Original Article
NEAT1 regulates pancreatic cancer cell growth, invasion and migration though microRNA-335-5p/c-met axis

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Abstract: NEAT1 has been reported to affect cancer progression, which was subsequently confirmed in multiple cancers. Hsa-miRNA-335-5p (miR-335-5p) has recently been identified as an anticancer agent in various organs. However, the relationship between NEAT1 and miR-335-5p remains poorly understood. In this study, we investigated the effects of NEAT1 and miR-335-5p on development of pancreatic cancer. The ectopic expression of miR-335-5p in pancreatic cancer cell lines significantly suppressed cell growth by inhibiting c-met. In addition, downregulating NEAT1 upregulates miR-335-5p. Taken together, our results demonstrate that the NEAT1/miR-335-5p/c-met axis plays a pivotal role in pancreatic cancer by regulating the proliferation, metastasis, and apoptosis of pancreatic cancer cells in vivo and in vitro.

Keywords: Pancreatic cancer, NEAT1, microRNA-335-5p, c-met

Introduction

Pancreatic cancer ranks among the most malignant of human cancers [1]. Its prognosis is extremely poor, with a 5-year relative survival rate of 5% [2] and a median survival of 3.5 months for non-resectable tumors [3]. Surgical resection is the only potentially curative therapy [4], but relapses are common even in these cases [5]. Therefore, the pathological mechanisms of pancreatic cancer urgently need to be understood to facilitate early diagnosis and advance therapeutic modalities and agents.

Recent evidence has suggested a relationship between several long non-coding RNAs (lncRNAs) and metastasis, drug resistance and other clinical outcomes in several types of cancers [6-10]. NEAT1, a nuclear-restricted long non-coding RNA, is known as a transcriptional regulator for numerous genes. NEAT1 was first transcribed from the multiple endocrine neoplasia locus [11], suggesting that this lncRNA affects cancer progression, which was subsequently confirmed in multiple cancers and various studies [12-18]. However, the emerging potential role of NEAT1 in pancreatic cancer remains unclear.

MicroRNAs (miRNAs) are a class of small non-coding regulatory RNAs, which can broadly regulate target genes by binding to a complementary sequence in their 3'UTR [19, 20]. miRNAs play important roles in tumor development by regulating the expression of various oncogenes and tumor suppressor genes [21]. For example, miR-199a suppresses the tumorigenicity and multidrug resistance of ovarian cancer-initiating cells [22], whereas miR-27a reverses the multidrug resistance phenotype by regulating the expression of MDR1 and β-catenin [23]. Furthermore, miR-146b-5p suppresses the translation of EGFR, binds to the EGFR 3'UTR, and inhibits the migration of glioma cells [24]. Similarly, miR-335 activates the p53 tumor suppressor pathway to limit cell proliferation and neoplastic cell transformation [25]. miR-335 also targets Bcl-w and negatively regulates the invasiveness of ovarian cancer cells [26]. In addition, miR-335 inhibits the proliferation and migration of human mesenchymal stem cells by targeting RUNX2 [27] and is involved in regulating target genes in several oncogenic signal-pathways, such as p53, MAPK, TGF-β, Wnt, ERβ, mTOR, Toll-like receptor and FAK (focal adhesion kinase) [28]. However, the mechanism and the role of miR-335 in regulation of pancreatic cancer remain unknown.
The molecular targeting of oncogenes as a therapeutic approach is currently being intensively investigated. Specifically, the identification of deregulated oncogenic pathways in pancreatic cancer will lead to new therapeutic options. To this end, c-Met, a tyrosine kinase receptor, is overexpressed in a subset of human epithelial malignancies [29] including colorectal [30, 31], gastric [32, 33], ovarian [34, 35], endometrial [36], breast [37, 38], prostate [39] and hepatocellular [40] carcinomas. This overexpression may be the result of c-Met amplification [31].

