miR-21 and KLF4 jointly augment epithelial-mesenchymal transition via the Akt/ERK1/2 pathway

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Abstract. miR-21 induces epithelial-mesenchymal transition (EMT) of human cholangiocarcinoma (CCA) cells. However, the mechanism by which this occurs remains unclear. In the present study, high throughput platform was employed to detect the genes that are differential expressed in QBC939 cells transfected with a hsa-miR-21 antagomir or control vectors. The EMT-related Krüppel-like factor 4 (KLF4) gene was downregulated after miR-21 was knocked down. Overexpression of miR-21 upregulated KLF4, Akt, ERK and mesenchymal cell markers (N-cadherin and vimentin), downregulated the expression of epithelial cell marker E-cadherin and reduced cell migration and invasion. Immunohistochemistry showed that KLF4, pAkt and pERK were upregulated in tumor xenografts transfected with miR-21 mimics. Inhibitors of the PI3K-Akt and ERK1/2 pathways, LY294002 and U0126, significantly suppressed the EMT phenotype. The present data demonstrated that overexpression of miR-21, accompanied with KLF4, augmented the EMT via inactivation of Akt and ERK1/2 pathways. In conclusion, we have identified a novel mechanism that may be targeted in an attempt to relieve the malignant biological behavior of CCA cells.

Introduction

Cancer metastasis is a highly coordinated and sequential process. An epithelial-mesenchymal transition (EMT) is important for disseminating cancer cells by endowing them with greater motility and invasiveness (1,2). Transformed cells downregulate epithelial cell marker E-cadherin whilst upregulating mesenchymal markers N-cadherin and vimentin (3-6).

MicroRNAs (miRNAs) are a class of small, endogenous non-coding RNAs that regulate target gene expression through pairing to its 3'-UTR (7). Accumulating evidence demonstrates that miRNAs are integral to the progression and metastasis of many human malignancies (8,9), including cholangiocarcinoma (CCA).

miR-21 is a major oncogenic miRNA, which is associated with several biomarkers and therapeutic targets in a variety of tumor types, such as renal (10), bladder (11), colorectal cancer (12) and others. miR-21 is an important critical regulator of the biological behavior of cancer cells and plays a role in apoptosis (10) and proliferation (13). Increasing evidence has demonstrated that miR-21 is implicated with induction and regulation of the EMT program (14-17). In addition, recent studies have suggested that miR-21 increases the proliferation, migration and invasion of cancer cells of various origin through activation of the phosphatidylinositol 3-kinase (PI3K)/protein kinase B (Akt) and mitogen-activated protein kinase (MAPK)/extracellular signal regulated kinase1/2 (ERK1/2) pathways (18,19), both of which are dysregulated during the maintenance of EMT (20-22). Notably, Yan et al (23) found that PI3K/AKT inhibition by miR-21 knockdown reversed EMT in breast cancer in vitro.

We previously demonstrated that miR-21 is involved in the EMT of CCA cells (24). Krüppel-like factor 4 (KLF4) has been considered to be critical for EMT (25), however, the interactions between miR-21 and KLF4 that trigger the EMT of CCA cells remain unclear. The present study aimed to identify if miR-21, jointly with KLF4, is involved in the maintenance of EMT of CCA cells via Akt and ERK1/2 pathways.

Materials and methods

Reagents and antibodies. Human hsa-miR-21 (MIMAT 0000076) antagomir, mimics, scramble control and negative control were purchased from Guangzhou Land Unicommed Biotechnology, Co., Ltd. (Guangzhou, China). PCR primers were synthesized by Sangon Biotech, Co., Ltd. (Shanghai, China). Real-time PCR assay kits were purchased from Takara Bio (Dalian, China). Specific siKLF4 duplexes and a scramble control siRNA sequence (SsiKLF4) were purchased from Guangzhou Land Unicommed Biotechnology. LY294002 and
U0126 were purchased from Sigma-Aldrich (St. Louis, MO, USA). Lipofectamine 2000 was purchased from Invitrogen (Carlsbad, CA, USA). Rabbit anti-human raised against E-cadherin, N-cadherin, vimentin, KLF4, Akt, phospho-Akt (p-Akt), ERK1/2, phospho-ERK1/2 (p-ERK1/2), GAPDH and an HRP-conjugated anti-rabbit IgG secondary antibody were purchased from Cell Signaling Technology (Danvers, MA, USA).

