

LncRNA *NEAT1*/*miR-204*/*NUAK1* Axis is a Potential Therapeutic Target for Non-Small Cell Lung Cancer

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Background: Long non-coding RNA (lncRNA) is a key part of non-coding RNA, and more and more evidence has revealed that it plays a vital role in tumors. *NEAT1* is a lncRNA discovered in the early stage. However, it is still unclear whether *NEAT1* and *miR-204* play a regulatory role in lung cancer (LC). This research aimed to determine the biological function of *NEAT1*/*miR-204* in non-small cell lung cancer (NSCLC).

Materials and Methods: In order to research the function of *NEAT1* in NSCLC, RT-PCR, Western blot, luciferase assay and RNA immunoprecipitation assay were used to determine the relationship between *NEAT1*, *miR-204* and *NUAK1*. CCK8 test, cell migration and invasion test were used to explore the influence of *NEAT1* on proliferation and metastasis of LC cells. Tumor allotransplantation was used to detect the influence of *NEAT1* on the growth of LC.

Results: The results revealed that *NEAT1* was obviously enhanced in LC cell lines. Further functional analysis showed that low expression of *NEAT1* obviously suppressed the growth, migration and invasion of NSCLC and facilitated cell apoptosis. Determination of luciferase reporter gene revealed that *miR-204* was the direct target of *NEAT1* in LC. In addition, *NUAK1* was called the direct target of *miR-204*, and *miR-204*/*NUAK1* had saved the role of *NEAT1* in NSCLC cells. Tumor allotransplantation experiments showed that knocking down *NEAT1* could inhibit the growth of LC.

Conclusion: In summary, our results showed that the down-regulation of *NEAT1* in NSCLC inhibited its growth, migration and invasion through the *miR-204*/*NUAK1* axis.

Keywords: *NEAT1*, *miR-204*, *NUAK1*, non-small cell lung cancer

Introduction

Lung cancer (LC), as the most common malignant respiratory tumor clinically, is still the main reason of tumor-related morbidity and mortality all over the world.¹ According to the global statistics in 2018,² there are more than 2,000,000 new patients with LC and 1,700,000 dead patients in 2018. It also indicates that the new patients with LC are gradually getting younger. According to the type of LC, it can be separated into small cell lung cancer (SCLC) and non-small cell lung cancer (NSCLC), in which NSCLC accounts for more than 85%.³ At present, the treatment of NSCLC is mainly through surgery combined with radiotherapy and chemotherapy. However, due to the relatively hidden incidence of NSCLC, most of the patients are in the middle and advanced stages when they are admitted to the hospital, while the prognosis of patients with advanced NSCLC is not ideal.

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Data show that the 5-year survival rate of patients with late NSCLC is less than 15%.^{4,5} In recent years, targeted therapy has played a vital role in improving the prognosis of patients.⁶ However, there are few therapeutic targets for NSCLC at present. It is the key to seek potential targets and further explore the molecular mechanism of NSCLC for improving the present situation.

At present, only 1% to 2% of the genome and transcriptome data studied by humans are genome encoded proteins, while most mammalian genomes contain this large amount of non-encoded RNA.⁷ In recent years, long-chain non-coding RNA (lncRNA) has become a research hotspot in various fields.⁸ lncRNA is a non-coding RNA with a length of more than 200 nt. Since lncRNA were initially found to have no protein-coding capacity,⁹ they were considered as metabolic wastes during transcription. However, recent studies have revealed that lncRNA has great differences in various diseases.^{10,11} Especially in tumors, researches^{12,13} have shown that part of lncRNA affect the development and progression of tumors. Nuclear para specific assembly transcript 1 (*NEAT1*) (also known as LINC00084), as a member of lncRNA family, is located on human 11q13.1 chromosome.^{14,15} Previous studies have revealed that *NEAT1* is differentially expressed in liver carcinoma,¹⁶ gastric carcinoma¹⁷ and breast carcinoma.¹⁸ Many other studies have revealed that^{19,20} *NEAT1* participates in the development of LC by regulating downstream microRNA (miR), but there is no report that *NEAT1* can participate in the development of LC by regulating *miR-204*. In this study, we found a targeted binding site between *NEAT1* and *miR-204* through the joint prediction of LncBase,²¹ starbase,²² mircode²³ and miRDB.²⁴

Therefore, this study aimed to explore whether *NEAT1* could regulate *miR-204* and its downstream target genes to participate in the development of LC and provide potential targets for clinical treatment.

Methods and Materials

Online Website Analysis

The expression of *miR-204* in lung adenocarcinoma (LA) was analyzed by starbase website (<http://starbase.sysu.edu.cn/>), and the expression of *NEAT1* in LA was analyzed by GEPIA2²⁵ (<http://gepia2.cancer-pku.cn/#index>), and pictures were obtained to show the expression.

Clinical Data

A total of 68 patients with NSCLC treated in People's Hospital of Gaochun from January 2015 to January 2017 were selected. The tumor and adjacent tissues of the patients were collected during surgery. The collected tissues were transported by liquid nitrogen and sent to the laboratory for testing. The samples collected from patients in this study had not been treated with chemoradiotherapy or anti-tumor therapy before this study. All samples of the patients were sent to the pathology department for examination and patients were confirmed as LA. All patients affixed informed consent forms. This research was ratified by the Medical Ethics Committee of People's Hospital of Gaochun and it was consistent with the Helsinki Declaration.²⁶

Cell Culturing and Transfection

Human bronchial epithelial cell strain 16HBE, as well as LC cell strains A549, SPC-A1, NCI-H23, NCI-H520 were acquired from the American Center for Typical Culture Collection (Manassas, Virginia, the United States). The purchased cells were cultivated using Dulbecco modified Eagle Medium (DMEM), which supplemented 10% of fetal bovine serum (Gibco, Grand Island, New York, the United States), penicillin (100 U/mL) and streptomycin (100 mg/mL) in a humid environment comprising 5% CO₂ at 37 °C. In this study, Lipofectamine 3000 (Invitrogen, CA, USA) was used for transfection. *miR-204* mimetic (*miR-204-mimics*) /inhibitor (*miR-204-inhibit*), siRNA targeting *NEAT1* (si-*NEAT1*#1:5'-GGTCTGTGTGGAAGGAGGAAGGCAG-3', si-*NEAT1*#2:5'-GCCAUCAGCUUUGAAUAAU-3', si-*NEAT1*#3:5'-GGUGUUAUCAAGUGAAUUAU-3'), lentivirus and corresponding negative control were all from GenePharma (Shanghai, China).

