

The RNA N6-Methyladenosine Demethylase FTO Promotes Head and Neck Squamous Cell Carcinoma Proliferation and Migration by Increasing CTNNB1

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Objective: In this study, we aimed to investigate the role of RNA N6-methyladenosine demethylase fat mass and obesity-associated protein (*FTO*) in head and neck squamous cell carcinoma (HNSCC).

Methods: Clinical data downloaded from The Cancer Genome Atlas (TCGA) database were used to analyze the relationship between mRNA levels of *FTO*, *METTL3*, *METTL14*, and *ALKBH5*, and the overall survival in cancer and para-cancer datasets. *FTO* expression in tumor and normal tissues was compared using immunohistochemistry, and its relationship with overall survival was analyzed based on the Kaplan–Meier method. The FaDu cell line with high *FTO* levels was chosen from five HNSCC cell lines for further experiments. *FTO* was verified as an oncogene in HNSCC by in vitro loss-of-function and overexpression studies, cell proliferation assay, wound healing assay, and identification of expression changes of epithelial–mesenchymal transition (EMT)-related markers. Catenin beta 1 (*CTNNB1*) was confirmed as a downstream target gene of *FTO* with additional methods like the GEPIA online tool, qRT-PCR, Western blotting, and dot blot assay.

Results: We found that *FTO* expression was significantly upregulated in HNSCC datasets and tissues. Increased *FTO* expression indicated a trend towards poor prognosis and was found to promote disease proliferation and migration. Mechanistically, cell proliferation assay, wound healing assay, and identification of expression changes of EMT-related markers demonstrated that *FTO* could act as an oncogene in HNSCC. *FTO* expression was significantly correlated with *CTNNB1* expression. Moreover, it exerted a tumorigenic effect by increasing *CTNNB1* expression in an m⁶A-dependent manner.

Conclusion: *FTO* promotes head and neck squamous cell carcinoma proliferation and migration by increasing *CTNNB1* in an m⁶A-dependent manner.

Keywords: *FTO*, m⁶A, HNSCC, *CTNNB1*

Introduction

Head and neck squamous cell carcinoma (HNSCC) is the sixth most common type of malignant tumor in the world, with approximately 600,000 newly diagnosed cases per year. The reason for its high mortality is HNSCC recurrence and metastasis, and most patients are diagnosed with HNSCC at an advanced stage.^{1,2} Therefore, exploring the molecular mechanisms regulating HNSCC recurrence and metastasis is likely to be of great significance for improving patient prognosis.

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N6-methyladenosine (m⁶A) modification, which is methylation at the nitrogen-6 position of adenosine bases in mRNA, is reversible and occurs widely (>25%) as a post-transcriptional modification of mRNA in eukaryotes.³ This modification has an important role in regulating RNA transactions, including RNA splicing, stability, transport, and translation.⁴⁻⁷ The m⁶A modification is mainly mediated by the “writers” methyltransferase-like (*METTL3/14*),⁸⁻¹⁰ Wilms tumor 1-associated protein (WTAP),¹¹ RNA binding motif protein 15/15B (RBM15/15B),¹² and KIAA1429,¹³ whereas “erasers” include fat mass and obesity-associated (*FTO*)¹⁴ and alkB homolog 5 (ALKBH5).¹⁵ “Readers” that recognize m⁶A methylated transcripts include YTH domain family proteins (YTHDF1-3, YTHDC1-2), heterogeneous nuclear ribonucleoprotein (HNRNP), and IGF2 mRNA binding proteins (IGF2BPs).^{16,17} The m⁶A modification was proven to regulate the occurrence and development of various cancers and was found to play an important role in stem cell differentiation and embryonic development.^{4,5} To date, some studies have showed that the alterations of m⁶A on specific RNA sites in HNSCC and several m⁶A regulatory genes could influence the tumorigenesis and act as prognostic biomarkers.^{18,19} Positive correlation was found between selected RNA methylation machinery gene expression and m⁶A abundance on total RNA.²⁰ RBM15, as a m⁶A-related regulator, was significantly upregulated in Laryngeal squamous cell carcinoma and relevant to the prognosis.²¹ Identified as m⁶A regulatory gene family, the expression of IGF2BP family could be an important role in tumor growth of HNSCC.²² However, relevant mechanisms of m⁶A RNA modification in HNSCC remain largely unknown.

FTO was first identified as a gene associated with obesity and energy metabolism and recently was proven to be the RNA m⁶A eraser.²³ The specific role of *FTO* in tumor recurrence and metastasis varies among different types of cancers. Some studies have shown that *FTO* has tumor suppressor activity and is inversely correlated with tumor development in renal cell carcinoma, intrahepatic cholangiocarcinoma, hepatocellular carcinoma, ovarian cancer, and colorectal cancer.²⁴⁻²⁸ In contrast, *FTO* has also been proven to play an oncogenic role in acute leukemia, cervical cancer, melanoma, and breast cancer through its mRNA maturation regulatory function.^{4,29-31} However, the role of *FTO* in HNSCC as an m⁶A demethylase remains poorly studied.

In this study, we first demonstrated that *FTO* expression was upregulated in HNSCC tissues and that increased *FTO* expression indicated a trend towards poor prognosis.

Then, we ascertained that *FTO* promotes tumor proliferation and migration in vitro by increasing the expression of the downstream target gene *CTNNB1* by increasing its mRNA stability. Overall, our findings indicate the critical role of *FTO* in HNSCC proliferation and migration.

Materials and Methods

Bioinformatics Analysis

Clinical data, comprising that of 44 normal and 520 HNSCC tissues, were downloaded from The Cancer Genome Atlas (TCGA) database (<https://portal.gdc.cancer.gov/>). The data were used to analyze the relationship between *FTO*, *METTL3*, *METTL14*, and *ALKBH5* mRNA levels; the overall survival in cancer and para-cancer datasets was used by the UALCAN database (<http://ualcan.path.uab.edu/cgi-bin/ualcan-res.pl>). We used the GEPIA website (<http://gepia.cancer-pku.cn/>) for correlation analysis between *FTO* and *CTNNB1* in HNSCC.

