Original Article
The FOXM1/ATX signaling contributes to pancreatic cancer development

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Abstract: Both autotaxin (ATX) and Forkhead Box M1 (FOXM1) have been commonly recognized as oncogenes in multiple types of human malignancies. However, the expression and biological functions of ATX in pancreatic ductal adenocarcinoma (PDAC), and its correlation with FOXM1 are poorly understood. The present study aimed to investigate their correlation and biological consequences in PDAC development. By dual luciferase reporter and chromatin immunoprecipitation assays, we found that ATX was a downstream transcriptional target gene of FOXM1. Further cellular functional experiments indicated that ATX was required for FOXM1-mediated PDAC cell proliferation and migration. Data from molecular biological experiments showed that ATX could enhance FOXM1 expression in turn by inhibiting the Hippo signaling pathway, suggesting that ATX and FOXM1 formed a positive feedback loop to facilitate PDAC progression. Using immunohistochemistry (IHC) method, both ATX and FOXM1 expression were found to be frequently up-regulated in PDAC tumor tissues when compared with adjacent normal tissues, and elevated ATX and FOXM1 expression were positively correlated with each other. In conclusion, the present work identified a positive feedback loop between ATX and FOXM1 which promotes PDAC cell proliferation and migration.

Keywords: ATX, FOXM1, pancreatic cancer, proliferation, migration

Introduction

As the most common type of pancreatic cancer, pancreatic ductal adenocarcinoma (PDAC) is one of the most lethal human malignancies worldwide, with an extremely low 5-year survival rate [1, 2]. Although big efforts have been made to improve the prognosis of patients with PDAC in the past decades, little improvement has been achieved due to the high tendency of PDAC to metastasize and resistance to chemoradiation therapy [3]. Therefore, a better understanding of the molecular mechanisms underlying the tumorigenesis and metastasis of PDAC is needed to identify novel potential therapeutic targets for the clinical management of PDAC.

Autotaxin (ATX), also known as nucleotide pyrophosphatase-phosphodiesterase 2 (ENPP2) [4], is a secreted glycoprotein that primarily catalyzes the hydrolysis of lysophosphatidylcholine to produce lysophosphaticid acid (LPA) [5], which is well-known to stimulate the proliferation, migration, and survival of many human malignancies [6, 7]. High expression of ATX has been reported in different tumors and ATX overexpression contributes to tumor proliferation and invasion mainly through the induction of LPA [6, 8, 9]. In PDAC, increased serum ATX activity was found in patients with pancreatic cancer [10], and higher expression of ATX was observed in a highly invasive PANC-1 cell line [11]. Although these studies strongly suggest an aggressive role of ATX in PDAC development, its expression and biological functions in PDAC are largely unknown, and the molecular mechanisms that regulate ATX expression in PDAC as of yet remain unclear.

Forkhead box M1 (FOXM1) is a key member of the forkhead family of transcription factors [12, 13]. Numerous studies have reported that FOXM1 plays a critical role in regulating cell cycle transition, cell proliferation and chromosome stability [13-16]. Like ATX, FOXM1 is also highly expressed in various types of human malignancies including PDAC [17, 18], and overexpression of FOXM1 can stimulate tumor cell
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proliferation, invasion and angiogenesis in different tumors [17, 19]. Although over-activation of both ATX and FOXM1 occur in many human cancers, it is unclear whether they are functionally correlated with each other. In the present study, we resorted to investigate their correlation and underlying molecular mechanisms in PDAC development.

Methods

Tissue microarray (TMA) and immunohistochemistry (IHC) analysis

For IHC analysis, a human PDAC TMA (#HLiv-HPanA150Su01) containing 60 tumor tissues and paired adjacent normal tissues as well as additional 30 tumor tissues derived from 90 patients with PDAC was purchased from Shanghai Outdo Biotech Co., LTD., China. Standard IHC procedures were performed using specific antibodies against ATX (1:500, Abcam, #ab77104) and FOXM1 (1:200, Santa Cruz Biotechnology, #sc-502). The staining results were evaluated by two pathologists blinded to the clinical information independently according to the percentage of ATX/FOXM1-positive cells and staining intensity. Specifically, the percentage of ATX/FOXM1-positive cells was classified into 5 groups: < 10% (0), 10-25% (1), 25-50% (2), 50-75% (3), and > 75% (4). The staining intensity was divided into 4 groups: no staining (0), light brown (1), brown (2), and dark brown (3). Finally, the overall scores were calculated using the following formula: overall score = percentage score × intensity score. The samples with an overall score ≤ 6 were defined as weak, and > 6 were defined as strong.