In this study, we showed for the first time that miR-335-5p directly targets and regulates human c-met gene. Collectively, we discovered that NEAT1 promotes pancreatic cancer cell growth, migration and invasion while inhibiting cell apoptosis. Moreover, miR-335-5p also inhibits pancreatic cancer cell growth, migration and invasion and promotes cell apoptosis by targeting the 3'UTR of c-Met. Thus, we uncovered a pathway that is established by NEAT1/miRNAs/c-met axis, which promotes pancreatic cancer malignancy.

Materials and methods

Cell culture

Human pancreatic cancer cell lines (PANC-1, SW1990, CAPAN-1, JF305 and PC-3) and the nonmalignant HPC-Y5 cell line were obtained from the American Type Culture Collection (ATCC, Manassas, VA, USA) and cultured in Dulbecco's modified Eagle's medium (DMEM; Sigma-Aldrich, St. Louis, MO, USA) supplemented with 10% fetal bovine serum (FBS; Sijiqing biochemical, Hangzhou, China) at 37°C in a humidified 5% CO₂ atmosphere. The cells were transfected using Lipofectamine 2000 reagent (Invitrogen, USA) following the manufacturer's instructions.

Clinical samples and RNA isolation

Fifteen paired human pancreatic cancer and matched adjacent normal tissue samples from the same patient were collected with patient consent at the time of surgery. The tumors were graded according to the WHO criteria (World Health Organization, 2008). The tissue specimens and clinical information were obtained as part of a study approved by the Institutional Review Board at Xinhua Hospital of Shanghai Jiaotong University, China. Total RNA was extracted from pancreatic cancer cells using TRIzol Reagent according to the manufacturer's instructions (Invitrogen, Carlsbad, CA, USA).

Real-time quantitative PCR analysis

Total RNA (5 µg) was reverse transcribed into cDNA using M-MLV reverse transcriptase (Promega, USA) with specific primers. The cDNA was used as template to amplify either mature miR-335-5p or an endogenous control U6 snRNA by PCR. NEAT1 or an endogenous control, GAPDH, were also amplified. This PCR was followed by SYBR-Green real-time PCR (Qiagen, Hilden, Germany). The PCR procedure was as follows: 94°C for 3 min, followed by 40 cycles of 94°C for 30 s, 50°C for 30 s and 72°C for 30 s. The relative gene expression levels were calculated using the 2ΔΔct method using U6 snRNA or GAPDH mRNA as an internal control.

Plasmid construction and cell transfection

We commercially synthesized the 2'-O-methyl-modified antisense oligonucleotides of miR-335-5p (ASO-miR-335-5p) to inhibit miR-335-5p. The 3'UTRs of c-met and NEAT1, which contain the miR-335-5p binding site, and the mutant 3'UTR fragment, which contains the mutant binding site of miR-335-5p, were obtained by annealing double-strand DNA and inserted into the pmirGLO vector at the BamHI and EcoRI sites.

The pSilencer/shR-NEAT1 plasmid, which expresses a siRNA that targets NEAT1 transcription, was constructed by annealing single-strand hairpin cDNA and inserting it into a pSilencer2.1-U6 neo vector (Ambion, Austin, TX, USA) at the BamHI and HindIII sites. The full-length sequences of human NEAT1 and c-met cDNA deposited in Genbank were cloned into the EcoRI/Xhol restriction sites of pcDNA3. The resultant plasmids were named pcDNA3/NEAT1 and pcDNA3/c-met. The promoter of miR-101 was amplified from genomic DNA and cloned into the KpnI/EcoRI restriction sites of pGL3-Basic (Promega) upstream of the firefly luciferase gene. At 60% confluency, the cells were transfected with plasmids using Lipofectamine 2000 Transfection Reagent (Invitrogen, Carlsbad, CA, USA) according to the manufacturer's instructions.
Cell proliferation assay

Cell proliferation was measured using an MTT assay. Transiently transfected cells were seeded in a 96-well plate at a cell density of 1.0 × 10⁴ cells/well and then cultured for 12 h intervals for a total of 2 days. Subsequently, MTT solution (0.2 mg/ml, Sigma-Aldrich, St Louis, MO, USA) was added to each well, and the cells were incubated for an additional 4 h. The solution was then carefully aspirated, and 150 μL DMSO was added to each well to dissolve the crystal. The absorbance was determined at 570 nm using a microplate reader (Bio-Tek Instruments, Winooski, VT, USA).