Clinical Samples

Fifty-eight samples of hypopharyngeal and laryngeal squamous cell carcinomas, and adjacent normal tissues were collected from January 2016 to June 2019 at the Shanghai General Hospital. The samples were frozen at -80°C before RNA extraction and immunohistochemistry (IHC) analysis. We compared the expression levels of *FTO* in tumor tissues and normal tissues according to IHC results. The relationship between overall survival and the expression level of *FTO* was also analyzed. This study was conducted in accordance with the Declaration of Helsinki and approved by the ethics committee of Shanghai General Hospital (2016KY175-4). Patients had given their signed informed consent prior to the study.

Immunohistochemistry Assay

The tissue was first embedded in paraffin. Next, a paraffin microtome was used to cut the tissue to sections with uniform thickness, and the sections were fixed on a clean glass slide. The sections were put into xylene and graded concentrations of alcohol for the removal of paraffin. After dewaxing was completed, sections were cleaned and treated in citrate buffer for antigen retrieval. Next, 3% H₂O₂ was added for 10 min at 25°C to eliminate the endogenous peroxidase activity. The sections were rinsed with distilled water and PBS, and then were blocked with 5–10% normal goat serum (diluted in PBS). Subsequently, they were incubated with the

primary antibody at 4°C overnight. Following washing in PBS, sections were incubated with secondary antibody at 25°C for 20 min. After an additional wash with PBS, sections were treated with an appropriate amount of alkaline phosphatase-labeled streptavidin working solution at 37°C for 10–30 min. The color was developed with the developer for 3–15 min and the sections were rinsed with distilled water, followed by counterstaining, dehydration, transparentizing, and mounting. Evaluation of *FTO* expression was performed with the histological scoring method by three independent pathologists using the light microscope. To evaluate the IHC staining, the staining intensity was stratified into four levels, namely, 0, 1, 2, and 3, which were denominated as absent, weak, moderate, and strong signals, respectively. The percentage of stained cells was recorded (0% to 100%). The final IHC score was the sum of each staining intensity multiplied by the corresponding percentage, and the score scale is 0 to 300. The classification of high and low *FTO* expression was carried out according to the median score (≤ 150).

Cell Culture and Transfection

Head and neck squamous cell carcinoma cell lines, Cal27, FaDu, Tu686, HN6, and Hep-2, were purchased from the Cell Bank of the Type Culture Collection of the Chinese Academy of Science, Shanghai Institute of Cell Biology. The cell lines were maintained in Dulbecco's Modified Eagle's Medium supplemented with 10% fetal bovine serum (Gibco, Grand Island, NY, USA) and 100 U/mL penicillin-streptomycin (Gibco) in a humidified atmosphere with 5% CO₂ at 37°C. Lentivirus for shFTO-1 (TRCN0000246247), shFTO-2 (TRCN0000246248), *FTO* overexpression plasmid, double mutant (R316Q/R322Q) *FTO* plasmid as well as their controls were packaged with pMD2.G and psPAX2 (purchased from Addgene, Watertown, MA, USA). Briefly, 1.5 µg pMD2.G, 3 µg psPAX2, and 4.5 µg construct for overexpression or knockdown of specific genes were co-transfected into HEK-293T cells in 100 mm cell culture dish with Effectene Transfection Reagent (301427, QIAGEN, Valencia, CA). The lentivirus particles were harvested at 48 and 72 hours after transfection. Finally, the lentivirus particles were directly added into target cells before incubating at 37°C for 24–48 hours. Lastly, puromycin (1 µg/mL) was added to the cells for 5 days for establishing stable cell lines.

RNA Extraction, Reverse Transcription, and Amplification

Total RNA was extracted from tissues and cell lines using RNAiso Plus (code No. 9108, Takara, Shiga, Japan). The RT-PCR kit (code No. RR036A, Takara) was used to reverse transcribe total RNA into cDNA according to the manufacturer's protocol. Quantitative real-time fluorescence quantitative polymerase chain reaction (qRT-PCR) was performed using the TB Green PCR Premix Ex Taq™ II Kit (code no. RR820A, Takara) on an Applied Biosystems 7500 device (Foster City, CA, USA). All qRT-PCR tests were repeated three times. The standard for gene expression was determined based on the expression of the internal control *GAPDH*. The primer sequences were as follows: *GAPDH* forward primer, 5'-ACAACCTTTGGTATCGTGGAAGG-3'; reverse primer, 5'-GCCATCACGCCACAGTTTC-3'; *METTL3* forward primer, 5'-TTGTCTCCAACCTTCCGTAGT-3'; reverse primer, 5'-CCAGATCAGAGAGGTGGTGTAG-3'; *METTL14* forward primer, 5'-AGTGCCGACAGCATTGGTG-3'; reverse primer, 5'-GGAGCAGAGGTATCATAGGAAGC-3'; *FTO* forward primer, 5'-GCTGCTTATTTCGGGACCTG-3'; reverse primer, 5'-AGCCTGGATTACCAATGAGGA-3'; *ALKBH5* forward primer, 5'-CGGCGAAGGCTACACTTACG-3'; reverse primer, 5'-CCACCAGCTTTTGGATCACCA-3'; *CTNNB1* forward primer, 5'-CTGAGGAGCAGCTTCAGTCC-3'; reverse primer, 5'-GGCCATGTCCAACCTCATCA-3'; m⁶A PCR forward primer, 5'-CAGGGGA-GAACCCCTTGATA-3'; reverse primer, 5'-GTTCCATCATGGGGTCCATA-3'.