Detection of Cell Proliferation

In this study, CCK-8 kit was used to test cell proliferation. The transfected cells (2.5 × 10³) were inoculated into 96-well plate (Corning Inc, United States). According to the manufacturer's regulations, cell growth was evaluated at various time points (0, 24, 48, 72 and 96 h) through CCK-8 assay (Dojindo, Tokyo, Japan). The absorbance value at 450 nm wavelength was measured using an enzyme-labeling instrument (Molecular Devices, USA), which was positively correlated with cell proliferation.

Cell Invasion and Migration

In this study, Transwell was used to measure cell migration and invasion. For invasive analysis, the Transwell chamber (8 μ m, Corning, Inc., Corning, NY, the United States) was precoated with Matrigel (BD Biosciences, Franklin Lake, NJ, the United States). Transfected cells were resuspended in serum-free medium and inoculated into the upper Transwell chamber. The DMEM (500 μ L) comprising 10% fetal bovine serum was put in the lower chamber as chemotactic inducer. After culturing at 37 °C for 24 h, the cells remaining in the upper chamber were eliminated with cotton swabs, and the cells invading the lower chamber were fastened in 100% methanol and dyed with 0.05% crystal violet (Sigma-Aldrich). The amount of invasive cells was counted from 5 randomly selected areas and imaged using an inverted microscope (Olympus, Tokyo, Japan). The study of cell migration was conducted in a cross-well chamber without matrix and followed similar instructions.

Apoptosis Detection

Flow cytometry was used to test apoptosis in this study. Transfected cells were obtained by trypsin transfection. After double staining with FITC-Annexin V and PI, FITC-Annexin V apoptosis test kit (BD Biosciences) was used on the basis of kit instructions. Flow cytometer (FACScan, BD biosciences) of Cell Quest 3.0 software was used to analyze cells.

qRT-PCR

In this research, qRT-PCR was used to detect the relative expression of lncRNA/miR/mRNA. The cells were obtained and total RNA was abstracted from the tissue through TRIzol reagent (Invitrogen). The extracted total RNA was reverse transcribed into cDNA through RNA reverse transcription kit (Takara). Real-time PCR analysis was conducted with SYBR Green (Takara), the amplification was performed with ABI7500, and the operation was conducted on the basis of the manufacturer's specifications. GAPDH (for lncRNAs) or U6 (for miRNAs) were used as internal parameters. The multiple change of relative expression was calculated by $2^{-\Delta\Delta CT}$.²⁷ The primer sequences are shown in Table 1.

WB Detection

Radioimmunoprecipitation assay (RIPA) of protease inhibitor was used to break cells with lysis buffer. After high-speed centrifugation, the supernatant was obtained and heated in a water bath to denature the protein. The protein was quantified by double caprylic acid (BCA). Then, the protein was separated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and sealed with 5% skim milk and 0.05%TBS-Tween-20 (TBST) for 1 h at room temperature, and the first antibodies (cleave-caspase 9, Bax, Bcl-2, MMP-2, MMP-9) were added and cultured at 4 °C for 12 h. The polyvinylidene fluoride (PVDF) film was washed with TBST solution, and then cultured with a secondary antibody (Hubeibiosci Biotechnology Co., Ltd., 1: 2000) at ambient temperature for 1 hour. After that, the chemiluminescence method was developed by hypersensitivity ECL (Hubei Biotechnology Co., Ltd.). β -actin was used as an internal reference.

Double Fluorescein Report

The *miR-204* mutation site in *NEAT1* luciferase reporter vector was produced through site-directed mutagenesis. PCR products were cloned into PMIR-GLO luciferase vector downstream of the firefly luciferase coding region (Invitrogen, USA). The construct was called pMIR-*NEAT1*-wt. The mutation of *miR-204* binding site was introduced through site-directed mutagenesis, and the obtained plasmid was called pMIR-*NEAT1*-Mut. Luciferase assay was performed with 1×10^4 , and HEK-293 cells were inoculated in a 48-hole plate. The co-transfection was conducted through 200 ng pMIR-*NEAT1*-wt or pMIR-*NEAT1*-Mut vector and 80 ng *miR-204* mimetic or inhibitor. After transfection for 48 hours, the cells were obtained, and the contents of fireflies and algal luciferases were determined using Dual-Glo luciferase assay kit (Promega, Madison, Wisconsin, United States). All transfection tests were repeated three times. *miR-204* and NUA family kinase 1 (*NUAK1*) were carried out according to the above steps.

Table 1 Primer Sequences

Primer Sequences	Upstream Primers	Downstream Primers
<i>NEAT1</i>	5'-GGATATGCCCTTCATCCTGTGACGC-3'	5'-TCCACACGTCCACTGTCCCAAAATC-3'
<i>miR-204</i>	5'-CCCATCGTTAAGCAATGCATGAC-3'	5'-GAGGGCCTCCTGATCATTTACC-3'
<i>NUAK1</i>	5'-CCGCTCACTGATGTAATCGT-3'	5'-GTCATCTCTCAACCATCCTCAT-3'
GAPDH	5'-ATTCCACCCATGGCAAATTC-3'	5'-TGGGATTTCCATTGATGACAAG-3'
U6	5'-CTCGCTTCGGCAGCAC-3'	5'-AACGCTTCACGAATTTGCGT-3'

RIP Experiment

In this study, EZ-Magna RIP RNA binding protein immunoprecipitation kit (Millipore, Bedford, MA, USA) was applied for RNA immunoprecipitation (RIP) analysis. The specific steps were carried out on the basis of the manufacturer's kit. The method was as below: A549 cells with fusion degree of 80–90% were assimilated, obtained and lysed in RIP lysis buffer. Next, 100 μ L of whole cell lysate was cultivated with RIP buffer comprising magnetic beads coupled to human antibody against Argonaute 2 (Ago2) (ABCam, Cambridge, MA, USA) or normal mouse IgG (Millipore) for 6 hours at 4 °C. The beads were processed with protease K buffer to digest the protein, and then the target RNA was abstracted. Finally, the coprecipitated RNA was analyzed by qRT-PCR for further study.