Cell lines and xenografts. Human QBC939 cholangiocarcinoma cells were purchased from the Cell Bank of the Chinese Academy of Sciences. Cells were maintained in high-glucose Dulbecco's modified Eagle's medium (DMEM; Gibco, Waltham, MA, USA), supplemented with 10% fetal bovine serum (FBS; Zhejiang Tianhang Biotechnology, Co. Ltd., Huzhou, China), 100 units/ml penicillin, 100 mg/ml streptomycin and 1% glutamine (Invitrogen) and were grown at 37˚C in a humidified atmosphere of 95% air and 5% CO2. Xenografts, procured from our previous study (24), were embedded within consecutive paraffin sections and sliced prior to immunohistochemical analysis.

Vectors and transfection. Specific siKLF-4 duplexes and a scramble control siRNA sequence (SstKLF-4) were designed, synthesized and annealed by Sigma-Aldrich. The selected RNA duplex (siKLF-4) corresponded to nucleotide sequences of KLF-4 mRNA sense: 5'-CCC ATC TTC ATC CAC GTT CG-3' and antisense: 5'-TGC TTG ATC TTG GGG CAC GT-3'. LY294002 (inhibitor of PI3K/Akt) and U0126 (inhibitor of ERK1/2) were purchased from Sigma-Aldrich. Human miR-21 mimics, inhibitor and negative control were purchased from Guangzhou RiboBio, Co., Ltd. (Guangzhou, China). QBC939 cells were transfected with either: i) miR-21 mimics (mimic group); ii) miR-21 negative control (control group); or ii) miR-21 inhibitor (inhibitor group) using Lipofectamine 2000 transfection reagent (Invitrogen) following the manufacturer's guidelines. Briefly, 5x104 cells were seeded into 6-well plates and grown to 60% confluence. Human miR-21 antagonist or its negative control were transfected directly into QBC939 cells at a final concentration of 50 nmol/l. Human miR-21 mimics or its negative control were allowed to form transfection complexes with Lipofectamin™ 2000 in Opti-MEMH I serum-free medium (Invitrogen) at a final concentration of 40 nmol/l. Transfections were performed in triplicate. The transfection efficiency was determined by fluorescent microscopy 24 h post transfection. Finally, the transfected cells were harvested for total RNA and protein extraction.

Real-time RT-PCR analysis. We investigated the expression of miR-21, KLF4 and ERK in transfected QBC939 CCA cells. Initially, total RNA was extracted using TRIzol reagent (Invitrogen) following the manufacturer's guidelines. Next, RNA was reverse transcribed into cDNA using a PrimeScripts RT reagent kit (Takara Bio). For the reverse transcription of miR-21, a miR-21 reverse transcription (RT) primer (5'-GTC GTA TCC AGT GCA GGG TCC GAG GTA TCC GCA CTG GAT ACG ACT CAA CA-3') and quantitative real-time RT-PCR (qPCR) primers (forward, 5'-GCC CGC TAG CTT ATC AGA CTG ATG-3' and reverse, 5'-GTC CAG GGT CCG AGG T-3') were used. qPCR reactions were carried out using SYBRH Premix Ex Taq™ II (Takara Bio) using an Applied Biosystems real-time PCR system (Life Technologies, Carlsbad, CA, USA). U6 small nuclear RNA was used as an internal control for miR-21. Glyceraldehyde-3-phosphate dehydrogenase (GAPDH) mRNA was used as an internal control for KLF4 and ERK genes. The following primers were used: miR-21: forward, 5'-ACA CTC CAG CGT GAG TTA TCA GAC TGA TG-3' and reverse, 5'-CTC AAC TGG TGT CGT GGA-3'; U6: forward, 5'-CTC CGT TCG GCA GCA CA-3' and reverse, 5'-AAC GCT TCA CGA ATT TGC GT-3'. PCR reactions were done in triplicate. All samples were normalized to internal controls and fold changes were calculated using the relative quantification method (2-ΔΔCT).

Western blotting. Western blotting of five proteins was conducted as described in our previous study (24). Briefly, cells and xenografts were washed in phosphate-buffered saline (PBS) and incubated in lysis buffer (Sangon Biotech, Co., Ltd., Shanghai, China). Protein was separated by 12% SDS-PAGE and transferred to a PVDF membrane (Sangon Biotech). Non-specific binding sites were blocked by incubation with TBST containing 5% (w/v) non-fat dried milk. Next, the membrane was incubated with rabbit anti-human Akt, p-Akt, KLF-4, ERK or p-ERK (1:1,000 dilution) primary antibodies at 37˚C for 1 h. Subsequently, membranes were incubated overnight with an HRP-conjugated anti-rabbit IgG secondary antibody (1:1,000 dilution) at 4˚C. Signals were visualized by an ECL chemiluminescence detection kit and semi-quantitated using ImageJ software. Equal protein loading was assessed by the expression of GAPDH.