Western Blotting

RIPA lysis buffer (PC101, EpiZyme, Shanghai, China) was used to extract the proteins from cells. Protein concentrations were determined using a bicinchoninic acid assay reagent kit (Beyotime, Shanghai, China). Protein lysates were separated using 10% sodium dodecyl sulfate polyacrylamide gel electrophoresis and subsequently transferred onto a polyvinylidene fluoride (PVDF, WJ002, EpiZyme) membrane. The PVDF membrane was blocked with 5% skimmed milk at 25°C for 2 h followed by overnight incubation with primary antibodies (anti-FTO, 1:1000, Abcam, USA, ab92821; anti-GAPDH, 1:1000, Abcam, USA, ab8245; anti-CTNNB1 1:1000; 8480S; Cell Signaling Technology, USA, anti-slug 1:2000, Abcam, USA, ab51772; anti-ZEB1, 1:500, Abcam, USA, ab203829; anti-SOX2, 1:500, Abcam, USA, ab92494) at 4°C. The membrane was then incubated with

secondary antibodies at 25°C for 2 h and binding signals were detected using an enhanced chemiluminescent kit (Beyotime, China). Afterwards, a multi-imaging system (Tanon 5200, Tanon Science & Technology Inc., Shanghai, China) was used for image acquisition and ImageJ program was used for data quantification.

Measurement of m⁶A Levels

Total RNA from tissues and cell lines was extracted using RNAiso Plus (code no. 9108, Takara, Shiga, Japan) with deoxyribonuclease I (Sigma, Shanghai, China). The m⁶A RNA methylation quantification kit (ab185912; Abcam, USA) was used to detect m⁶A levels in the total RNA as per the manufacturer's protocol. The m⁶A level was determined by measuring the absorbance at 450 nm (Thermo Fisher, USA).

Cell Proliferation Assay

Cells were added to a 96-well plate at approximately 1500 cells per well, followed by incubation in a humidified atmosphere of 5% CO₂ at 37°C. A cell counting kit-8 (CCK-8; Obio Technology, Shanghai, China) was used for cell proliferation assays as per the manufacturer's protocol, and the absorbances at 450 nm were noted at 0, 24, 48, 72, and 96 h.

Wound Healing Assay

The digested cells were plated with appropriate confluence to ensure that the confluence was close to 100% overnight. A linear wound was scratched along the surface by a 10ul pipette tip. The serum-free medium was added for incubation in a humidified atmosphere of 5% CO₂ at 37°C after the cells were rinsed with PBS. At 0 and 24 hours after the incubation, the images were acquired with an inverted microscope after the incubation and analyzed with image J. The percentage of wound closure (%) = (width on day 0 - with on day 24)/width on day 0 × 100%.

Dot Blot Assay

The procedures were carried out according to the operation manual. The cell clone solution was fixed on the nitrocellulose membrane, and was dried and incubated with sealing solution at 25°C for 1 h. The nitrocellulose membrane was soaked in the primary antibody and incubated at room temperature for 1 h, and the nitrocellulose film was then soaked in the second antibody again after washing with TBST solution. It was then incubated at room temperature for 1 h and then washed in TBST solution again. Finally,

the nitrocellulose film was immersed in fluorescent chromogenic reagent and exposed.

Statistical Analysis

All experiments were repeated three times and GraphPad Prism (version 8; GraphPad Software, Inc., San Diego, CA, USA) was used for statistical analysis. Data are presented as the mean ± SD. The significance of differences between two groups was determined using a Student's *t*-test. One-way ANOVA was performed to compare the results of more than two groups. Data were considered statistically significant at *P* < 0.05.

Results

Prognostic Value of m⁶A

Methyltransferases and Demethylases in HNSCC

m⁶A modification is a dynamic process mediated by genes known as writers and erasers.^{32,33} To investigate the potential roles of m⁶A modulators, which might result in poor clinical outcomes in HNSCC, we first compared m⁶A modulator transcriptome expression between normal and tumor tissues in TCGA. We found that all four m⁶A modulators showed higher mRNA expression in HNSCC tissues (Figure 1A). Furthermore, we analyzed their prognostic values in TCGA cohort. Based on the Kaplan–Meier method and the Log rank test, we found that patients with high FTO expression had a trend towards poor prognosis. Although there was no significant difference in this correlation, with *P* > 0.05, the relationship between FTO and prognosis was relatively more important than that of the other three genes (Figure 1B).

We also performed an IHC staining assay to investigate FTO protein expression levels in tumor and adjacent normal tissue pairs from patients with HNSCC (Figure 1C). Consistent with the results based on the TCGA database, FTO was expressed significantly higher in HNSCC tissues compared to normal tissues, according to IHC results (Figure 1C). We further conducted survival analysis based on the expression level of FTO and the overall survival of patients; the results showed that the overall survival of patients with high FTO expression was significantly lower than that of patients with low FTO expression (Figure 1D). In addition, we found from Table 1 that there exists a correlation between smoking status and FTO expression. Among the 37 patients with a non-smoking history, the proportion of patients with low FTO expression was higher. Meanwhile, the proportion of patients