Statistical Analysis

In this study, GraphPad 7 was used to draw the required pictures and data analysis. SPSS version 20.0 was used to analyze the independent prognostic factors of patients.

The measurement data was represented by mean number \pm standard deviation (Mean \pm SD). The comparison between groups was conducted through independent sample *t* test. The counting data were represented by percentage (%). The chi-square test was expressed by 2. One-way ANOVA was applied for comparison among multiple groups, expressed as F. LSD-*t* test was applied for pairwise comparison afterwards. Repetitive measurement and analysis of variance was applied for expression at multiple time points, expressed as F. Bonferroni was used for post test. Pearson test was used to analyze the correlation of each gene. There was a statistical difference with $P < 0.05$.

Results

Expression of *NEAT1* Increased in LA

First of all, we analyzed it to determine the *NEAT1* expression in LA through GEPIA2 online website. The results showed that the expression of *NEAT1* in LA increased in the TCGA database (Figure 1A). Then, qRT-PCR was used to detect the expression in LA

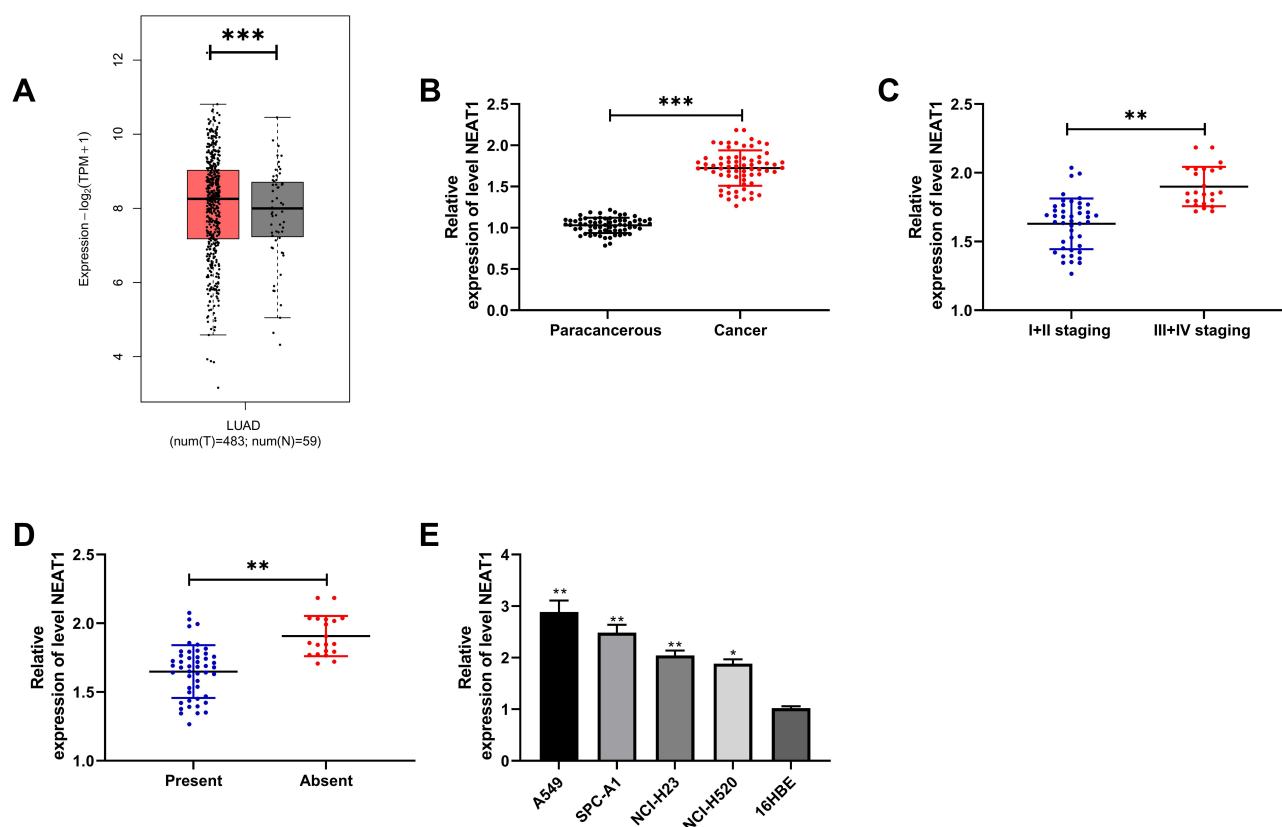


Figure 1 Expression of *NEAT1* in LA. (A) GEPIA2 was used to analyze the *NEAT1* relative expression in TCGA database. (B) qRT-PCR was used to detect the *NEAT1* relative expression in tumor tissues of LA patients. (C) qRT-PCR was used to detect the relationship between *NEAT1* relative expression and TNM staging in tumor tissues of LA patients. (D) qRT-PCR was used to detect the relationship between *NEAT1* relative expression and lymph node metastasis in tumor tissues of LA patients. (E) qRT-PCR was used to detect the *NEAT1* relative expression in LA cell line. * $P < 0.05$; ** $P < 0.01$; *** $P < 0.001$.

Table 2 Relationship Between *NEAT1* and Pathological Data of LA Patients

Factors		NEAT1		P
		High Expression (n=34)	Low Expression (n=34)	
Age	≥55 years old (n=42) <55 years old (n=26)	20 14	22 12	0.618
Gender	Male (n=28) Female (n=40)	15 19	13 21	0.622
Smoking history	Yes (n=38) No (n=30)	22 12	16 18	0.143
TNM stages	I+II (n=44) III+IV (n=24)	18 16	26 8	0.042
Lymph node metastasis	Present (n=20) Absent (n=48)	14 20	6 28	0.033
Differentiation	Poorly differentiated (n=19) Middle+well differentiated (n=49)	12 22	7 27	0.177

patients' tissues and it revealed that the relative expression was obviously enhanced compared with the paracarcinoma tissues (Figure 1B). Patients were further separated into high and low expression groups in accordance with the median expression value of *NEAT1*. By observing the connection of *NEAT1* with the pathological data of patients, it was found that there was lymph node metastasis and TNM high stage probability increased in patients with high expression of *NEAT1* (Table 2). Further analysis uncovered that *NEAT1* in patients with lymph node metastasis (Figure 1C) and high TNM stage (Figure 1D) was higher than in patients without lymph node metastasis and low TNM stage. In addition, we found that the *NEAT1* in LC cell line was obviously higher than that in human bronchial epithelial cell line (Figure 1E) through qRT-PCR detection, suggesting that *NEAT1* was highly expressed in LA.