Immunohistochemical staining. Tissues were obtained from mouse xenografts as previously described (24), and were fixed in formalin, embedded in paraffin and sectioned (2 μm). Slides were stained with KLF-4, p-Akt and p-ERK antibodies, and developed with a streptavidin-peroxidase (S-P) kit (Fuzhou Maixin Biotechnology Development, Co., Ltd., Fuzhou, China). The proportion of positive cells was determined using five x200 magnification fields per slide.

Migration and invasion assays. Migration and invasion of cells were determined with the Transwell chamber assay (8-μm pore size; Millipore, Billerica, MA, USA) completed as per the manufacturer's instructions. To determine invasion, the chamber was coated with 80 μl Matrigel (BD Biosciences, San Jose, CA, USA) hydrated with 50 μl serum-free medium. Next, 200 μl transfected cell suspension (1x105 cells/ml) was added to the upper chamber, and media supplemented with 30% fetal bovine serum (FBS) was added to the bottom wells. After culturing for 24 h, cells were fixed for 15 min in 4% formaldehyde and stained with 1% crystal violet. Cell numbers were counted under an optical microscope. Cell migration was also evaluated using the Transwell chamber assay with the exception that the chamber was not coated with Matrigel. Each experiment was repeated at least three times.

Construction and transfection of siKLF4. Based on the human KLF4 mRNA sequence (NM_000314), specific siRNA duplexes were designed, synthesized and annealed by Land Unicom Biotechnology. The selected RNA duplex (siKLF4)
corresponding to nucleotides 1567-1585 of KLF4 mRNA was defined as sense: 5’-GCU ACC UGU UAA AGA AUC AdTdT-3’ and antisense: 5’-UGA UUC UUU AAC AGG UAG CdTdT-3’. The scramble-control siRNA sequence (SsiKLF4) was also designed by Land Unicomed Biotechnology, and has no significant homology to any known human gene sequence. Prior to transfection, 5x10^5 cells were seeded into 6-well plates and grown to 70% confluence. Human siKLF4 or SsiKLF4 (final concentration of 50 nmol/l) was allowed to form transfection complexes with the Lipofectamine 2000 reagent (Invitrogen) in Opti-MEMI serum-free medium (Invitrogen) as per the manufacturer’s guidelines.

Scratch wound migration assay. For the scratch wound migration assay, 1x10^6 cells were seeded into 6-well plates and grown to 70% confluence. Next, the cells were transfected with a miR-21 mimic and co-incubated with LY294002 or U0126 inhibitors. A ‘wound’ was then created by scratching the cell cultures with a plastic filter tip. The cells were rinsed three times with PBS and incubated at 37˚C. The average distance that cells migrated after 0, 24 and 48 h was determined using an inverted microscope with x8 vision. Migration distance was calculated using image processing software. From the data, the migration rate = (1-distance at indicated time/distance at 0 h)%.$\,\text{Experiments were performed in triplicate.}$

Statistical analysis. All data are expressed as the mean ± SD. Data were analyzed using SPSS V20.0 software. The Student's t-test was used to determine statistical differences between treatment groups. A P≤0.05 was considered statistically significant.

Results

Transfected miR-21 and its inhibitor change the expression of miR-21 in QBC939 cells. To evaluate whether miR-21 mimics and inhibitors can modify the expression of miR-21, QBC939 cells were transfected with a miR-21 mimic or inhibitor for 48 h after which the relative expression of miR-21 was measured by RT-PCR. The relative expression of miR-21 was significantly upregulated in QBC939/miR-21 cells relative to NC cells (P<0.01; Fig. 1A). However, the relative expression of miR-21 was significantly downregulated in QBC939/miR-21 inhibitor cells relative to NC cells (P<0.01; Fig. 1A). These results suggested that hsa-miR-21 upregulates expression of miR-21 within QBC939 cells, whereas the hsa-miR-21 inhibitor could downregulate miR-21 in QBC939/miR-21 inhibitor cells.