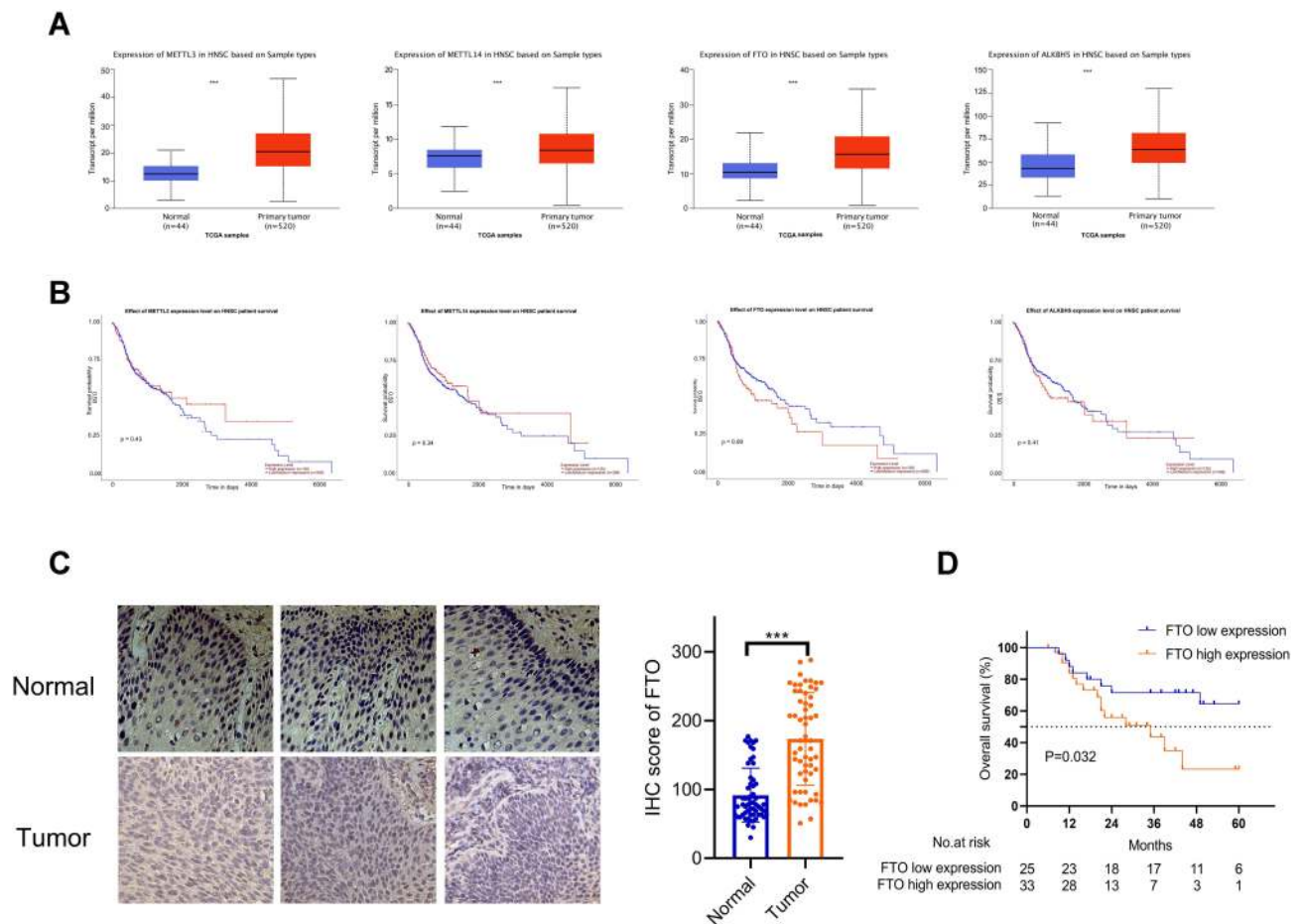


Figure 1 (A) The mRNA levels of *METTL3*, *METTL14*, *FTO*, and *ALKBH5* in the TCGA HNSCC cohort. $***P < 0.001$. (B) Kaplan–Meier analysis of the prognostic value of expression of m⁶A methyltransferase and demethylase family members in the TCGA HNSCC cohort. The survival curve for patients grouped based on their expression: *METTL3*, *METTL14*, *FTO*, and *ALKBH5*. (C) Expression of *FTO* in tumor and normal tissues in HNSCC patients, as determined by immunohistochemistry. Data are presented as the mean \pm SD, $***P < 0.001$. (D) Survival analysis of 58 patients with HNSCC showed that patients with high *FTO* expression levels had a poorer prognosis ($P < 0.05$). **Abbreviations:** TCGA, The Cancer Genome Atlas; *FTO*, fat mass- and obesity-associated protein; HNSCC, head and neck squamous cell carcinoma.

with high *FTO* expression was higher in 21 patients with a history of smoking. The relevant clinical information of patients is shown in Table 1.

FTO Acts as an Oncogene in HNSCC

To investigate the relationship between *FTO* and the phenotype of HNSCC, we first evaluated the mRNA and protein levels of *FTO* in five HNSCC cell lines, Cal27, FaDu, Tu686, HN6 and Hep-2 (Figure 2A), and chose the FaDu cell line with high *FTO* levels for further experiments. Based on loss-of-function studies in vitro, we knocked down *FTO* and constructed stable *FTO*-overexpressing FaDu cell lines (Figure 2B). The proliferation rate of FaDu cells was effectively retarded by *FTO* knock-down and accelerated by *FTO* overexpression (Figure 2C). Based on wound healing assay, *FTO* knockdown inhibited cell migration; however, *FTO* overexpression enhanced cell migration in FaDu cell line

(Figure 2D). In addition, *FTO* knockdown decreased the expression of epithelial–mesenchymal transition (EMT)-related markers and *FTO* overexpression increased EMT-related markers protein levels (Sox2, Slug, and ZEB1; Figure 2E).