Knocking Down *NEAT1* Could Inhibit the Growth, Invasion and Migration of LA Cells and Induce Cell Apoptosis

To further determine the influence of *NEAT1* on LA, we constructed si-*NEAT1* vector (Figure 2A), and

selected si-*NEAT1*#2 with significant difference for cell transfection (Figure 2B). Subsequently, CCK-8 assay revealed that the proliferation activity of cells transfected with si-*NEAT1*#2 was significantly inhibited compared with that of si-NC (Figure 2C-D), while Transwell experiment revealed that the invasion and migration activity of cells transfected with si-*NEAT1*#2 were significantly decreased compared with that of si-NC (Figure 2E-F). In addition, flow cytometry showed that apoptosis was significantly induced after transfection of si-*NEAT1*#2 (Figure 2G). These results implied that *NEAT1* might be a latent target for the therapy of LA.

NEAT1 Acted as an *miR-204* Sponge

In order to further determine the mechanism of *NEAT1* in LA, we predicted that *NEAT1* could bind miR. Previous studies showed that lncRNA could regulate miR to participate in tumor development. Through the online prediction website, we predicted and analyzed the downstream of *NEAT1* in combination with miR. It was found that *NEAT1* and *miR-204* had a targeted binding relationship (Figure 3A-B). Subsequently, we concluded that *miR-204* was weakly expressed in LA through starbase analysis of

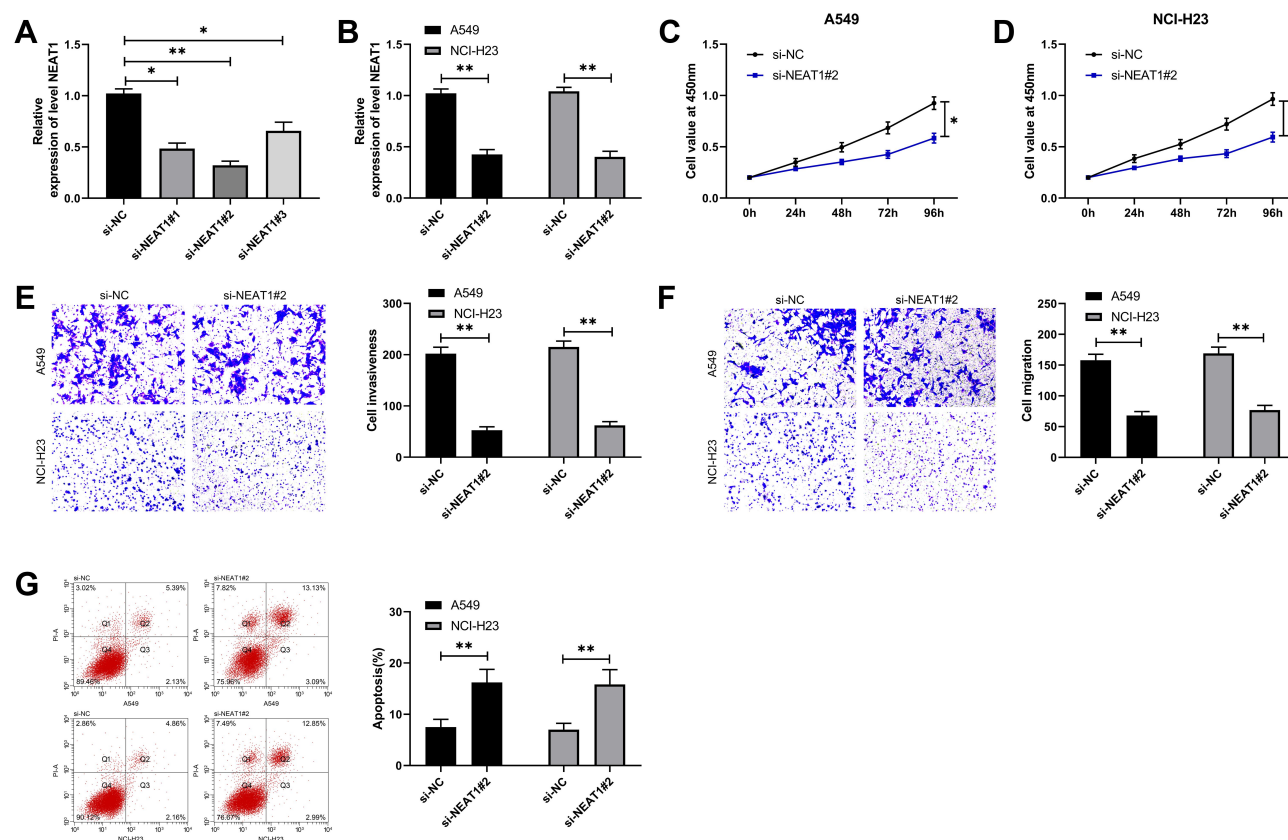


Figure 2 Knocking down NEAT1 inhibited the proliferation, invasion, migration and induced apoptosis of LC cells. (A) qRT-PCR was used to detect the relative expression of NEAT1 in si-NEAT1 vector. (B) qRT-PCR was used to detect the relative expression of NEAT1 after si-NEAT1#2 transfection into LC cells. (C-D) CCK-8 test was used to detect the proliferation of cells transfected with si-NEAT1#2. (E-F) Transwell test was used to detect the change of cell invasion and migration ability after transfection of si-NEAT1#2. (G) Flow cytometry was used to detect the change of apoptosis rate after transfection of si-NEAT1#2. * $P < 0.05$, ** $P < 0.01$.

miR-204 expression in LA (Figure 3C). In addition, qRT-PCR detection revealed that *miR-204* in cancer tissue of LA patients was lower than that in para-carcinoma tissues (Figure 3D). The correlation analysis also revealed that the *NEAT1* was negatively correlated with *miR-204* in LA tissues (Figure 3E). This indicated that there was a latent targeting connection between *NEAT1* and *miR-204*. Then, we found that *miR-204*-mimics could inhibit *NEAT1*-WT fluorescence activity (Figure 3F) through double luciferase report and RIP experiment analysis respectively, while RIP experiment showed that *NEAT1* and *miR-204* could be precipitated by Ago2 antibody (Figure 3G). It confirmed that *NEAT1* could target *miR-204*.