Inhibition of miR-21 downregulates KLF4 in QBC939 cells. To investigate the role of miR-21 on EMT, QBC939 cells transfected with the hsa-miR-21 inhibitor or control differentially expressed genes and EMT-related genes were analyzed by Human GeneChip PrimeView. The data showed that 400 genes were differentially expressed (115 upregulated and 285 downregulated genes) in miR-21-knockdown cells compared to NC cells (Fig. 1B). These differentially expressed genes

Figure 1. Altered expression of miR-21, EMT-related genes, KLF4, Akt and ERK after transfection with miR-21 mimic, miR-21 inhibitor, or control. (A) QBC939 cells treated with miR-21 mimic showed increased expression of miR-21, whereas cells treated with the miR-21 inhibitor showed decreased expression of miR-21 compared to the NC group (P<0.01). (B) Clustered and (C) pathway enrichment of differentially expressed genes induced by miR-21 knockdown. (D and E) Altered levels of KLF4 and ERK by qPCR. The qPCR reactions were performed in triplicate. As loading control, 18S rRNA was used. (**P<0.01, compared with NC group).
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were mainly enriched for the glycolysis gluconeogenesis, pentose phosphate, fructose and mannose metabolism and p53 signaling pathways (Fig. 1C). Further analysis of these genes using IPA software showed that genes KLF4 and ERK are related to EMT. KLF4 and ERK expression were verified by qPCR (Fig. 1E).

miR-21 augments levels of KLF4, Akt and ERK in QBC939 cells. To investigate the mechanism of miR-21 on EMT in QBC939 cells, the relative mRNA expression of KLF4, Akt and ERK was determined. miR-21 increased mRNAs levels of KLF4, Akt and ERK relative to corresponding control cells (P<0.01; Fig. 1E). Conversely, the miR-21 inhibitor decreased KLF4, Akt and ERK levels (P<0.01). These results suggest that miR-21 affects mRNA expression of KLF4, Akt and ERK.

miR-21 increases expression of KLF4, and Akt and ERK activation in xenografts. To evaluate whether miR-21 upregulates the expression of KLF4 and of Akt and ERK activation, we used immunohistochemistry and western blotting to determine miR-21 protein levels in tissues obtained from mouse xenografts as previously described (24). As expected, the protein expression of KLF4 in the miR-21 mimic group was markedly upregulated compared to the control group (P<0.01; Fig. 2A, D and I). In the miR-21 mimic group, levels of pAkt and pERK were also significantly upregulated (P<0.01; Fig. 2A-C and E-G). These results demonstrate that miR-21 mimics upregulate the expression of KLF4 and activate Akt and ERK.

miR-21 antagonism accompanies KLF4 knockdown inactivated AKT and ERK1/2. Akt and ERK1/2 are two major components of signaling pathways that are involved in the regulation of cell proliferation, migration and survival. To determine the signaling molecules involved in the antagonism of KLF4 and miR-21-induced EMT phenotype, protein levels of phosphorylated Akt (p-Akt), Akt, phosphorylated ERK1/2 (p-ERK1/2) and ERK1/2 were determined by western blot analysis. The protein levels of p-Akt and p-ERK1/2 in QBC939/anti-KLF4/miR-21 inhibitor cells were markedly lower compared to levels in anti-KLF4 cells (P<0.01; Fig. 3A-C). miR-21 antagonism, in combination with KLF4 knockdown, could reverse EMT via suppressing Akt and ERK1/2 activation.

Overexpression of miR-21 induces an EMT phenotype accompanied with upregulation of KLF4 and activation of AKT and ERK1/2. To confirm the role of miR-21 in regulating an EMT phenotype, hsa-miR-21 mimic or inhibitor were transfected into QBC939 cells. The protein expression of EMT biomarkers (N-cadherin, vimentin and E-cadherin), KLF4, p-Akt, Akt, p-ERK1/2 and ERK1/2 were measured by western blot analysis. Compared to NC cells, overexpression of miR-21 increased the protein expression of N-cadherin and vimentin, but decreased the expression of E-cadherin (P<0.01; Fig. 3E and I). In addition, low expression of miR-21, induced by the miR-21 inhibitor, decreased the levels of N-cadherin and vimentin and increased the expression of E-cadherin (P<0.01; Fig. 3E and I). This suggests that overexpression of miR-21...
induces an EMT phenotype in QBC939 cells. Overexpression of miR-21 increased the expression of KLF4 (P<0.01; Fig. 3E and H). In addition, re-expression of miR-21 increased the expression of p-Akt and p-ERK1/2 in QBC939 cells (P<0.01; Fig. 3F and G), thereby suggesting that an overexpression of miR-21 could activate Akt and ERK1/2 pathways. Together, these data imply that miR-21 regulates EMT and is related to alterations in KLF4 expression and Akt/ERK1/2 pathways.
miR-21 regulates an EMT phenotype and increases invasion and migration through AKT and ERK1/2 pathways. To further investigate the involvement of Akt and ERK1/2 pathways in miR-21 regulating an EMT, QBC939 cells were transfected with miR-21 mimics for 72 h. Cells were treated for 24 h with either the LY294002 inhibitor of Akt or the U0126 inhibitor of ERK1/2. Our data confirm that treatment with LY294002 and U0126 resulted in abolished overexpression of miR-21, induced inhibition of E-cadherin, and activation of N-cadherin and vimentin (P<0.01; Fig. 4). Thus, we have demonstrated that miR-21 overexpression and its ability to induce the EMT phenotype were inhibited by LY294002 or U0126 in QBC939 cells.