Cell apoptosis assay

The fraction of apoptotic cells was determined with the Annexin V-7-ADD apoptosis detection kit (Roche, Switzerland). Briefly, 48 hours (h) after transfection, the cells were collected and washed twice with cold PBS buffer, resuspended in 200 μl of binding buffer, and incubated with 20 μl of Annexin-V-R-PE for 20 minutes in a dark ice bath. Subsequently, 10 μl of 7-AAD was added, and the cells were then analyzed by flow cytometry. Cells treated with DMSO were used as the negative control.

Cell migration and invasion assays

For the migration assay, 1.0 × 10⁵ cells were suspended in serum-free medium and plated in Transwell chambers chambers (Corning Costar, NY, USA). Briefly, medium containing 10% FBS was added to the lower chamber as a chemoattractant. After incubating the cells for 24 h at 37°C in 5% CO₂, they were fixed in methanol for 15 min and stained with 0.05% crystal violet in PBS for 15 min before being counted under a microscope (Olympus, Tokyo, Japan). Cells that had not invaded upper chamber were removed by wiping the surface with a cotton swab, and invasive cells were fixed with 4% formaldehyde in PBS and subsequently stained with 1% crystal violet in 2% ethanol. The cells on the lower surface of the filter were photographed under a light microscope (100 × magnification).

These procedures were also followed for the invasion assay, but the filters were pre-coated with 100 μl of Matrigel (BD Biosciences, CA, USA) at a 1:4 dilution in DMEM to mimic a basement membrane.

DNA hybridizations

DNA fragments were transferred from agarose gels to nylon membranes (Amersham) by vacuum blotting. The probes were labeled by nick translation and hybridizations with [α-32P] dCTP (Amersham) according to the manufacturer’s instructions (Promega, Madison, Wisconsin, USA). The partial nucleotide fragments internal to the 16S rDNA (Accession No. AY509240.1) and cat gene (GenBank accession AJ132968.1) were used as probes in this work. The sequences were analyzed with the MEGALIGN program (DNA Star, Madison, WI).

Western blot analysis

Total cellular extracts were obtained using RIPA buffer on ice. Equal amounts of protein (30 μg protein per lane) were separated by SDS-PAGE and transferred to PVDF membranes (Millipore, Boston, MA, USA). The immunoblots were blocked with 5% skim milk in TBS/Tween 20 (0.05%, v/v) at room temperature for 1 h. The membrane was then probed primary antibodies overnight at 4°C. The following primary antibodies were used: anti-c-met, anti-actin and HRP-conjugated anti-rabbit (Santa Cruz Biotechnology, Santa Cruz, CA, USA). Subsequently, the membranes were incubated with horseradish peroxidase-conjugated secondary antibody for 1 h at room temperature. The blots were then developed using an enhanced chemiluminescence western blotting detection system (Amersham Bioscience, UK).

Target prediction and dual luciferase reporter assay

Based on bioinformatic predictions (TargetScan (http://www.targetscan.org/mamm_31/), miRDB (http://www.mirdb.org/miRDB/) and PicTar (http://www.pictar.org/)), c-met was selected as candidate target of miR-335-5p. The 3’UTR segments of c-met containing putative binding sites for miR-335-5p were obtained by annealing and inserted into the pmirGLO vector. The wild-type reporter construct pmirGLO/c-met-3’UTR and the mutant reporter construct pmirGLO/c-met-3’UTR mut, in which a site that perfectly complements miR-335-5p was mutated by annealing, were used for miRNA functional analysis. The wild-type and mutant insertions were confirmed by DNA sequencing. All primer information is available.
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For the luciferase reporter experiments, PANC-1 and SW990 cells were co-transfected with the miR-335-5p in a 48-well plate followed by the pmirGLO/c-met-3'UTR reporter vector or the pmirGLO/c-met-3'UTR mut. The firefly luciferase and Renilla luciferase levels were measured 48 h after transfection. Each experiment was repeated at least three times.