miR-204 Targeted Regulation of *NUAK1*

It has been proved that miR regulates downstream target genes to participate in the occurrence of diseases. In order to further study the potential mechanism of *miR-204*, we predicted the target gene downstream of *miR-204*. Through prediction, we found that there was a target

site between *miR-204* and *NUAK1* (Figure 4A). Subsequently, we detected the *NUAK1* in LA. qRT-PCR showed that the *NUAK1* was increased in LA (Figure 4B). The correlation analysis also showed that *miR-204* was negatively correlated with *NUAK1* and positively correlated with *NEAT1* in tumor tissue of patients with LA (Figure 4C). In addition, the double luciferase report experiment showed that the fluorescence activity of *NUAK1*-WT could be suppressed by *miR-204*-mimics (Figure 4D), and the relative expression of *NUAK1* in the cells after transfection with *miR-204*-mimics was also determined through qRT-PCR and WB experiments (Figure 4E-F). This indicated that *miR-204* targeted regulation of *NUAK1*.

NEAT1 Inhibited the Growth and Metastasis of LC Cells by Regulating *miR-204/NUAK1* Axis

At the end of the study, we conducted a rescue experiment to further determine whether *NEAT1* could participate in the

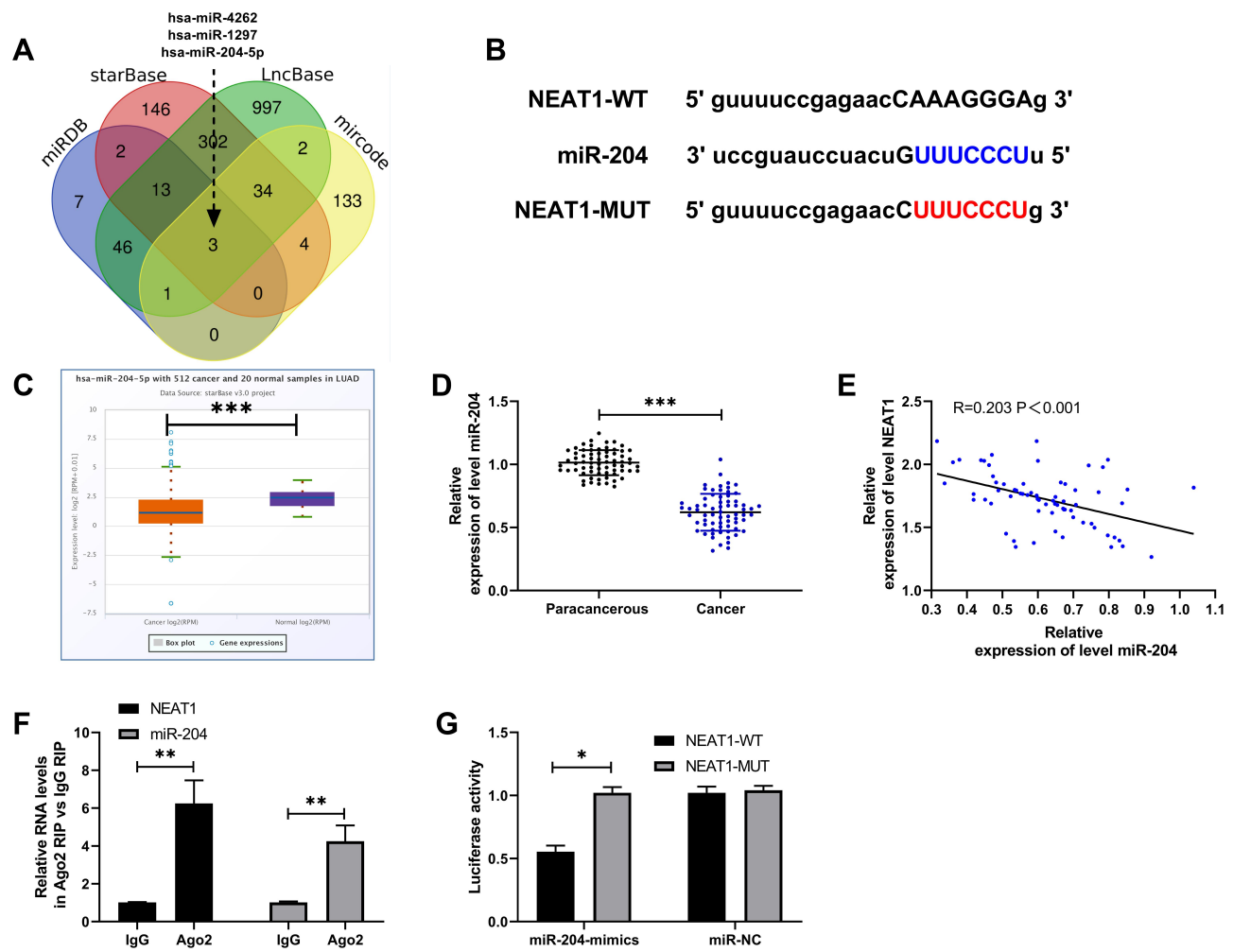


Figure 3 NEAT1 regulated miR-204. (A) miRDB, starBase, LncBase and miRcode online prediction websites were used to predict NEAT1 potential miR. (B) Targeted binding site of NEAT1 and miR-204. (C) starBase was used to analyze the relative expression of miR-204 in LA in TCGA database. (D) qRT-PCR was used to detect the relative expression of miR-204 in tumor tissues of LA patients. (E) Pearson test was used to analyze the correlation between NEAT1 and miR-204 in tumor tissues of LA patients. (F) RIP experiment confirmed that NEAT1 and miR-204 could be precipitated by Ago2 antibody. (G) Double luciferase report was used to detect the effect of miR-204-mimics on NEAT1-WT fluorescence activity. *P<0.05, **P<0.01, ***P<0.001.

development of LC by adjusting the *miR-204/NUAK1* axis. Through the experiment, we concluded that the cell growth, invasion and migration abilities were significantly faster than those of pcDNA-NC+miR-inhibit, and the cell apoptosis was inhibited since *miR-204*-inhibit and pcDNA-*NUAK1* were transfected into LA cells. However, the above results were reversed since si-*NEAT1*#2 was co-transfected with *miR-204*-inhibit or pcDNA-*NUAK1* (Figure 5A-C, 6A). In addition, WB detection showed that the relative expression of cleaved-caspase 3, cleaved-caspase 9, Bax, MMP-2, MMP-9 protein in cells transfected with *miR-204*-inhibit and pcDNA-*NUAK1* was obviously decreased, while the relative expression of Bcl-2 protein was obviously increased. However, the above results were reversed after co-transfection (Figure 6B), suggesting that

NEAT1 inhibited the growth and metastasis of LC cells by regulating *miR-204/NUAK1* axis.

