the EMT phenotype change into MET phenotype (Figure 3(c)).

STAT3 is a transcription activator of HNF1A-AS1

All findings above indicated the oncogenic role of HNF1A-AS1 in OSCC. Mechanism experiment and bioinformatics analysis were used to detect the mechanism which contributed to the upregulation of HNF1A-AS1. According to the data of UCSC (http://genome.ucsc.edu/index.html), STAT3 is a potential upstream transcription factor of HNF1A-AS1. STAT3 was found to be upregulated in OSCC tissues (Figure 4(a)). The relevance between HNF1A-AS1 and STAT3 in OSCC tissues was found to be positive in accordance with the correlation analysis (Figure 4(b)). Furthermore, we found that both levels of STAT3 and p-STAT3 were much higher in OSCC cell lines than that in normal cell line (Figure 4(c)). Subsequently, the mRNA level of STAT3 was found to be higher in OSCC cell lines (Figure 4(d)). Intriguingly, STAT3 positively regulated the HNF1A-AS1 expression (Figure 4(e)). Therefore, we hypothesized that STAT3 might upregulate HNF1A-AS1 by activating the transcription of HNF1A-AS1 in OSCC cells. The motif of STAT3 obtained from JASPAR (http://jaspar.genereg.net/) was shown in Figure 4(f). These five binding sites in part 1 (P1) or part 2 (P2) of HNF1A-AS1 promoter were chosen to do subsequent experiments (Figure 4(g)). ChiP assay further demonstrated the affinity of STAT3 to the P1 of HNF1A-AS1 promoter (Figure 4(h)). Since there are three sites in P1, the luciferase reporter assay was conducted to determine the binding condition of these three sites. As shown in Figure 4(i), the

| Variable       | Category       | P-value  |
|----------------|----------------|----------|
| HNF1A-AS1      | Low            | 0.014*   |
| Gender         | Male           | 0.125    |
| Age            | < 50           | 0.917    |
| Differentiation| Well/moderately| 0.048*   |
| Tumor Stage    | < 2            | 0.109    |
| Nodal Invasion | Positive       | 0.224    |
| Smoking        | Yes            | 0.057    |
| Drinking       | Yes            | 0.840    |

Proportional hazards method analysis showed a positive, independent prognostic importance of HNF1A-AS1 expression (*P = 0.014*), in addition to the independent prognostic impacts of T Stage (*P = 0.010*) and Differentiation (*P = 0.048*). *P < 0.05 was considered statistically significant.

Table 1. Correlation between the expression of HNF1A-AS1 and clinicopathological features of OSCC patients. (n = 62).

Table 2. Multivariate analysis of prognostic parameters in patients with OSCC by Cox regression analysis.

CANCER BIOLOGY & THERAPY
luciferase activity of three sites was decreased by STAT3 knockdown but increased by STAT3 overexpression, indicating the transcription activation of STAT3 on HNF1A-AS1. JAKs are known as activators of STAT3. Here, we examined the effect of silenced JAK1 or JAK2 on STAT3 activation. As displayed in Figure 4(j), both levels of STAT3 and p-STAT3 were significantly decreased by silenced JAK1, indicating the positive effect of JAK1 on STAT3 activation. Furthermore, we detected the effect of JAK1 knockdown on STAT3-mediated luciferase activity. The result showed that the positive effect of STAT3 on the luciferase activity of three binding sites was partially attenuated by JAK1 knockdown (Figure 4(k)). All these data showed that STAT3 upregulated HNF1A-AS1 by promoting the transcription activity of HNF1A-AS1.

**HNF1A-AS1 enhanced the activity of notch signaling in OSCC cells**

Notch signaling pathway has been reported to be a crucial factor in the progression of human cancers. In this study, we tried to investigate the interaction between HNF1A-AS1 and Notch signaling. The protein levels of Notch1 and Hes1 (two factors of Notch signaling pathway) were obviously higher in tumor tissues than that in the normal tissues (Figure 5(a)). Moreover, the protein levels of Notch1 and Hes1 were higher in SCC-15 and Tca8113 cells than that in NHOK cell (Figure 5(b)). The protein levels of Notch1 and Hes1 were decreased in SCC-15 and Tca8113 cells transfected with sh-HNF1A-AS1 compared with cells transfected with sh-NC (Figure 5(c)). Therefore, we confirmed that HNF1A-AS1 enhanced the Notch signaling in OSCC cells.

**HNF1A-AS1 facilitated OSCC progression by activating notch signaling pathway**

To detect interaction between HNF1A-AS1 and Notch signaling pathway in OSCC progression, rescue assays were conducted in Tca8113 cells. Firstly, MTT and colony formation assay demonstrated that the proliferation ability weakened by sh-HNF1A-AS1 was rescued by Jagged-1/Fc (activator of Notch signaling) (Figure 6(a-b)). Additionally, stagnant cell cycle caused by sh-HNF1A-AS1 was recovered by Jagged-1/Fc.
Similarly, increased apoptosis rate of HNF1A-AS1-downregulated Tca8113 cell was reduced again by adding Jagged-1/Fc (Figure 6(d)). Transwell assay was also carried out in Tca8113 cells which had been treated with sh-HNF1A-AS1 and Jagged-1/FC. It is obvious that the inhibitory effects of sh-HNF1A-AS1 in cell migration were mediated by Jagged-1/FC (Figure 6(e)). Furthermore, we uncovered that the inhibitory effect of sh-HNF1A-AS1 on EMT progress was partially abrogated by Jagged-1/FC (Figure 6(f)).