**In vivo metastatic assay and bioluminescent imaging**

The animal procedures were performed in accordance with the Institutional Animal Care and Use Committee guidelines of the Experiment Animal center of the Fourth Military Medical University. Athymic 4-week-old female nude mice were obtained from the Shanghai Laboratory Animal Center of China and housed under standard conditions. Each mouse was injected via the tail vein with $2 \times 10^6$ cells suspended in 100 μl of phosphate-buffered saline. To track cells in vivo, the cells were stably transfected with firefly luciferase. Pancreas metastases was quantitated in vivo by imaging based on bioluminescence. Mice to be imaged were injected with 150 mg/kg of D-luciferin (Xenogen, Hopkinton, MA) intraperitoneally in 100 μl of phosphate-buffered saline and then anesthetized with a continuous flow of pento-barbital. Ten minutes later, bioluminescence images of the mice were obtained with the IVIS Imaging System (Xenogen) and then analyzed using the IVIS Living Image software (Xenogen) software.

**Statistical analysis**

Each experiment was repeated at least three times. The quantitative data between groups were compared and analyzed by Student’s t-test (two tailed) or a one-way analysis of variance. The data are expressed as the means ± standard deviation (SD), and P ≤ 0.05 was considered to indicate a significant difference using the Students-Newman-Keuls test.

**Results**

**NEAT1 is up-regulated in pancreatic cancer**

We used quantitative real time RT-PCR to determine the expression of NEAT1 in 15 human pancreatic cancer tissues and adjacent normal tissues (Figure 1A). Moreover, real-time PCR showed that NEAT1 expression was significantly overexpression in the pancreatic cancer cell lines PANC-1, SW1990, CAPAN-1, JF305 and PC-3 relative to the prostate cell line HPC-Y5 (Figure 1B). In general, NEAT1 expression was significantly upregulated in pancreatic cancer.

**NEAT1 promotes the long-term proliferation and inhibits the apoptosis of pancreatic cancer cell lines**

To research the function of NEAT1 in pancreatic cancer, we constructed si-NEAT1 to knock
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A

Relative expression of NEAT1

B

Absorbance (OD 595nm)

SW1990

0h 12h 24h 48h

PANC-1

0 1.0 2.0 3.0 4.0

si-NC si-NEAT1

C

si-NC

si-NEAT1

PANC-1

Migration

Invasion

SW1990

Migration

Invasion

D

Apoptotic cells (%)

PANC-1

0 5 10 15 20 25

si-NC si-NEAT1

SW1990

0 5 10 15 20

si-NC si-NEAT1

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Figure 2. NEAT1 promotes long-term proliferation and inhibits the apoptosis of the pancreatic cancer cell lines. A. NEAT1 expression levels measured by real-time RT-PCR. Total RNA was extracted from PANC-1 and SW1990 cells transfected with si-NEAT1 or si-NC, and GAPDH served as an endogenous control. The relative NEAT1 expression level (mean ± SD) is shown (*P < 0.05). B. Cell proliferation was assessed with an MTT assay. After PANC-1 and SW1990 cells were transfected with the si-NEAT1 or si-NC, the MTT assay was used to determine relative cell growth 0, 12, 24 and 48 h after transfection. The relative cell growth was normalized to the growth activity of PANC-1 and SW1990 cells in the control groups (*P < 0.05). C. Transwell migration and Matrigel invasion assays were used to evaluate the migration and invasion of PANC-1 and SW1990 cells transfected with si-NEAT1 or si-NC. Representative fields of migrating or invasive cells on the membrane. The data represent three independent experiments (*P < 0.05). D. The incidence of apoptosis was studied by flow cytometry. The cells were stained with annexin V-fluorescein isothiocyanate and counterstained with 7-ADD. The data represent three independent experiments (*P < 0.05).