Discussion

In this research, we concluded that *NEAT1* expression increased in patients with LA and the cell strains, and the probability of TNM high staging and lymphatic metastasis increased significantly in patients with high expression of *NEAT1*. In addition, we found that knock-down of *NEAT1* could inhibit the growth, invasion and migration of LA cells, and induce apoptosis of LA cells, so it was a potential therapy target for LA. Besides, we found for the first time that *NEAT1* could suppress the growth and metastasis of LC through *miR-204/NUAK1* axis, which was expected to become a clinical therapeutic target.

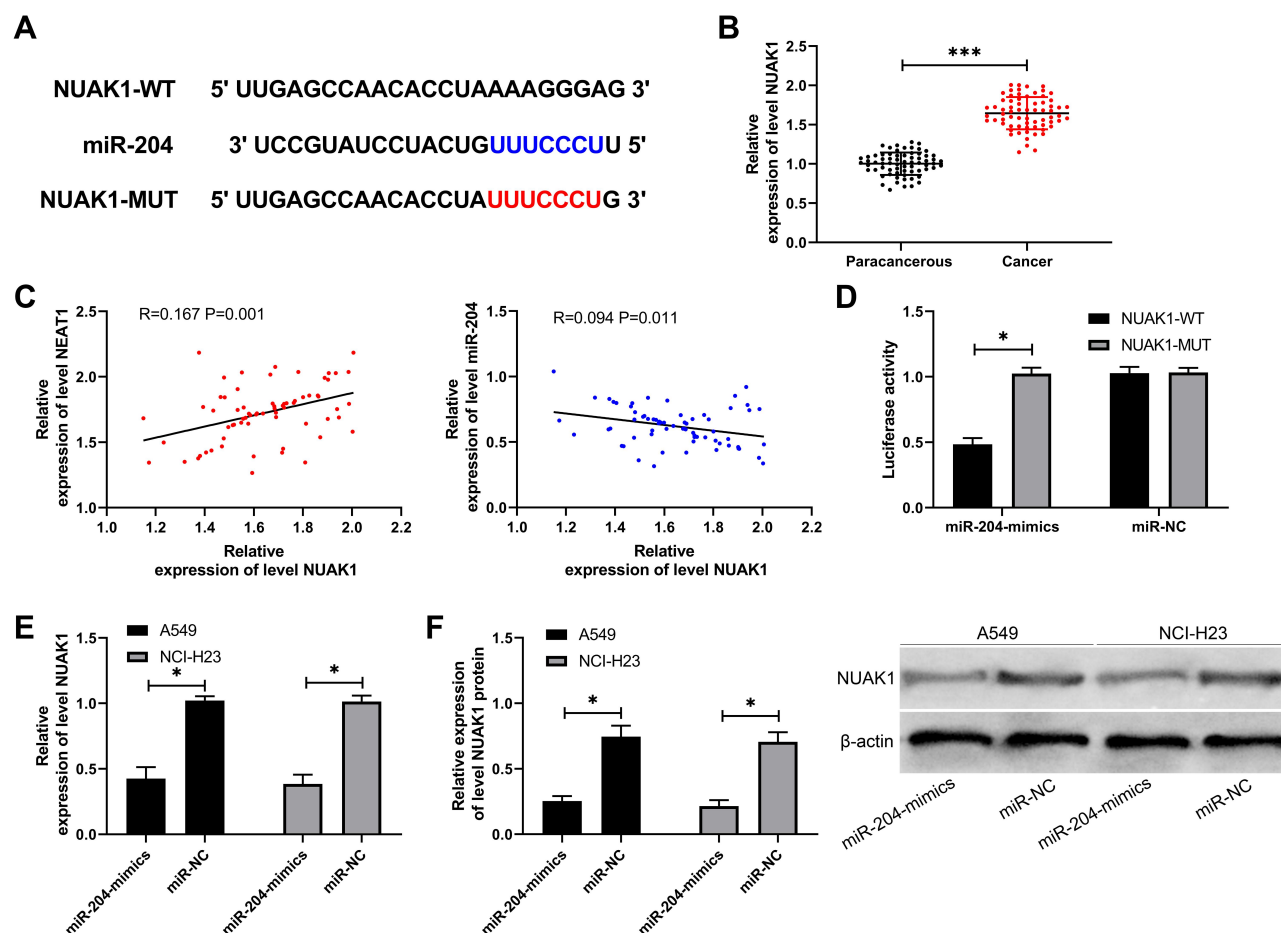


Figure 4 NUA1 was a downstream target gene of miR-204. **(A)** Targeted binding site of miR-204 and NUA1. **(B)** qRT-PCR was used to detect the relative expression of NUA1 in tumor tissue of LA patients. **(C)** Pearson test was used to analyze the correlation between NUA1 and miR-204/NEAT1. **(D)** Double luciferase report confirmed that the fluorescence activity of NUA1-WT was inhibited by miR-204-mimics. **(E-F)** qRT-PCR and WB experiments were used to detect the relative expression of NUA1 in cells transfected with miR-204-mimics. * $P < 0.05$, *** $P < 0.001$.

NEAT1, as an early lncRNA, has been found to affect the progression of many diseases. For instance, studies by Pang et al²⁸ have revealed that *NEAT1* could promote breast cancer progression through miR-124/STAT3 axis. In addition, the researches of Zhang et al²⁹ have shown that *NEAT1* can promote the progress of colorectal cancer by activating Wnt/ β -catenin signal and interacting with DDX5. In this research, we first studied the *NEAT1* in LA through GEPIA2 web analytics, and found that the *NEAT1* in LA was higher than that in control samples. Then, we found that the expression of *NEAT1* in tumor tissues of patients with LA and LA cell strains was higher than that in adjacent tissues and human bronchial epithelial cell lines through qRT-PCR detection, which was consistent with the previous research results, indicating that *NEAT1* was expected to become a potential diagnostic indicator of LA. At present, the strategy for clinical treatment of tumors is to achieve therapeutic effect by

suppressing the growth of tumor cells and facilitating tumor cell apoptosis.³⁰ Therefore, we transfected the knocked-down *NEAT1* vector into LA cells. Through observation, we found that the growth, invasion and migration of cells transfected with si-*NEAT1*#2 were significantly weakened compared with the control, and the apoptosis was further promoted. However, studies of Li et al³¹ have also shown that knocking down *NEAT1* can suppress the proliferation and invasion of LC cells, which verified our study.

