MicroRNA-206 inhibits the viability and migration of human lung adenocarcinoma cells partly by targeting MET

XI CHEN*, ZHONG-KAI TONG*, JIAN-YA ZHOU, YA-KE YAO, SHU-MENG ZHANG and JIAN-YING ZHOU

Department of Respiratory Diseases, Thoracic Disease Diagnosis and Treatment Center, The First Affiliated Hospital, School of Medicine, Zhejiang University, Hangzhou, Zhejiang 310003, P.R. China

Received March 31, 2015; Accepted May 13, 2016

DOI: 10.3892/ol.2016.4735

Abstract. MicroRNA (miRNA)-based targeting in cancer has emerged as a potential therapeutic strategy. miR-206 has recently been implicated in cancer. However, the role and molecular mechanism of miR-206 in lung adenocarcinoma are still unclear. The present study revealed that miR-206 was downregulated in human lung adenocarcinoma tissues. Overexpression of miR-206 in human lung adenocarcinoma-derived cells significantly inhibited cell viability and migration. Further experiments indicated that the overexpression of miR-206 decreased the expression of MET at the messenger RNA and protein levels via direct targeting of MET in a 3'-untranslated region-dependent manner. The knockdown of MET by small interfering RNA partly led to a phenocopy effect of miR-206. In conclusion, the present study identified miR-206 as a potential tumor suppressor of lung adenocarcinoma that exerts its functions, in part, by negative regulation of MET.

Introduction

Lung cancer is the leading cause of cancer-associated mortalities worldwide, and adenocarcinoma is one of the most common histological types of lung cancer in multiple countries (1). In the recent past, various major therapeutic advances have improved the prognosis of specific subgroups of patients (2). However, the 5-year survival rate of patients with lung cancer remains <15% (3), and therefore, novel therapeutic strategies are required for better management of this disease.

Materials and methods

Human tissue. Formalin-fixed and paraffin-embedded (FFPE) sections of paired lung adenocarcinoma tissues and adjacent normal tissues were obtained from patients who underwent radical resection for lung cancer. The samples were collected between October and December 2013 at The First Affiliated Hospital, School of Medicine, Zhejiang University (Zhejiang, China). The patients provided signed informed consent for participation in the study, which was approved by the Ethics Committee of Zhejiang University. The demographic and clinicopathological data are listed in Table I.
Table I. Clinical features of the 12 patients with lung adenocarcinoma.

<table>
<thead>
<tr>
<th>Patient no.</th>
<th>Gender</th>
<th>Age (years)</th>
<th>Histological type</th>
<th>Tobacco smoking history</th>
<th>pTNM stage</th>
<th>Tumor grade</th>
<th>EGFR mutation</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>M</td>
<td>47</td>
<td>Adenocarcinoma</td>
<td>Yes</td>
<td>T2aN0M0</td>
<td>IB</td>
<td>Exon 19 deletion</td>
</tr>
<tr>
<td>2</td>
<td>F</td>
<td>82</td>
<td>Adenocarcinoma</td>
<td>No</td>
<td>T2aN0M0</td>
<td>IB</td>
<td>Unknown</td>
</tr>
<tr>
<td>3</td>
<td>F</td>
<td>41</td>
<td>Adenocarcinoma</td>
<td>No</td>
<td>T2aN1M1</td>
<td>IV</td>
<td>Unknown</td>
</tr>
<tr>
<td>4</td>
<td>F</td>
<td>50</td>
<td>Adenocarcinoma</td>
<td>No</td>
<td>T2aN0M0</td>
<td>IB</td>
<td>Unknown</td>
</tr>
<tr>
<td>5</td>
<td>F</td>
<td>47</td>
<td>Adenocarcinoma</td>
<td>No</td>
<td>T2aN0M0</td>
<td>IB</td>
<td>Unknown</td>
</tr>
<tr>
<td>6</td>
<td>F</td>
<td>69</td>
<td>Adenocarcinoma</td>
<td>No</td>
<td>T1aN2M0</td>
<td>IIIA</td>
<td>Exon 19 deletion</td>
</tr>
<tr>
<td>7</td>
<td>M</td>
<td>46</td>
<td>Adenocarcinoma</td>
<td>No</td>
<td>T2bN1M0</td>
<td>IIB</td>
<td>Unknown</td>
</tr>
<tr>
<td>8</td>
<td>F</td>
<td>73</td>
<td>Adenocarcinoma</td>
<td>No</td>
<td>T2aN1M0</td>
<td>IIA</td>
<td>No mutation</td>
</tr>
<tr>
<td>9</td>
<td>F</td>
<td>50</td>
<td>Adenocarcinoma</td>
<td>No</td>
<td>T1bN0M0</td>
<td>IA</td>
<td>Unknown</td>
</tr>
<tr>
<td>10</td>
<td>F</td>
<td>51</td>
<td>Adenocarcinoma</td>
<td>No</td>
<td>T2bN3M0</td>
<td>IIIB</td>
<td>No mutation</td>
</tr>
<tr>
<td>11</td>
<td>M</td>
<td>37</td>
<td>Adenocarcinoma</td>
<td>No</td>
<td>T1bN0M0</td>
<td>IA</td>
<td>Unknown</td>
</tr>
<tr>
<td>12</td>
<td>M</td>
<td>43</td>
<td>Adenocarcinoma</td>
<td>No</td>
<td>T1bN2M0</td>
<td>IIIA</td>
<td>Unknown</td>
</tr>
</tbody>
</table>