Cell culture and reagents

Human PDAC cell lines PANC-1 and SW1990 were purchased from Shanghai Cell Bank of the Chinese Academy of Sciences Shanghai, China. Cells were cultured in Dulbecco’s modified Eagle’s medium (DMEM, Corning, Inc., Corning, NY, USA) supplemented with 10% fetal bovine serum (FBS) and 1% penicillin/streptomycin (M&C Gene Technology Ltd., Beijing, China). All of these cell lines were maintained in a humidified incubator at 37°C with 5% CO₂. The Hippo signaling pathway (MST1/2) inhibitor XMU-MP-1 was purchased from Selleckchem, Houston, TX, USA.

Cell Transfection

The FOXM1-expressing plasmids pcDNA3.1-FOXM1 (isoform C) were gifts from Dr. Xie Keping (The University of Texas MD Anderson Cancer Center, Houston, TX, USA), and lentivirus carrying FOXM1-shRNA or FOXM1-ORF plasmids and the siRNAs against ATX (2’-O-methyl modified) were synthesized by GenePharma Co., Ltd., Shanghai, China. PDAC cells were transfected with plasmids or siRNAs using Lipofectamine 3000 (Invitrogen, USA) according to the protocol of the manufactures. To generate lentiviral stable cell lines, PDAC cells were infected with the lentiviruses described above with the assistance of polybrene (4 μg/ml), and puromycin (2 μg/ml, Sigma, MO, USA) was used to selected stably infected PDAC cells.

Western blots analysis

PDAC cells were lysed using a Radioimmunoprecipitation assay (RIPA) buffer supplemented with a protease inhibitor cocktail (Sigma-Aldrich; Merck KGaA, Darmstadt, Germany) to extract total proteins. Standard western blot analysis was performed using primary antibodies anti-FOXM1 (1:200, Santa Cruz Biotechnology, #sc-502), anti-ATX (1:500, Abcam, #ab77104), and anti-β-actin (1:1000, Santa Cruz Biotechnology, #sc-81178), anti-mouse (Cell Signaling Technology) and anti-rabbit (Cell Signaling Technology) antibodies were used as secondary antibodies. The bands were visualized and quantified using a Li-Cor Odyssey protein imaging system (LI-COR Biosciences, Lincoln, NE, USA).

Quantitative real-time polymerase chain reaction (qPCR) analysis

Total RNAs were extracted from PDAC cells with Trizol reagent (Sigma-Aldrich, MO, USA) and quantified by a Fisher Scientific NanoDrop™ 2000 spectrophotometer (NanoDrop; Thermo Fisher Scientific, Inc., Wilmington, DE, USA) according to the manufacturer’s specifications, after which 1 μg RNA was reverse-transcribed into cDNA using the PrimeScript™ RT reagent kit (Takara Bio, Inc., Otsu, Japan). Standard qPCR analysis was performed with an ABI QuantStudio™ 6 Flex system (Thermo Fisher Scientific, Inc.) using SYBR green reagent (TaKaRa, Japan). Relative mRNA expression levels of FOXM1 and ATX were determined.
using the 2-ΔΔCt method and β-actin was used as an internal control. The following qPCR primers were used: For ATX, 5'-ACAACGGAGGACCTGCAAT-3' (forward), 5'-AGAAGTCCAGCTTGGTGA-3' (reverse); for FOXM1, 5'-GGCCGCACGGCGGAAATGAGA-3' (Forward), 5'-CCACTCCTCTCAAGGAGGCTC-3' (Reverse); for β-actin: 5'-CTGGCAACCAGCAATG-3' (forward), 5'-GGGCCGACTGTCATATCT-3' (reverse).

Colony formation assay

Briefly, PDAC cells were seeded into a 6-well plate at a density of 1 × 10^3 cells per well and then incubated for 2 weeks at 37°C with 5% CO_2. 4% paraformaldehyde and crystal violet were used for the fixation and staining of the colonies, respectively. Next, the colonies were photographed, and the number of colonies was counted with image J software. Each experiment was performed in triplicate and repeated at least three times independently.