CTNNB1 is a Downstream Target Gene of FTO

To identify the underlying mechanisms by which *FTO* is involved in HNSCC progression and metastasis, a previous study reported that *FTO* could demethylate *CTNNB1* transcripts and promote *CTNNB1* expression in cervical squamous cell carcinoma.³⁴ First, as we expected, *FTO* expression was significantly correlated with *CTNNB1* expression in TCGA database (Figure 3A). *FTO* knockdown increased m⁶A levels, and replenishing wild-type (WT) *FTO* but not mutant *FTO* decreased m⁶A levels in

Table 1 Summary of Patient Characteristics

		FTO Expression		P-value*
		Low Expression	High Expression	
Age	≤60 years	15 (25.9%)	17 (29.3%)	0.520
	>60 years	10 (17.2%)	16 (27.6%)	
Drinking	Never	6 (10.3%)	7 (12.1%)	0.983
	Current or former	19 (32.8%)	26 (44.8%)	
Smoking	Never	22 (37.9%)	15 (25.9%)	0.001
	Current or former	3 (5.2%)	18 (31.0%)	
Subsite	Laryngeal carcinoma	21 (36.2%)	21 (36.2%)	0.086
	Hypopharyngeal carcinoma	4 (6.9%)	12 (20.7%)	
T category	T1	5 (8.6%)	6 (10.3%)	0.816
	T2	6 (10.3%)	9 (15.5%)	
	T3	10 (17.2%)	10 (17.2%)	
	T4	4 (6.9%)	8 (13.8%)	
N category	N0	7 (12.1%)	5 (8.6%)	0.309
	N1	4 (6.9%)	10 (17.2%)	
	N2	14 (24.1%)	18 (31.0%)	
Differentiation of tumor cells	Poorly differentiated	6 (10.3%)	10 (17.2%)	0.595
	Well differentiated	19 (32.8%)	23 (39.7%)	

Note: *Significant at the 0.05 level.

Abbreviation: FTO, fat mass- and obesity-associated protein.

the FaDu cell line, based on a dot blot assay (Figure 3B). By qRT-PCR, we found that *CTNNB1* was remarkably inhibited by *FTO* deficiency in FaDu cells (Figure 3C). *CTNNB1* mutations, via constitutive activation of the Wnt signaling pathway, are involved in the development of cancer. Next, we constructed WT *FTO*- and mutant *FTO*-overexpression FaDu cells and found that WT *FTO* significantly increased *CTNNB1* expression, but not the mutant *FTO* (Figure 3D). By performing gene-specific m⁶A qRT-PCR validation, after *FTO* knockdown in HNSCC cells, m⁶A levels were significantly increased compared with those in normal control cells (Figure 3E).

CTNNB1 Reverses the Effects of FTO in HNSCC Cell Lines

We then explored whether *CTNNB1* could reverse the effects of *FTO* deficiency and overexpression on HNSCC cell proliferation and migration. We first constructed *CTNNB1*-knock-down FaDu cell line and verified that *CTNNB1* deficiency could inhibit FaDu cell growth and migration (Figure 4A and B). Then, we overexpressed *CTNNB1* in the *FTO*-knockdown FaDu cell line. As

expected, *CTNNB1* overexpression notably reversed the inhibition of HNSCC proliferation and migration resulting from *FTO* deficiency (Figure 4C and D). Together, these results showed that that *CTNNB1* is the downstream target gene of *FTO* and that *FTO* could accelerate HNSCC proliferation and migration by increasing *CTNNB1* expression.

Discussion

N6-methyladenosine RNA modification is a post-transcriptional regulation of gene expression, which has wide-ranging effects on biological processes.^{35–38} The m⁶A modification has been shown to be involved in the occurrence and progression of various cancers by regulating different downstream genes.^{16,39,40} Several studies till date have focused on the role of m⁶A regulatory genes in HNSCC,^{18–22} especially on prognosis prediction and guidance for therapeutic strategies selection and yet, little is known about the underlying mechanisms of m⁶A RNA modification in HNSCC. As RNA m⁶A eraser, FTO is involved in the demethylation of m⁶A in various cancers. For example, in ovarian cancer,