M, male; F, female; TNM, tumor-node-metastasis; EGFR, epidermal growth factor receptor.

**Cell culture and reagents.** The HCC827 and A549 cell lines were purchased from the Committee on Type Culture Collection of the Chinese Academy of Sciences (Shanghai, China). The PC-9 cell line was a gift from Dr Caicun Zhou (Department of Oncology, Shanghai Pulmonary Hospital, Shanghai, China) and was verified by short tandem repeats analysis. HCC827 and A549 cells were cultured in RPMI-1640 (Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA) with 10% fetal bovine serum (FBS; Gibco; Thermo Fisher Scientific, Inc.) at 37°C and 5% CO₂. The PC-9 cells were maintained in Dulbecco’s modified Eagle’s medium (DMEM; Gibco; Thermo Fisher Scientific, Inc.) supplemented with 10% FBS.

**Oligonucleotides and transfection.** miR-206 mimics (double-stranded RNA oligonucleotides) and a negative control (NC) duplex that lacks significant homology to all known human sequences were purchased from Shanghai GenePharma Co., Ltd. (Shanghai, China). Small interfering RNA (siRNA) targeting MET (si-MET) and NC siRNA were purchased from Shanghai GenePharma Co., Ltd. The sequences are listed in Table II. The transfection was performed using Lipofectamine 2000 reagent (Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer’s protocol.

**Cell viability assay.** The day prior to transfection, 3,000 cells/well were plated in 96-well plates. Upon overnight incubation, the cells were transfected with miR-206 mimics or with the NC miRNA at 20 or 40-nM concentration. After the culture period, 10 µl cell counting solution (water-soluble tetrazolium-8; Dojindo Molecular Technologies, Inc., Kumamoto, Japan) was added to each well, followed by an additional incubation for 2 h at 24 h post-transfection, 1.5 h at 48 h post-transfection or 1 h at 72 h post-transfection. The absorbance of the solution was measured spectrophotometrically at 450 nm with an MRX II absorbance reader (Dynex Technologies, Chantilly, VA, USA). Each experiment was conducted three times in triplicate.

**Cell migration assays.** Transwell chambers (8-µm pore size; EMD Millipore, Billerica, MA, USA) were used to determine the migration of tumor cells. A total of 5x10⁴ cells were resuspended in 0.2 ml fresh medium without FBS 48 h after transfection, and then added to the inserts. Next, 0.6 ml culture medium with 10% FBS was added to the lower compartment. After 24 h, the non-migrating cells and culture medium were removed using cotton buds, while the cells on the lower surface of the inserts were fixed and stained with 0.1% crystal violet. Four random fields per insert were counted. The experiment was conducted three times in triplicate.

**Cell apoptosis assay.** Cell apoptosis was detected with an annexin V-fluorescein isothiocyanate (FITC) apoptosis detection kit (BD Biosciences, Franklin Lakes, NJ, USA). The cells were harvested 72 h after NC or miR-206 treatment. The cells were then washed twice with pre-chilled phosphate-buffered saline and resuspended in 500 µl binding buffer at a concentration of 1x10⁶ cells/ml. Then, cells were stained with 5 µl annexin V-FITC and 5 µl propidium iodide for 10 min in the dark. The apoptosis analysis was performed by the BD FACS Verse flow cytometer with CellQuest Pro 6.0 software (BD Biosciences).