Discussion

According to the difference in size, noncoding RNAs (ncRNAs) can be divided into two types, one is small ncRNAs (< 200 nt), the other one is lncRNAs (> 200 nt). In recent years, lncRNAs has gradually become a hotspot in biological research. Although more and more lncRNAs have been unmasked, countless lncRNAs still need to be further explored. It has been revealed that lncRNAs can regulate cellular processes, therefore contributing to the progression of human cancers. LncRNA HNF1A-AS1 has been reported in human malignancies due to its oncogenic properties. In this study, we examined the expression of HNF1A-AS1 in OSCC tissues and cell lines. Unsurprisingly, HNF1A-AS1 was upregulated in OSCC tissues and cell lines. Kaplan Meier methods was used to determine the prognostic value of HNF1A-AS1 for patients with OSCC. Since high expression of HNF1A-AS1 predicted poor prognosis for OSCC patients, the potential oncogenic role of HNF1A-AS1 in OSCC was uncovered. To validate the oncogenic function of HNF1A-AS1 in OSCC, loss-of-function assays were carried out. According to the experimental results, silenced HNF1A-AS1 inhibited cell proliferation by inducing cell apoptosis and causing cell cycle arrest. Moreover, cell migration and EMT was suppressed by the knockdown of HNF1A-AS1. Hereeto, we confirmed that HNF1A-AS1 acted as an oncogene in OSCC.

More and more studies have revealed that the dysregulation of lncRNAs contributes to the progression and development of human cancers. In this study, we uncovered that upregulation of HNF1A-AS1 contributes to the malignancy of OSCC. It is necessary to investigate the mechanism contributed to the upregulation of HNF1A-AS1 in OSCC. Based on the bioinformatics analysis, STAT3 was found to be a potential transcription activator for HNF1A-AS1. Mechanism experiments further demonstrated the affinity of STAT3 to
Figure 4. STAT3 is a transcription activator of HNF1A-AS1. A. The expression of STAT3 in OSCC tissues was examined by qRT-PCR. B. The relevance between the expression of STAT3 and that of HNF1A-AS1 in OSCC tissues was analyzed by Pearson correlation analysis. C. The levels of STAT3 and p-STAT3 were detected in OSCC cells and normal cell line. D. The mRNA level of STAT3 in OSCC cell lines and one normal cell line was detected with qRT-PCR. E. The expression of HNF1A-AS1 was examined in response to STAT3 overexpression or knockdown. F. The motif of STAT3 binding sequence was obtained from JASPAR. G. Top five binding sites of STAT3 in part 1 (P1) and part 2 (P2) of HNF1A-AS1 promoter were predicted and obtained from JASPAR. H. ChIP assay further demonstrated the affinity of STAT3 to the part 1 (P1) of HNF1A-AS1 promoter. I. The luciferase reporter assay was conducted to determine the binding condition of three sites of P1. J. The levels of STAT3 and p-STAT3 were detected in cells transfected with si-JAK1 or si-JAK2. K. The luciferase reporter assay was conducted in two OSCC cells transfected with STAT3 expression vector and si-JAK1. **P < 0.01 vs. control group.
the promoter region of HNF1A-AS1. Therefore, we confirmed that STAT3-induced lncRNA HNF1A-AS1 promoted OSCC progression.

It has been reported that lncRNAs can affect cell activities in human cancers by regulating the signal of some signaling pathway. For instance, lncRNA HOTAIR could activate the AKT/
mTOR signaling pathway, thereby enhancing the proliferative or migratory ability of osteosarcoma cells. LncRNA MRCCAT1 improves cell metastasis in clear cell renal cell carcinoma by activating p38-MAPK signaling. LncRNA HNF1A-AS1 has also been proved to be able to interact with Wnt/beta-catenin signaling pathway in both colorectal cancer and osteosarcoma. Notch signaling pathway is widely considered as a highly conserved cell signaling pathway. It was capable of participating in multiple aspects of cell growth and development, including proliferation, apoptosis and metastasis. Additionally, Notch signaling pathway has been revealed to be one of the commonest activated signaling pathway in tumors, especially in gastrointestinal malignancies, liver cancer and pancreatic cancer. Some researchers found that the aberrant activation of Notch signaling might be induced by mutation or amplification of Notch signaling components. The regulatory function of Notch signaling in genes’ expression often promote tumor formation. Therefore, in this study, we explored the regulatory relationship between HNF1A-AS1 and Notch signaling in OSCC. We found that both HNF1A-AS1 and Notch signaling were upregulated in OSCC cells. Silenced HNF1A-AS1 could inactivate Notch signaling, this result means that HNF1A-AS1 might positively regulate Notch signaling in OSCC cells. Rescue assays were further conducted to examine the interaction between HNF1A-AS1 and Notch signaling pathway. The inhibitory effects of sh-HNF1A-AS1 on cell proliferation, migration and EMT progress was rescued by Notch signaling activator. Therefore, we confirmed that HNF1A-AS1 positively modulated Notch signaling, thus promoting OSCC progression. In conclusion, upregulation of HNF1A-AS1 induced by STAT3 accelerated the progression of OSCC by activating Notch signaling pathway. Our findings will be helpful for finding novel potential therapeutic targets for OSCC.

Disclosure of Potential Conflict of interest

No potential conflicts of interest were disclosed.

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