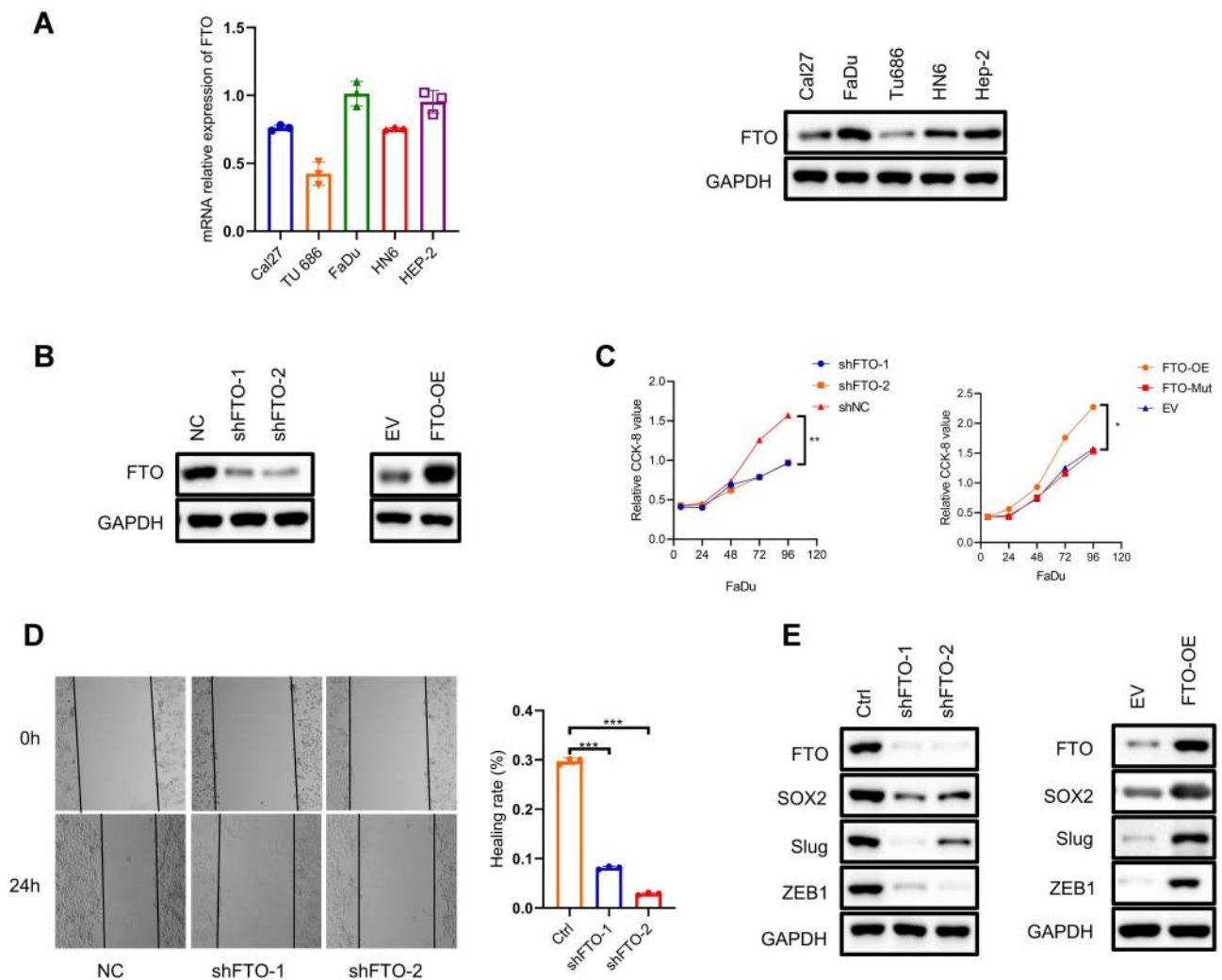


Figure 2 (A) qRT-PCR assay showing mRNA levels of *FTO* in five human HNSCC cells (Cal27, FaDu, Tu686, HN6 and Hep-2). (B) Immunoblotting to show *FTO* expression in normal control (NC) FaDu cells versus the same cells with *FTO* knockdown with shFTO-1, shFTO-2 and empty vector (EV) versus *FTO*-overexpressing (FTO-OE). Samples are normalized to GAPDH. (C) CCK-8 assay demonstrating the proliferative ability of *FTO* knockdown and replenishment wt or mutant *FTO* cell lines. * $P < 0.05$, ** $P < 0.01$. (D) Wound healing assay shows the migration ability of *FTO* knockdown cell lines. Data are presented as the mean \pm SD; *** $P < 0.001$. (E) Immunoblotting of EMT markers (SOX2, Slug, ZEB1) in FaDu cells after *FTO* knockdown and overexpression.

Abbreviations: *FTO*, fat mass- and obesity-associated protein; CCK-8, cell counting kit 8.

FTO plays a tumor suppressor role by constraining cancer stem cell self-renewal.²⁴ In hepatocellular carcinoma and intrahepatic cholangitis carcinoma, *FTO* was proven to inhibit tumorigenesis.^{25,26} However, *FTO* promotes glioblastoma tumorigenesis and cancer stem cell renewal by regulating target genes.⁴¹ *FTO*, via the m⁶A regulatory machinery, degrades the pro-apoptosis gene *BNIP3*, resulting in the promotion of breast cancer progression.⁴² In addition to these conflicting studies about the different roles in various cancers, the function of *FTO* in HNSCC was unknown to date. Our report showed that *FTO* exerts an oncogenic effect on HNSCC progression, meaning that *FTO* might be an important marker for predicting HNSCC

progression. Combined with our previous work on NSUN2,⁴³ these findings lead us to propose a mechanism through which mRNA modification affects the tumorigenesis, development, and metastasis of HNSCC. The β -catenin protein encoded by the *CTNNB1* gene is an important effector of the Wnt signaling pathway, and the pathway can regulate cell cycle, cell apoptosis, cell transformation and other processes, participating in tumorigenesis and development.⁴⁴ *CTNNB1* ectopic expression exists in liver cancer, gastric cancer, colon cancer, and breast cancer with high incidence.⁴⁵ *CTNNB1*, combining with T cell factor/lymph enhancer factor, can enter the cell nucleus and activate genes that promote cancer