**Reverse transcription-quantitative polymerase chain reaction (RT-qPCR).** Total RNA was extracted from the cultured cells with TRIzol reagent (Invitrogen; Thermo Fisher Scientific, Inc.). RNA was reverse transcribed into complementary DNA (cDNA) using PrimeScript™ RT reagent kit (Takara Biotechnology Co., Ltd., Dalian, China). miRNAs isolated from FFPE tissues were extracted using the RNeasy FFPE kit (Qiagen GmbH, Hilden, Germany). miRNAs that were isolated from the cultured cells were extracted with RNAiso Plus (Takara Biotechnology Co., Ltd.) and reverse transcribed into cDNA using an RT kit (Qiagen GmbH). PCR amplification and detection were performed on a 7500 Fast Real-Time PCR System (Applied Biosystems; Thermo Fisher Scientific, Inc.) using miScript SYBR Green PCR kit (Qiagen GmbH).
Table II. Sequences of the oligonucleotides used in the present study.

<table>
<thead>
<tr>
<th>Name</th>
<th>Sequence (5'-3')</th>
</tr>
</thead>
<tbody>
<tr>
<td>miR-206 mimics</td>
<td>UGGAAGUGUAAGGAAGUGUGUGG</td>
</tr>
<tr>
<td></td>
<td>ACACACUUCCCUUACAUUCAAUU</td>
</tr>
<tr>
<td>si-MET</td>
<td>GAGGGUGUUUGGAAAGAUATdT</td>
</tr>
<tr>
<td></td>
<td>UAUUUUCCCAACACCCUC4dT</td>
</tr>
<tr>
<td>NC</td>
<td>UUCUCGGAACGUACGGUTT</td>
</tr>
<tr>
<td></td>
<td>ACGUGACAGUUUGGAGAATT</td>
</tr>
<tr>
<td>GAPDH F</td>
<td>GGTCTCCTCTGACTTCAACA</td>
</tr>
<tr>
<td>GAPDH R</td>
<td>AGCAAAATTCGTGTCATAC</td>
</tr>
<tr>
<td>MET F</td>
<td>GTTTGTCCACAGAGACTTGGCTG</td>
</tr>
<tr>
<td>MET R</td>
<td>ATCCACTTCACTGGCAGCTTTG</td>
</tr>
</tbody>
</table>

F, forward; R, reverse; miR, microRNA; si, small interfering; GAPDH, glyceraldehyde 3-phosphate dehydrogenase; NC, negative control.

The amplification protocol was as follows: Initial denaturation at 95°C for 30 sec, and then 40 cycles of denaturation at 95°C for 5 sec, annealing and extension at 60°C for 34 sec. The relative amounts of mRNA and miRNA gene transcripts were normalized to the levels of glyceraldehyde-3-phosphate dehydrogenase (GAPDH) and RNA, U6 small nuclear 2 (RNU6-2). The sequences of the primers for GAPDH and MET are listed in Table II. The primers for RNU6-2 and miR-206 were purchased from Qiagen Gmbh.

Vector construction and dual-luciferase reporter assay. The 3' untranslated regions (UTRs) and the mutant 3'-UTR of MET, which contained putative binding sites for miR-206, were cloned downstream of the pmirGLO Dual-Luciferase miRNA Target Expression vector (Promega Corporation, Madison, WI, USA). The cells were plated in 24-well plates and were transfected with the constructed luciferase vector (100 ng) and 40 nM miR-206 or NC. The relative luciferase activity was measured 48 h after transfection as described previously (25).

Immunoblotting. Following the various treatments, the cells were harvested, lysed and quantified. Equivalent quantities (50 µg) of protein were separated by 8% sodium dodecyl sulfate-polyacrylamide gel electrophoresis and transferred onto polyvinylidene difluoride membranes. Membranes were blocked for 1 h with 5% non-fat milk and then incubated overnight with a primary rabbit polyclonal antibody against MET (1:1,000 dilution; catalog no., 4560; Cell Signaling Technology, Inc., Danvers, MA, USA) and a rabbit polyclonal antibody against β-actin (1:2,000 dilution; catalog no., sc-130656; Santa Cruz Biotechnology, Inc., Dallas, TX, USA). Next, the membranes were incubated with a chicken anti-rabbit horseradish peroxidase-conjugated secondary antibody (catalog no., sc-516087, Santa Cruz Biotechnology, Inc.) at a 1:2,000 dilution for 2 h. Immunoreactive bands were detected with an enhanced chemiluminescence system (Pierce; Thermo Fisher Scientific, Inc.) as described previously (23).