Cell proliferation assay

Cell viability of PDAC cells were evaluated using Cell Counting Kit-8 (CCK-8, Dojindo, Kumamoto, Japan) according to the manufacturer’s protocol. Briefly, the selected PDAC cells were seeded into a 96-well plate (3 × 10^3 cells/well) and incubated under normal conditions. At each scheduled time point (0, 24, 48, 72, 96 h), 10 μl of CCK-8 reagents were added into each well, and after incubation for 1 h at 37°C, the absorbance was measured using an automated microplate reader (SpectraMax M5, Molecular Devices LLC, CA, USA) at a wavelength of 450 nm.

Cell migration assay

Cell migration assay was conducted using a 24-well Transwell filter (pore size, 8 μm; Costar; Corning, Inc.). In brief, 800 μl DMEM containing 10% FBS was added to the bottom chamber, and 3 x 10^4 cells maintained in 400 μl DMEM without FBS were added into the upper chamber. After incubation under normal conditions for 24 h, the cells that stayed in the upper chamber were removed with a cotton swab, and the migrated cells were fixed in 4% paraformaldehyde for 15 min followed by staining with crystal violet for additional 15 min. Finally, five random fields were selected and the migrated cells were photographed and counted under an upright metallurgical microscope (Leica, Germany).

Dual luciferase reporter assay

To construct ATX promoter-activated luciferase reporter plasmid, wild or mutant ATX promoter was cloned into pGL3 basic luciferase reporter vector and the products were verified by DNA sequencing. A FOXM1 promoter-activated luciferase reporter plasmid containing 5' FOXM1 sequences from -2430 to +66 bp relative to the transcription initiation site (designated as pFOX-M1-2496) was constructed as described previously [20]. To conduct luciferase reporter assay, PDAC cells were seeded into a 24-well plate and transfected with the indicated promoter reporters. 24 h after transfection, the Dual-Glo® Luciferase Assay Kit (Promega, WI, USA) was used to evaluate the promoter activities of ATX or FOXM1 by quantifying both firefly and Renilla luciferase activity on a GloMax® 96 Microplate Luminometer (Promega, WI, USA).

Chromatin immunoprecipitation (ChIP) assay

ChIP assay was performed on PANC-1 cells using a ChIP assay kit (Millipore, CA, USA) according to the manufacturer’s instructions. Briefly, an aliquot of the immunoprecipitated DNA samples was subjected to PCR and two primers were used to amplify the FOXM1-binding sites in the ATX promoter. Afterwards, the PCR products were resolved electrophoretically with a 2% agarose gel in 1 x TAE and the results were visualized using the ChemiDoc MP system (Bio-Rad Laboratories, Hercules, CA, USA). ChIP-qPCR analysis was performed using ChIP samples and SYBR green reagent (TaKaRa, Japan) according to the manufactures’ protocols. The primers used for ChIP assays were as follows: #1 site: 5'-AGCCATTGCACTCCAGCCT-3' (forward) and 5'CTGGTTGATCCCAATATCC-3' (reverse); #2 site: 5'-GAATCTGTAATGAAACCCA-3' (forward) and 5'-TGCGTAAACAAATACGAGAC-3' (reverse).

Statistical analysis

Data was shown as mean ± standard error of the mean (SEM). Categorical data was analyzed using χ^2 test, the significance of the data from independent experiments were determined using the two-tailed Student t-test or one-way analysis of variance (ANOVA), and the correla-
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Figure 1. Regulation of ATX expression by FOXM1 in PDAC cells. (A and B) Proteins and RNAs were isolated from PANC-1 and SW1990 cells stably infected with LV-FOXM1 lentivirus or control lentivirus, and western blot (A) and qPCR (B) analyses were performed, respectively. (C and D) Proteins and RNAs were extracted from PANC-1 and SW1990 cells with stable expression of FOXM1, respectively. And western blot (C) and qPCR (D) analyses were carried out to determine the effects of FOXM1 on ATX expression. * P < 0.05, ** P < 0.01.