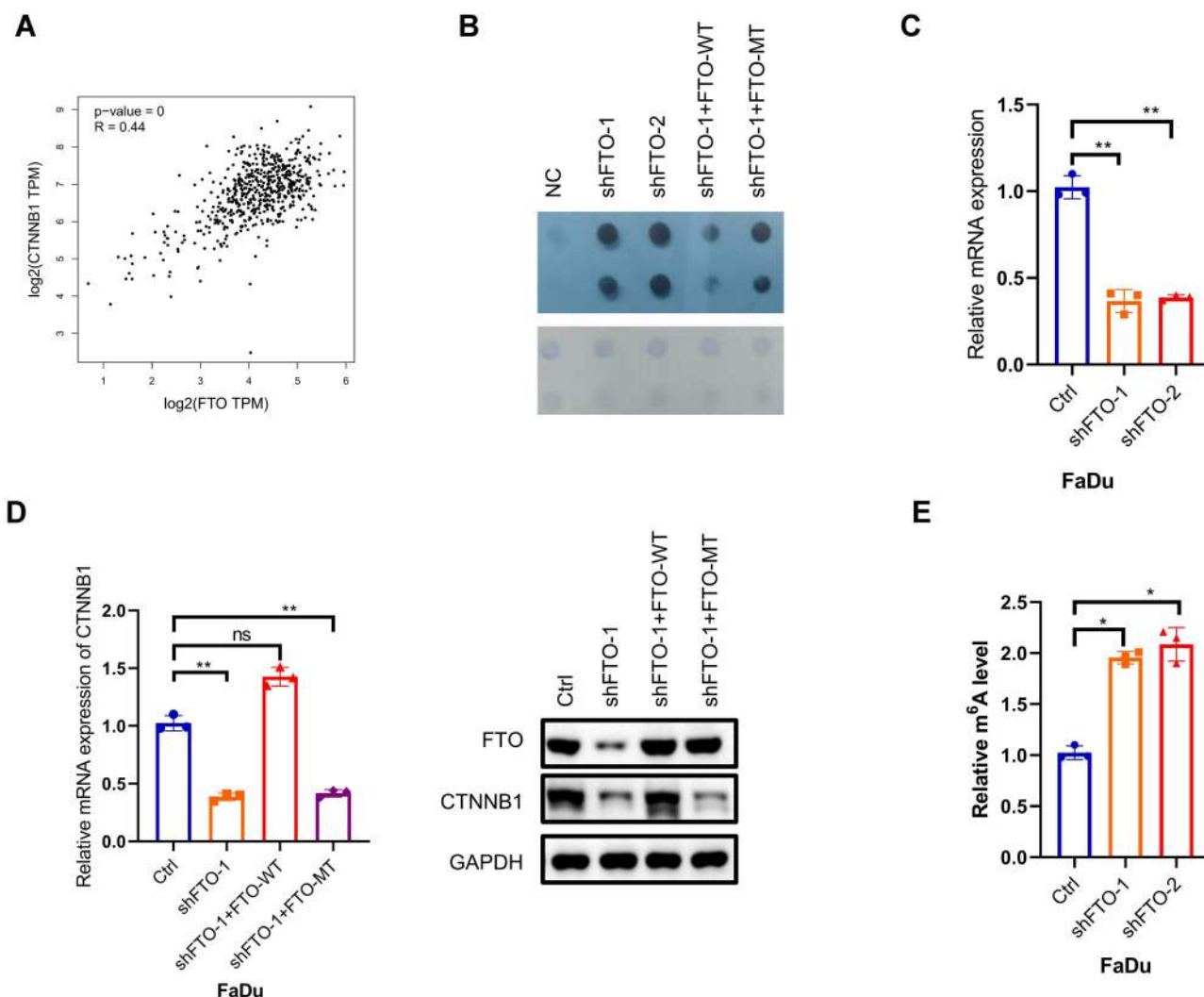


Figure 3 (A) The correlation between *FTO* and *CTNNB1* RNA expression levels in the TCGA dataset for HNSCC assessed by using the GEPIA online tool (<http://gepia.cancer-pku.cn/>). (B) m⁶A dot blot assay of knockdown and replenishment of wild-type (WT) and mutant (MT) *FTO* in FaDu cells. Methylene blue staining was used as the control. (C) RT-qPCR arrays were used to detect the RNA expression levels of *CTNNB1* after *FTO* knockdown in FaDu cell, normalized to GAPDH. Data are presented as the mean \pm SD; ** $P < 0.01$. (D) *CTNNB1* RNA expression levels were analyzed in *FTO*-WT and *FTO*-mutant FaDu cells, normalized to GAPDH. Immunoblot assay of *CTNNB1* protein levels in FaDu cells upon *FTO* knockdown and *FTO*-WT and *FTO*-mutant. Data are presented as the mean \pm SD, *** $P < 0.01$. (E) m⁶A RIP and qRT-PCR analysis of m⁶A levels in *CTNNB1* mRNA in sh*FTO*-1/2 or corresponding wild-type cells. Data are expressed as the mean \pm SD; * $P < 0.05$.

Abbreviations: ns, not significant; *FTO*, fat mass- and obesity-associated protein; *CTNNB1*, Catenin beta 1.

proliferation, thereby increasing the malignant behavior of cervical cancer cells.⁴⁶ Kurnit et al show that *CTNNB1* gene mutations can predict the risk of endometrial cancer recurrence, suggesting that *CTNNB1* may be related to tumor recurrence.⁴⁷

In our study, we found that *FTO* was highly expressed in HNSCC compared to that in normal tissues and that patients with high *FTO* expression had a trend toward poor prognosis, indicating the role of *FTO* in the regulation of HNSCC development. Regarding the relationship between smoking status and *FTO* expression, we found that among patients with a history of smoking, there were more patients with high *FTO* expression. Smoking has been

considered an unfavorable prognostic factor for HNSCC.^{48–50} Therefore, high *FTO* expression in patients with a history of smoking was compatible with our research view that patients with high *FTO* expression had a trend towards poor prognosis. A recent study on lung cancer mentioned that some m⁶A regulatory factors, including *FTO*, were related to smoking status.⁵¹ In later studies, we will continue to focus on the potential relationship between smoking status and *FTO* in HNSCC. We also designed serial biological functional assays using the FaDu cell line to demonstrate that *FTO* plays a pivotal role in promoting cell proliferation and metastasis in vitro. To explore the molecular mechanism by which *FTO*