Statistical analysis. All data are presented as the mean ± standard deviation of three independent experiments. All statistical analyses were performed using SPSS 16.0 statistical software (SPSS, Inc., Chicago, IL, USA). P<0.05 was considered to indicate a statistically significant difference according to Student's t-test.

Results

miR-206 was frequently downregulated in lung adenocarcinoma. In the present study, the expression levels of miR-206 were detected in 12 pairs of human lung adenocarcinoma tissues and adjacent normal tissues. In total, 8 of 12 (75%) lung adenocarcinoma tissues displayed a decreased expression of miR-206 compared with adjacent normal tissues (Fig. 1). This indicated that miR-206 is frequently downregulated in human lung adenocarcinoma.

Overexpression of miR-206 inhibited the viability of lung adenocarcinoma cells in vitro. In order to determine if miR-206 could function as a tumor suppressor in human lung adenocarcinoma cells, synthetic miR-206 mimics and NC miRNA were independently transfected into A549, PC-9 and HCC827 cells. The overexpression of miR-206 exerted a potent inhibitory effect in a time- and dose-dependent manner. Compared with the NC group, miR-206 caused an increased percentage of inhibition of cell viability at a concentration of 40 nM in the three cell lines tested at 48 or 72 h after transfection (Fig. 2). These data suggested that miR-206 negatively modulates the viability of lung adenocarcinoma cells. The apoptosis analysis by flow cytometry revealed that the early apoptosis rate was not affected in A549 or HCC827 cells, whereas the late apoptosis rate was higher in HCC827 cells following miR-206 treatment (Fig. 3).

Forced expression of miR-206 repressed the migration of lung adenocarcinoma cells. To better understand the function of miR-206 in lung adenocarcinoma, the present study tested...
whether miR-206 could affect the migration of lung adenocarcinoma cells. A549, PC-9 and HCC827 cells were transfected with different concentrations of negative control or miR-206 mimics for 24, 48 or 72 h. Cell viability was determined using the water-soluble tetrazolium-8 assay. Each experiment was performed three times in triplicate. The results are presented as the mean ± standard deviation (˚P<0.05, **P<0.01). miR, microRNA; NC, negative control.

**Figure 2.** Overexpression of miR-206 inhibited the viability of lung adenocarcinoma cells *in vitro*. A549, PC-9 and HCC827 cells were transfected with different concentrations of negative control or miR-206 mimics for 24, 48 or 72 h. Cell viability was determined using the water-soluble tetrazolium-8 assay. Each experiment was performed three times in triplicate. The results are presented as the mean ± standard deviation (˚P<0.05, **P<0.01). miR, microRNA; NC, negative control.

**Figure 3.** Early apoptosis was not induced by overexpression of miR-206 in A549 or HCC827 cells. A549 and HCC827 cells were transfected with 40 nM negative control or miR-206 mimics for 48 h. Cells were subjected to flow cytometry for cell apoptosis analysis. The results are presented as the mean ± standard deviation (˚P<0.05). miR, microRNA; NC, negative control; A, annexin V; FITC, fluorescein isothiocyanate; Q, quadrant; Comp, composition.

miR-206-binding sites, was previously demonstrated to be a specific target of miR-206 in rhabdomyosarcoma (24). In the present study, it was observed that MET was downregulated significantly at both the mRNA and protein levels (Fig. 5A and B). A luciferase reporter assay confirmed that MET was a direct target of miR-206 in human lung adenocarcinoma cells (Fig. 5C and D). These results suggested a targeted downregulation of MET by miR-206 in lung adenocarcinoma.

**Figure 3.** Early apoptosis was not induced by overexpression of miR-206 in A549 or HCC827 cells. A549 and HCC827 cells were transfected with 40 nM negative control or miR-206 mimics for 48 h. Cells were subjected to flow cytometry for cell apoptosis analysis. The results are presented as the mean ± standard deviation (˚P<0.05). miR, microRNA; NC, negative control; A, annexin V; FITC, fluorescein isothiocyanate; Q, quadrant; Comp, composition.