Competing endogenous RNAs (ceRNA) are considered as a new mechanism of RNA interaction.^{32,33} CeRNA can regulate gene expression by competitively binding microRNA.³⁴ For example, researches by Yuan et al³⁵ have shown that *NEAT1*/miR-133a axis participates in promoting the progression of cervical carcinoma by targeting SOX4. Other studies have revealed that³⁶ *NEAT1*/miR-1224/KLF3 participates in proliferation, apoptosis and

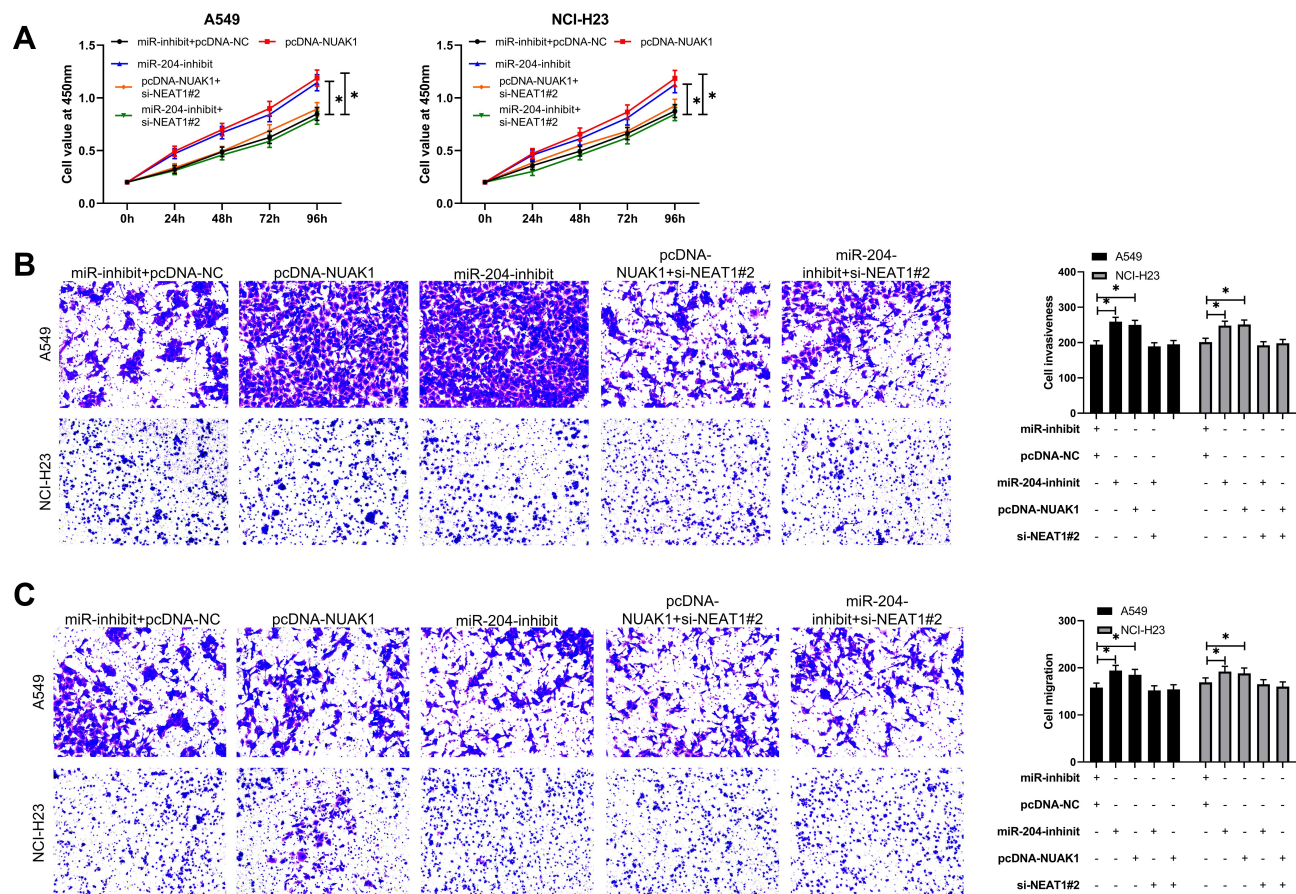


Figure 5 NEAT1 inhibited the proliferation, invasion and migration of LC cells by regulating miR-204/NUAK1 axis. **(A)** CCK-8 test was used to detect cell proliferation after co-transfection. **(B-C)** Transwell test was used to detect the changes of invasion and migration ability of cells after co-transfection. * $P < 0.05$.

invasion of LC cells. In this study, we predicted the potential binding miR of *NEAT1* through 4 online prediction websites and found that there was a targeted connection of *NEAT1* with *miR-204*. Subsequently, we analyzed the *miR-204* in LA through starbase and qRT-PCR. The results showed that the *miR-204* in tumor samples was lower than that in control samples both in the database and in LA tumor tissues. The correlation analysis showed that there was a negative correlation of *miR-204* with *NEAT1*. This indicated that there was a latent targeting connection of *miR-204* with *NEAT1*. Then, we verified it by RIP and double luciferase reporting experiments. Previous studies have revealed that³⁷ *miR-204* is low in LC and is expected to become a potential diagnostic indicator for LC. Moreover, other studies have revealed that³⁸ *miR-204* participates in the development of LC through targeted regulation of downstream genes. *NUAK1* is also known as adenosine monophosphate-related kinase 5 (ARK5). Studies have revealed that *NUAK1* is differentially expressed in nasopharyngeal carcinoma³⁹ and ovarian

carcinoma,⁴⁰ and it is expected to become a potential prognostic indicator. In this study, we found that there was a targeted binding site of *miR-204* with *NUAK1*. The targeted relationship between *miR-204* and *NUAK1* has been confirmed in liver cancer in previous studies by Yu et al.⁴¹ However, it is not clear whether there is the same regulatory relationship in LC. In this research, we concluded that the *NUAK1* in LA patients increased significantly and was negatively correlated with *miR-204*. In addition, we confirmed the targeting relationship between *miR-204* and *NUAK1* in LC through a double luciferase report.