Results

ATX expression is positively regulated by FOXM1 in PDAC cells

To evaluate the relationship between ATX and FOXM1 expression in PDAC, we generated stable PANC-1 and SW1990 cells with FOXM1 knockdown, and ATX expression was examined by western blots and qPCR analyses. As shown in Figure 1A and 1B, both protein and mRNA levels of ATX in PANC-1 and SW1990 cells were significantly decreased when FOXM1 was knocked down. On the contrary, ATX expression was enhanced in FOXM1 stably overexpressed PANC-1 and SW1990 cells (Figure 1C, 1D). These findings suggested that FOXM1 has a positive effect on ATX expression in PDAC cells.

FOXM1 is a transcriptional activator of ATX

Considering that FOXM1 functions as an important transcription factor, we next investigated whether FOXM1 regulates ATX expression at transcription level. -2 kb promoter region of human ATX gene was scanned to search potential FOXM1 DNA binding consensus sequence (5'-AT/CAAAT/CAA-3'), and two regions with putative FOXM1-binding sites were identified. Subsequently, we generated a series of luciferase reporter constructs containing wild-type (WT) and mutant (MUT) promoter region of ATX as presented in Figure 2A. The results of dual luciferase reporter assays indicated that FOXM1 knockdown decreased the promoter activity of WT-ATX, while no obvious effect was found on MUT-ATX promoter activity (Figure 2B). Concordantly, overexpression of FOXM1 significantly increased the WT-ATX but not MUT-ATX promoter activity in a dosage-dependent manner (Figure 2C), implying that FOXM1 regulates ATX expression via the putative FOXM1-binding sites. Furthermore, results of ChIP assay showed that the DNA fragments containing #1 site or
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#2 site sequence could be precipitated from PANC-1 cells with anti-FOXM1 antibody but not control IgG (Figure 2D, 2E), suggesting FOXM1 can directly bind to ATX promoter and activate ATX transcription in PDAC cells.

FOXM1 promotes PDAC cell proliferation and migration via ATX

To examine whether FOXM1 exerts its oncogenic function through ATX expression, CCK-8 assays were first employed to test cell growth in PDAC cells. The ATX knockdown was detected in PANC-1 cells by western blot analysis (Figure 3A). Results from CCK-8 assays revealed that ATX knockdown reduced PDAC cell growth compared with control cells. Meanwhile, ATX knockdown impaired the enhanced cell growth induced by FOXM1 overexpression in the two PDAC cell lines: PANC-1 and SW1990 (Figure 3B). Furthermore, we performed colony formation assays using PANC-1 cells and SW1990 cells stably overexpressing FOXM1, and consistently, promotion of colony formation by FOXM1 overexpression was also abolished by ATX knockdown (Figure 3C). On the other hand, cell migration results from transwell assays showed that FOXM1 promoted cell migration dependent on ATX expression in PANC-1 and SW1990 cells (Figure 3D). Therefore, these data above suggested that ATX may mediate the oncogenic role of FOXM1 in PDAC.
ATX stimulates FOXM1 expression via inhibiting the Hippo signaling in PDAC cells

Previous researches have clarified that FOXM1 is a downstream target gene of Hippo signaling pathway [21], and inhibition of Hippo pathway will induce dephosphorylation and nuclear localization of its main effector molecule Yes-Associated Protein (YAP), resulting in elevated FOXM1 transcription in tumor cells [22]. Interestingly, recent studies have shown that LPA, a primary downstream target gene of ATX, can suppress the Hippo/YAP signaling and thus mediates the up-regulation of FOXM1 mRNA and protein expression [23, 24]. Therefore, we assumed that ATX might regulate FOXM1 expression in PDAC cells through inhibiting the Hippo/YAP signaling. To validate our hypothesis, western blots and qPCR analyses were performed after ATX knockdown in PANC-1 cells. As shown in Figure 4A and 4B, ATX knockdown significantly decreased FOXM1 protein and mRNA expression in PANC-1 cells. Next, we investigated whether this effect could be reversed by XMU-MP-1, a selective Hippo pathway (MST1/2) inhibitor. Western blots analysis suggested that XMU-MP-1 treatment strongly reversed the ATX knockdown-mediated effects on FOXM1 expression (Figure 4C), which was validated by further qPCR analysis (Figure 4D). In addition, a dual luciferase reporter assay was employed to further explore the molecular mechanism via which ATX promoted FOXM1 expression. The results showed that FOXM1
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Promoter activity was significantly suppressed by knockdown of ATX, which was abolished by the treatment of XMU-MP-1 (Figure 4E), hinting to us that ATX mediated transcriptional activation of FOXM1 by inhibiting the activity of Hippo signaling pathway. These findings mentioned above demonstrated that ATX was critical for FOXM1 expression, and Hippo signaling plays important roles in this process.