**Overexpression of miR-206 downregulated the expression of MET in lung adenocarcinoma cells through targeting its 3’-UTR.** The mRNA of MET, whose 3’-UTR contains miR-206-binding sites, was previously demonstrated to be a specific target of miR-206 in rhabdomyosarcoma (24). In the present study, it was observed that MET was downregulated significantly at both the mRNA and protein levels (Fig. 5A and B). A luciferase reporter assay confirmed that MET was a direct target of miR-206 in human lung adenocarcinoma cells (Fig. 5C and D). These results suggested a targeted downregulation of MET by miR-206 in lung adenocarcinoma.

miR-206 triggered the inhibition of cell viability and migration in lung adenocarcinoma cells in part via targeting MET. To determine whether the downregulation of MET was involved in the miR-206-mediated inhibition of cell viability and migration, the physiological function of MET was evaluated by an RNA interference approach. The knockdown of MET suppressed cell viability and migration, but these effects were not as remarkable as those observed upon miR-206 overexpression (Fig. 6). These findings indicated that the anti-tumor effect of miR-206 is partly dependent on the targeted regulation of MET.
Discussion

In recent years, the idea of targeted molecular therapy has become one of the most attractive topics in the field of human cancer treatment (3). Thus, the identification of the molecular pathogenesis of cancer is crucial for the development of effective therapeutic approaches. Emerging evidence suggests that the dysregulation of miRNAs plays a pivotal role in...
the pathogenesis of human cancer development through the targeting of a variety of important molecules (25). Previous studies have demonstrated that miR-206 is downregulated in lung cancer, which is correlated with tumor invasion and metastasis (20). However, the functional role of miR-206 in lung adenocarcinoma is largely unknown. In the present study, the functional characterization and the mechanism of action of miR-206 in human lung adenocarcinoma were investigated. It was observed that tumor tissues from patients with lung adenocarcinoma exhibited a decrease in miR-206 expression. Furthermore, miR-206 expression caused significant suppression of cell viability and migration in lung adenocarcinoma cells in vitro. These results indicate that miR-206 may serve as a tumor-suppressor gene in lung adenocarcinoma.

MET is a receptor tyrosine kinase that is dysregulated in multiple cancer types, including lung cancer (26). Overexpression of MET protein is detected in 36.0-72.3% of lung adenocarcinoma tissues, and is associated with tumor metastasis and poor prognosis (27-29). Silencing or inactivation of MET has been demonstrated to be important for cell viability in vitro, as well as for cell motility and migration (11,30). In rhabdomyosarcoma, MET was demonstrated to be a direct target of miR-206 (15). In the present study, MET was downregulated by miR-206 in lung adenocarcinoma cells at both the mRNA and protein levels. The luciferase assay indicated that MET is a target of miR-206, as the specific knockdown of MET by siRNA led to a partial phenocopy of the inhibitory effect of cell viability and migration inhibition that was observed with miR-206 overexpression. Together with the present findings, these results suggested that MET acts as a potential oncogene and is involved in the anti-tumor effects of miR-206.

Figure 6. Knockdown of MET reduced the cell viability and migration of lung adenocarcinoma cells. (A) A549, PC-9 and HCC827 cells were transfected with 40 nM NC or si-MET for 48 h. The cell lysates were harvested, and the phosphorylation of the indicated proteins was determined by western blotting. Three independent experiments were performed. (B) A549, PC-9 and HCC827 cells were transfected with 40 nM NC or si-MET for 48 h. Cell viability was determined using water-soluble tetrazolium-8 assays. Each experiment was performed three times in triplicate. The results are presented as the mean ± SD (P<0.05, **P<0.01). (C) Quantification from three independent experiments and (D) representative micrographs of the cell migration assay (magnification, x100). The results are presented as the mean ± SD (P<0.05, **P<0.01). si, small interfering; NC, negative control; SD, standard deviation.
In conclusion, the present study confirms that miR-206 is frequently downregulated in lung adenocarcinoma. The ectopic expression of miR-206 can exert an inhibitory effect on cell viability and migration of lung adenocarcinoma cells, in part by targeting MET. These data suggest that miR-206 serves as a potential tumor suppressor and may be developed as a novel therapeutic strategy for patients with lung adenocarcinoma.

Acknowledgements

The present study was supported by grants from the National Natural Science Foundation of China (Beijing, China; grant no. 81101768), the State Scholarship Fund of China (Beijing, China; grant no. 201308330145) and the Medicine and Health Sci-Tech Project of Zhejiang Province (Hangzhou, China; grant no. 2015KYA074).

References