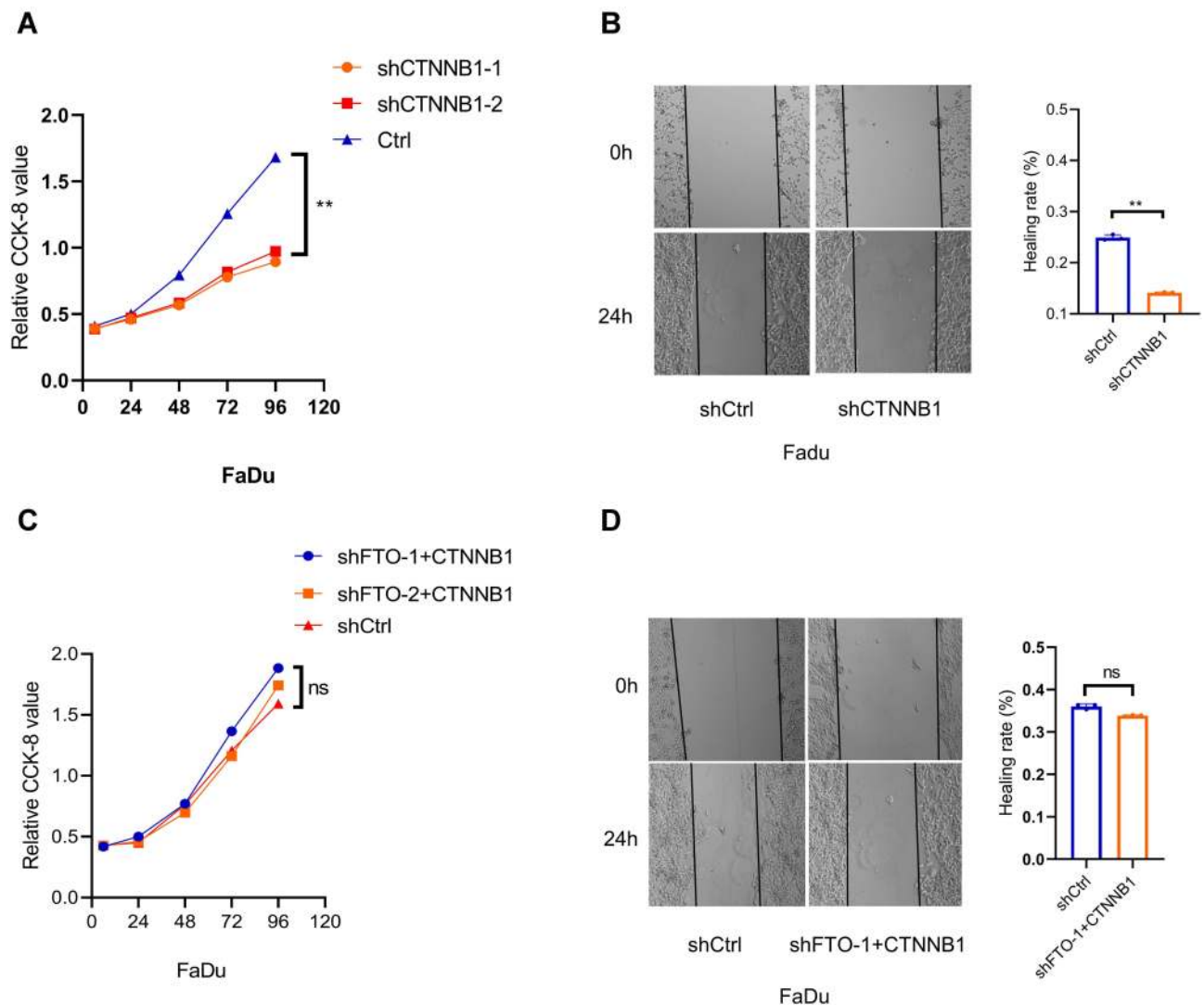


Figure 4 (A) CCK-8 assay demonstrating the proliferation ability of CTNNB1 knockdown cell lines. $**P < 0.01$. (B) Wound healing assay showing the migration ability of CTNNB1 knockdown cell lines. Data are presented as the mean \pm SD; $**P < 0.01$. (C) CCK-8 assay showing that the proliferation of control (Ctrl) and shFTO cell lines did not obviously differ after CTNNB1 overexpression in the shFTO cell lines. ns: no significance. (D) Wound healing assay showing that the migration of control (Ctrl) and shFTO cell lines was not significantly different after CTNNB1 overexpression in the shFTO cell lines. Data are presented as the mean \pm SD.

Abbreviations: ns, not significant; *FTO*, fat mass- and obesity-associated protein; *CTNNB1*, Catenin beta 1.

promotes tumor growth in HNSCC, we analyzed TCGA database for HNSCC. We found that the expression level of *FTO* was significantly correlated with one oncogenic transcription factor, *CTNNB1*. We then performed qRT-PCR and Western blot experiments, whose results suggest that *FTO* knockdown in the FaDu cell line would lead to a decrease in the mRNA and protein level of *CTNNB1*. We also investigated wild-type *FTO* and a control catalytically inactive mutant using an *FTO*-knockdown FaDu cell line, as control. We found that WT *FTO*, but not the inactive mutant, could stabilize mRNA and protein levels. These results indicated that *FTO* stabilizes the mRNA level in

FaDu cells to promote tumorigenesis and the development and metastasis of HNSCC.

There are some limitations in this study, such as the lack of in vivo experiments on *FTO* and *CTNNB1*, and the lack of m⁶A methylation sequencing to clarify *CTNNB1* as the target gene of *FTO*. In summary, we provide here the first report that *FTO*, the key m⁶A demethylase, is up-regulated in HNSCC patients and that patients with high expression had a trend towards poor prognosis. *FTO* was found to promote HNSCC cell line proliferation and migration by stabilizing mRNA encoding *CTNNB1*, which has oncogenic roles in various cancers. Altogether,

our findings suggest that *FTO* might serve as a novel potential therapeutic target for HNSCC.

Conclusion

Our studies demonstrated a novel mechanism that *FTO* promotes head and neck squamous cell carcinoma proliferation and migration by increasing *CTNNB1* in an m6A-dependent manner.

Abbreviations

Ctrl, control; HNSCC, head and neck squamous cell carcinoma; IHC, immunohistochemistry; m6A, N6-methyladenosine; *FTO*, fat mass and obesity-associated protein; *CTNNB1*, catenin beta 1; qRT-PCR, quantitative real-time fluorescence quantitative PCR; TBST, 0.1% Tris-HCl with Tween-20; TCGA, The Cancer Genome Atlas; *METTL3*, methyltransferase -like 3; *METTL14*, methyltransferase-like 14; *ALKBH5*, alkB homologue 5; EMT, epithelial-mesenchymal transition; *WTAP*, Wilms tumour 1-associated protein, *IGF2BP*, IGF2 mRNA binding proteins; m6A, N6-methyladenosine; *RBM15/15B*, RNA binding motif protein 15/15B, *YTHDF*, YTH521-B homology domain-containing protein family.

Data Sharing Statement

The original data of this article can be obtained from the corresponding author on reasonable request.

Author Contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

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Disclosure

Xiaoliang Wu is an employee of Tailai Bioscience, Shenzhen, China. The authors declare that they have no other conflict of interest.

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