down its expression in pancreatic cancer cell lines. First, we measured the NEAT1 mRNA levels in PANC-1 and SW1990 cells transfected with si-NEAT1 or si-NC by real time RT-PCR. The expression of NEAT1 was significantly decreased in PANC-1 and SW1990 cells transfected with si-NEAT1 compared with NC cells (Figure 2A). The viability of pancreatic cancer cells transfected with si-NEAT1 was evaluated with the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl-tetrazolium bromide (MTT) assay; si-NEAT1 reduced cell viability 12, 24 or 72 h after transfection (Figure 2B). In parallel, we analyzed the migration and invasiveness of the pancreatic cancer cell lines PANC-1 and SW1990. We examined the migration and invasiveness of pancreatic cancer cells transfected with si-NEAT1 and found that cells transfected with si-NEAT1 migrated less and were less invasive than control cells (Figure 2C). These data demonstrated that NEAT1 might promote migration and invasion in pancreatic cancer. The fluorescence-activated cell sorting (FACS) analysis showed that reduced NEAT1 expression resulted in pancreatic cancer cell apoptosis. NEAT1 suppression significantly decreased the percentage of total apoptotic cells (early apoptotic + late apoptotic) compared with si-NC in PANC-1 and SW1990 cells (Figure 2D). These results indicated that NEAT1 promoted proliferation and decreased apoptosis in pancreatic cancer cells.

NEAT1 decreases the expression of miR-335-5p in the pancreatic cancer cell lines PANC-1 and SW1990

To study the mechanisms by which NEAT1 promotes the proliferation and inhibits apoptosis of the pancreatic cancer cells, we selected miR-335-5p as the research target. The expression level of miR-335-5p in PANC-1 or SW1990 cells transfected with pcDNA3/NEAT1 or control was measured in a validation experiment by qRT-PCR, which indicated a clear decrease in the level of miR-335-5p in transfected PANC-1 and SW1990 cells (Figure 3D). The RT-PCR relative quantification procedure and the miR-335-5p and ncRNA sequences are shown (Figure 3A and 3B). Moreover, the miR-335-5p expression level in PANC-1 and SW1990 cells transfected with si-NEAT1 or si-NC was measured, and this experiment showed that a decrease in NEAT1 expression resulted in the upregulation of the NEAT1-targeting miRNA miR-335-5p (Figure 3C). Thus, we investigated the effect on miR-335-5p on NEAT1 expression. To this end, we transfected PANC-1 cells with the wild-type (WT) or mutated (Mut) version of the luciferase-NEAT1 3'-UTR reporter vector as well as the miR-335-5p or scramble. This experiment showed that miR-335-5p, but not the mutant, reduced the intensity of the luciferase-NEAT1 WT (Figure 3E). A Southern blot hybridizations assay was employed to further verify the relationship between NEAT1 and miR-335-5p, which indicated that NEAT1 was enriched more than 20-fold in miR-335-5p in PANC-1 or SW1990 cells (Figure 3F). These results show that miR-335-5p is suppressed by NEAT1 overexpression due to combination and interaction.