We also studied the effect of antagonizing miR-21 on the migration and invasion of CCA cells (Fig. 5A, B and E). The number of migrated and invaded QBC939/anti-miR-21 cells were significantly lower compared to that of NC cells (P<0.01; Fig. 5C, D and F), thereby suggesting that miR-21 antagonists reduced the migration and invasion of CCA cells.

Discussion

Increasing evidence suggests that the expression of miR-21 is upregulated in various cancer types, including cholangiocarcinoma (CCA), where it is frequently associated with EMT (15,16,26). Overexpression of miR-21 in CCA biopsies is predictive for a poor outcome (24), and is accompanied with low expression of E-cadherin and high expression of N-cadherin and vimentin (27-29). In our previous study, we showed that miR-21 complicates the EMT of CCA cells. We also showed that CCA xenografts have high levels of N-cadherin and vimentin and low levels of E-cadherin (24). In the present study, we first identified genes that are differentially expressed in a CCA xenograft model transfected with miR-21 knockdown or control lentivirus, and found that KLF4 and ERK were significantly downregulated. Both KLF4 and ERK genes are associated with EMT. Tiwari et al (25) previously showed that KLF4 was significantly downregulated during EMT of breast cancer cells. However, the role of KLF4 in the EMT of CCA cells has not been elucidated. Accordingly, we studied the effect of miR-21 on the expression of KLF4, its role in the EMT, and investigated the potential underlying mechanisms of action. We used immunohistochemistry and western blot analysis to show that, in QBC939 CCA cells, overexpression of miR-21 promoted upregulation of KLF4 and expression of Akt and ERK proteins. Therefore, our data suggest that miR-21 and KLF4 regulate EMT through AKT/ERK1/2 pathways.

To investigate the roles of miR-21 and KLF4 in regulating the EMT phenotype of CCA cells, miR-21 mimics or inhibitors were transfected into QBC939 cells. In addition, KLF4 was silenced by using siKLF4. The data showed that the EMT phenotype was maintained by miR-21 mimics and reversed by miR-21 inhibitor. The results are consistent with previous studies, in which was shown that miR-21 facilitated EMT of pancreatic cancers (15,30), thyroid (16) and CCA origin and decreased activation of Akt and ERK1/2 pathways (24). We also demonstrated that the downregulation of KLF4 by using RNAi reversed the EMT and suppressed the activation of Akt and ERK1/2 signaling. Studies have demonstrated that Akt and/or ERK1/2 pathways are required for augmenting the EMT of several different cancer cell lines, including MCF-7 breast cancer cells (20), breast epithelial cells (21,22), cervical cancer cells (20), ovarian carcinoma cells (31) and colon...
cancer cells (32). These results demonstrate that miR-21 and KLF4 play an important role in mediating EMT of CCA cells through the AKT and ERK1/2 pathway.

Finally, we evaluated the invasion and migration of QBC939 cells. Overexpression of miR-21 by miR-21 mimics promoted the activation of Akt and ERK1/2 pathways and resulted in an increase in invasive ability and migration rate. However, these effects were suppressed by treatment of the cells with PI3K-Akt or the ERK1/2 inhibitor LY294002 or U0126.

In summary, our data demonstrate that miR-21 and KLF4 synergistically regulated EMT of QBC939 CCA cells through Akt and ERK1/2 pathways. The present study illustrated a mechanism by which miR-21 and KLF4 maintain an EMT within CCA cells, which may be beneficial for the development of novel therapies to treat cancer.

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