Positive correlation of elevated expression of ATX and FOXM1 in PDAC

To verify the role of ATX in PDAC development and its relationship with FOXM1, a TMA was used to investigate ATX and FOXM1 protein expression by IHC analysis, and representative images were exhibited in Figure 5A. By statistical analysis, we found that both ATX (Table 1) and FOXM1 (Table 2) expression were significantly higher in tumor tissues as compared with adjacent normal tissues. Moreover, increased FOXM1 expression was positively correlated with elevated ATX expression in tumor tissues (R=0.66, P < 0.01, Figure 5B). These findings further confirmed the oncogenic role of ATX and its positive correlation with FOXM1 in PDAC.

Discussion

In the present study, we demonstrated for the first time that FOXM1 transcriptionally activated ATX, and ATX stimulated FOXM1 expression via inhibiting the Hippo/YAP signaling, thus forming a positive feedback loop and promoted PDAC cell proliferation and migration (Figure 5C).

ATX has been commonly recognized as an oncogene mainly by catalyzing the production of LPA, which enhances cell proliferation, migration and survival in different solid tumors such as prostate, breast, pancreatic and other...
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Previously, several researches have suggested ATX is related with the development of pancreatic cancer. For example, Yatomi et al. found that ATX level in the serum of patients with pancreatic cancer [10], and increased ATX expression was observed in a highly invasive PANC-1 cell line [11]. However, molecular mechanisms underlying aberrant activation of ATX/LPA signaling in cancers remains unclear yet. Herein, we reported that ATX is a downstream target gene of transcription factor FOXM1. Knockdown of FOXM1 significantly increased ATX expression whereas FOXM1 overexpression resulted in reduced ATX expression at both mRNA and protein levels. Furthermore, luciferase reporter assays and ChIP assays demonstrated that FOXM1 bound directly to specific regions of ATX promoter and positively regulated ATX promoter activity. Cell biological experiments suggested that the positive effects of FOXM1 on PDAC cell proliferation and migration abilities were dependent on the regulation of ATX expression. By IHC analysis, we confirmed that both ATX and FOXM1 were frequently up-regulated in PDAC tumor tissues when compared with adjacent normal tissues, and elevated ATX and FOXM1 expression were positively correlated with each other. Therefore, these findings strongly indicated that FOXM1 is a critical regulator of aberrant activation of ATX in PDAC.

As a key regulator of cell proliferation, invasion, metastasis, metabolism, angiogenesis and stem cell maintenance, FOXM1 has been identified as an oncogenic transcription factor in different human malignancies including PDAC [17, 18]. However, the molecular mechanisms underlying increased expression of FOXM1 and in PDAC still require further studies. Combined with previous studies that ATX/LPA signaling can inhibit the activity of the Hippo signaling pathway and that inhibition of the Hippo signaling promotes the expression and transcriptional activity of FOXM1 through increasing the nuclear localization of YAP [22-24], we assumed that ATX might influence FOXM1 expression in PDAC.
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cells by regulating the Hippo/YAP signaling. By molecular biological techniques we found that knockdown of ATX significantly reduced both mRNA and protein expression of FOXM1 in PDAC cells, which could be greatly reversed by treatment of a specific inhibitor of Hippo signaling pathway (MST1/2), XMU-MP-1. More importantly, results of dual luciferase reporter assays indicated that knockdown of ATX significantly inhibited the activity of FOXM1 promoter, which was strongly attenuated by treatment of XMU-MP-1. Hence, these data identified that transcriptional activation of ATX by FOXM1 increased the transcriptional activity of FOXM1 in turn via inhibiting the Hippo/YAP signaling, thus forming a positive feedback loop and promoted PDAC cell proliferation and migration.

Conclusions

In summary, the present study not only identified a positive feedback loop formed by ATX and FOXM1 and its important role in promoting PDAC cell proliferation and migration, but also indicated that inhibition of both ATX and FOXM1 might be a promising therapeutic strategy against human PDAC.

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Disclosure of conflict of interest

None.

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