miR-335-5p inhibitor counteracts NEAT1 suppression

First, we detected the miR-335-5p expression level in PANC-1 and SW1990 cells transfected with either pSilencer NC, pSilencer/si-NEAT1, or pSilencer/si-NEAT1 together with miR-335-5p inhibitor by qRT-PCR. Transfecting cells with pSilencer/si-NEAT1 increased the expression of miR-335-5p compared with NC cells; miR-335-5p inhibitor reversed this effect (Figure
An MTT assay was then used to characterize the effects of NEAT1 on tumor cell viability. The results of this assay indicated that knocking down NEAT1 inhibited the viability of PANC-1 and SW1990 cells, and this effect was counteracted by miR-335-5p inhibitor (Figure 4B and 4C). In parallel, cell invasion and migration assays were performed using Transwell chambers with or without Matrigel. Specifically, PANC-1 and SW1990 cells co-transfected with miR-335-5p inhibitor and pSilencer NC, pSilencer/si-NEAT1, or pSilencer/si-NEAT1 were seeded in Transwell chambers, and images were taken to count the number of cells. The data demonstrate that pSilencer/si-NEAT1 can suppress the migration and invasiveness of cells relative to the control, but this suppression is rescued by miR-335-5p inhibitor (Figure 4D and 4E). Next, we studied the effect of NEAT1 on apoptosis. The fluorescence-activated cell sorting (FACS) analysis showed that reduced NEAT1 expression resulted in pancreatic cancer cell apoptosis. Furthermore, miR-335-5p inhibitor counteracted the effect of si-NEAT1 (Figure 4F). Taken together, these data show that NEAT1 suppression inhibits the long-term proliferation and increases apoptosis of pancreatic cancer cell lines, and these effects are rescued by an miR-335-5p inhibitor.
c-met is a direct target of miR-335-5p

Based on the above discussion, NEAT1 suppression upregulates the expression of miR-335-5p and simultaneously regulates long-term proliferation and increases apoptosis in pancreatic cancer cell lines. To determine the mechanism of miR-335-5p-mediated cell dysregulation in pancreatic cells, we next identified target genes that could be responsible for the effect of miR-335-5p. Potential target genes of miR-335-5p were predicted using miRanda, TargetScan and PicTar, which suggested that miR-335-5p targets c-met (Figure 5A). We performed luciferase reporter assays to examine whether miR-335-5p interacts directly with its target c-met. We constructed a series of 3'UTR fragments, including the wild-type c-Met 3'UTR and a binding site mutant (Figure 5A). These fragments were then inserted into the pmirGLO luciferase reporter plasmid. Co-transfecting miR-335-5p and the wild-type c-met 3'UTR into PANC-1 and SW1990 cells significantly decreased the luciferase signal compared with the controls. However, co-transfecting mutant c-met 3'UTR and miR-335-5p mimics failed to alter the luciferase intensity (Figure 5B and 5C). Furthermore, the overexpression of miR-335-5p reduced c-met mRNA and protein expression in PANC-1 and SW1990 cells, whereas miR-335-5p inhibitor increased the level of c-met (Figure 5D-F). Taken together, these results suggest that miR-335-5p binds directly to the 3'UTR of c-met, thereby repressing gene expression.

NEAT1 and miR-335-5p regulate pancreatic cancer cell growth, invasion and migration via c-met

Next, we explored the relationship between NEAT1, miR-335-5p and c-met. The activation of c-Met promotes tumor cell proliferation, migration, invasion. Therefore, we transfected PANC-1 and SW1990 cells with pSilencer/shRN-NEAT1, pSilencer/shRN-NEAT1 and c-met, miR-
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335-5p mimic or miR-335-5p mimic and c-met. An MTT assay was used to assess cell growth at 0, 24 and 48 h after transfection. C-met increased cell proliferation relative to the pSilencer/shR-NEAT1 or miR-335-5p mimics group (Figure 6A and 6B). In addition, Colony formation assays were performed to confirm this tendency (Figure S1). Next, we studied the...
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The effect of c-met on apoptosis. The fluorescence activated cell sorting (FACS) analysis showed that the forced expression of c-met inhibited pancreatic cancer cell apoptosis (Figure 6C). Moreover, Transwell migration and Matrigel invasion assays revealed that c-met overexpression increased the number of migrating cells (Figure 6D and 6E). Thus, the downregulation of miR-335-5p induces c-met expression to inhibit pancreatic cancer malignancy. Furthermore, NEAT1 suppresses miR-335-5p transcription, which increases c-met translation.
and consequently promotes pancreatic carcinogenesis (Figure 6F). The data demonstrate that NEAT1 and miR-335-5p regulate pancreatic cancer cell growth, invasion and migration via c-met in vitro.