More and more studies have revealed that lncRNA can participate in tumor growth by regulating the miR/mRNA axis.⁴² In the above studies, we found that *NUAK1* might have the ability to regulate the *miR-204/NUAK1* axis, but whether it can play a role is still unclear. Therefore, we conducted a rescue experiment at the end of our research. Through experiments, we found that the growth, invasion and migration of cells transfected with *miR-204*-inhibit and pcDNA-*NUAK1*

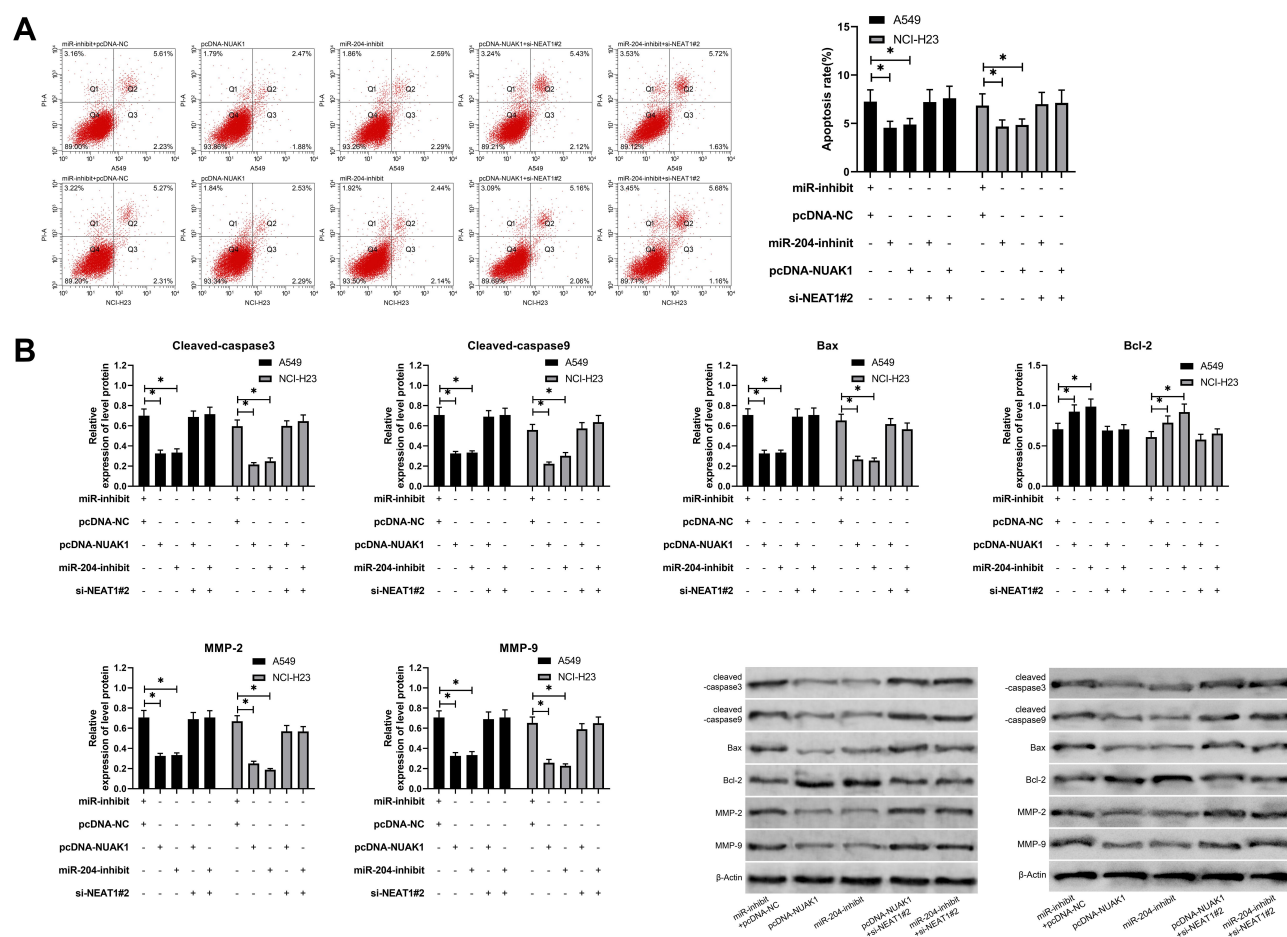


Figure 6 NEAT1 inhibited the apoptosis of LC cells by regulating miR-204/NUAK1 axis. (A) Flow cytometry was used to detect the change of apoptosis rate after co-transfection. (B) WB test was used to detect the relative expression of apoptosis-related proteins, MMP2 and MP9 after co-transfection. *P<0.05.

were significantly faster than those of the control, and the apoptosis of cells was inhibited. However, the above results were reversed since si-NEAT1#2 was co-transfected with miR-204-inhibit or pcDNA-NUAK1. In addition, our research also showed that the proteins of cleaved-caspase 3, cleaved-caspase 9, Bax, MMP-2 and MMP-9 in cells after co-transfection were obviously higher than those of cells transfected with miR-204-inhibit and pcDNA-NUAK1, while Bcl-2 was significantly lower, which suggested that NEAT1 could inhibit the growth of LA by regulating miR-204/NUAK1 axis to inhibit the invasion and migration of LA and promote the apoptosis of LA cells.

In this research, we confirmed the function of NEAT1/miR-204/NUAK1 axis in LA, but there are some limitations in this research. Firstly, we have not conducted in vivo experiments in this study, and whether NEAT1/miR-204/NUAK1 axis plays the same role in vivo still needs to be further verified. In addition, the samples that

we collected in this study are single, and some reports have shown that lncRNA is expressed in various samples. Whether the development of LA can be judged by detecting other samples is also of great significance, so we hope to carry out more research in the future to further supplement our research results.

Conclusion

To sum up, the expression of NEAT1 increases in patients with LA and cell strains, and inhibits the growth of LA by mediating miR-204/NUAK1 axis. It is a potential therapeutic target for LA.

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Disclosure

The authors report no conflicts of interest in this work.

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