**NEAT1 and miR-335-5p regulate pancreatic cancer cell tumor growth via c-met in vivo**

To further confirm the above assumptions, we treated PANC-1 and SW1990 xenograft tumor-bearing nude mice stably transfected with Firefly luciferase with pSilencer/shR-NEAT1, pSilencer/shR-NEAT1 and c-met, miR-335-5p mimic or miR-335-5p mimic and c-met. Ten minutes later, bioluminescence images of the mice were obtained using the IVIS Imaging System (Xenogen) 10 min later and then analyzed the images using the IVIS Living Image (Xenogen) software. Compared with the controls, high-density signals were captured in the c-met group (Figure 7A-D). These data reveal that NEAT1 and miR-335-5p regulate pancreatic cancer cell tumor growth via c-met in vivo.

**Discussion**

Here, we identified a NEAT1/miR-335-5p/c-met axis that is involved in pancreatic cancer development and progression. We found that knocking down NEAT1 IncRNA significantly impaired the growth and migration of pancreatic cancer cells in vitro. These findings corroborated previous findings in NSCLC [14] and HCC [41], which...
showed that knocking down NEAT1 expression in vitro inhibited the migration and invasiveness of cancer cells. Moreover, we found that the expression of NEAT1 increased the level of miR-335-5p. Conversely, aberrant miR-335-5p expression suppressed the intensity of the luciferase signal of NEAT1. Thus, the expression warrants further investigation with a dual luciferase reporter assay. The biological mechanisms of NEAT1 regulation in pancreatic cancer currently remain poorly understood.

To further investigate the mechanisms of NEAT1 in pancreatic cancer, we studied miR-335-5p and its target mRNA. First, we predicted putative targets of miR-335-5p using the prediction programs TargetScan, PicTar, and miRanda. Subsequently, the luciferase reporter assay identified c-met as the target because transfection with miR-335-5p reduced the luciferase activity of constructs carrying the target c-met fragment. Furthermore, the ectopic expression of miR-335-5p simultaneously reduced the mRNA and protein levels of c-met. Gao et al. also found that the oncogene c-met is a target gene of miR-335 in breast cancer cells [42]. MET can act as an oncogene, and its signaling plays essential roles in regulating tumorigenesis in various cancers, such as lung cancer [43], including NSCLC [44].

Next, we explored the relationship between NEAT1, miR-335-5p and c-met. By co-transfecting shR-NEAT1 and c-met or miR-335-5p mimic and c-met into PANC-1 and SW1990 cells, we found that c-met increased cell proliferation and inhibited pancreatic cancer cell apoptosis. Thus, miR-335-5p reduced c-met expression, and NEAT1 overexpression suppressed miR-335-5p transcription, leading to an increase in c-met translation, which promoted pancreatic cancer malignancy. Moreover, the xenograft tumor-bearing nude mice experiment revealed that NEAT1 and miR-335-5p regulate pancreatic cancer cell tumor growth via c-met in vivo.

In conclusion (Figure 6F), our findings suggest that NEAT1 regulates pancreatic cancer progression via the microRNA-335-5p/c-met axis. This axis can be triggered by other extrinsic signaling from the tumor microenvironment or cancer cell intrinsic signaling. Although many factors contribute to the development of pancreatic cancer, our study provides a new approach to delay the development of pancreatic cancer.

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Disclosure of conflict of interest

None.

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References


**Figure S1.** NEAT1 and miR-335-5p regulate pancreatic cancer cell proliferation via c-met. Effect of NEAT1 and miR-335-5p on cell proliferation as evaluated by a colony formation assay. PANC-1 (upper panel) and SW1990 (lower panel) cells transfected with si-NEAT1, si-NEAT1 and c-met, miR-335-5p mimic or miR-335-5p mimics and c-met were seeded in 12-well plates (left to right). The figure shows cells 10 days after seeding; the cells were stained with crystal violet.