**MicroRNA-454 inhibits the malignant biological behaviours of gastric cancer cells by directly targeting mitogen-activated protein kinase 1**

XIAODONG WANG¹, BAICHUN LIU¹, FUXING WEN² and YING SONG¹

¹Department of Digestive Endoscopy, The Second Hospital of Jilin University, Changchun, Jilin 130041; ²Department of General Medicine, The General Hospital of China National Petroleum Corporation in Jilin, Jilin 132021, P.R. China

Received July 13, 2017; Accepted December 12, 2017

DOI: 10.3892/or.2017.6171

**Abstract.** Gastric cancer (GC) is the fifth most commonly diagnosed malignant disease and the third leading cause of cancer-related deaths worldwide. Recently, numerous microRNAs (miRNAs) have been determined to contribute to GC initiation and progression, suggesting that miRNAs may be developed as effective diagnostic and prognostic molecular biomarkers and can be investigated as therapeutic targets for patients with this disease. Therefore, further investigation of the miRNAs involved in GC development represents an opportunity to improve the prognosis of GC patients. miRNA-454 (miR-454) is abnormally expressed in multiple types of human cancer. However, the expression pattern, biological roles and underlying mechanism of miR-454 in GC remain unclear and require further investigation. In the present study, we assessed miR-454 expression in GC tissues and cell lines. We also explored the effects of miR-454 on the biological behaviours of tumour cells and the underlying molecular mechanisms of miR-454. The results revealed that miR-454 was significantly downregulated in GC tissues and cell lines. Low miR-454 expression was positively associated with lymph node metastasis, invasive depth and TNM stage. Additionally, upregulation of miR-454 inhibited cell proliferation and invasion and induced the apoptosis of the GC cells. Subsequently, mitogen-activated protein kinase 1 (MAPK1) was identified as a direct target of miR-454. MAPK1 was upregulated in GC tissues and was found to be negatively correlated with the miR-454 expression level. Downregulation of MAPK1 also suppressed GC cell proliferation and invasion and increased apoptosis, thereby resembling the suppressive effects of miR-454 overexpression in GC. Moreover, upregulation of MAPK1 reversed the tumour-suppressive effects of miR-454 in GC. Collectively, our data demonstrated that miR-454 may play tumour-suppressing roles in GC through the regulation of MAPK1, suggesting that miR-454 may be a novel biomarker and therapeutic target for patients with this disease.

**Introduction**

Gastric cancer (GC) is the fifth most commonly diagnosed malignant disease and the third leading cause of cancer-related deaths worldwide (1). In China, GC is the most common type of cancer, and an estimated 679,100 new cases and 498,000 deaths occur each year (2). *Helicobacter pylori* infection, Epstein-Barr virus infection, poor diet, tobacco use and obesity are some of the identified risk factors that contribute to GC initiation and progression (3-5). Currently, the predominant treatments for patients with GC are surgery, chemotherapy, radiotherapy, immunotherapy and targeted therapy (6). Despite tremendous advances in the treatment strategies for patients with GC in recent years, the prognosis for advanced GC patients remains poor, and the 5-year survival rate for GC patients remains at approximately 4-5% (7). The poor prognosis of GC patients is mainly attributed to invasion and metastasis, which are the major causes of GC-related recurrence and death (8,9). Therefore, the molecular mechanisms underlying GC development and progression must be elucidated to develop novel biomarkers for the diagnosis and effective treatments for patients with this disease.

MicroRNAs (miRNAs) are a novel group of endogenous, single-strand, noncoding and short RNA molecules that are 20-25 nucleotides in length (10). miRNAs exert negative genetic regulation by directly binding to the 3'-untranslated regions (3'-UTRs) of their target genes in a base-pairing manner and subsequently inducing translational silencing or degradation of mRNAs (11). Substantial evidence indicates that miRNAs play important roles in most biological processes in both normal development and in various diseases, including cancer (12,13). In recent years, miRNA dysregulation has been reported in various types of human cancers, such as GC (14), bladder cancer (15), colorectal cancer (16), glioma (17) and...
lung cancer (18). Furthermore, aberrantly expressed miRNAs are associated with tumourigenesis and tumour development and affect cell proliferation, cell cycle distribution, apoptosis, migration, invasion, angiogenesis and epithelial-mesenchymal transition (19-21). Cancer-associated miRNAs may function as oncogenes or tumour-suppressor genes depending on the roles of their target genes and tumour types (22). Thus, miRNAs can serve as diagnostic markers or therapeutic targets for anticancer therapy.

miR-454 is abnormally upregulated in multiple types of human cancer including uveal melanoma (23) and colorectal cancer (24), whereas it is weakly expressed in glioblastoma (25) and osteosarcoma (26). However, the expression pattern, biological roles and underlying mechanism of miR-454 in GC remain unclear and require further investigation. In the present study, we examined miR-454 expression in GC tissues and cell lines. We also explored the effects of miR-454 on the biological behaviours of tumour cells and the underlying molecular mechanisms of miR-454.

Materials and methods

Human tissue samples. This study was approved by the Ethics Committee of The Second Hospital of Jilin University. Written informed consent was obtained from each patient who participated in this research. A total of 45 paired GC tissues (adenocarcinoma) and matched adjacent non-tumourous tissues were obtained from patients who were treated with surgical resection at The Second Hospital of Jilin University between June 2014 and March 2016. The matched adjacent non-tumourous tissues were obtained 5 cm away from the tumour tissues. In addition, histological examination was also performed to confirm the non-tumourous tissues. No patient had undergone any pre-operative treatment, such as radiotherapy, chemotherapy, immunotherapy or targeted therapy. All these tissues were immediately frozen in liquid nitrogen and stored in a refrigerator at -80°C until further use. Clinical features of the GC patients were collected and are summarised in Table I.

Cell lines, culture conditions and transfection. Human GC cell lines (MGC-803, MKN-1, SGC-7901, BGC-823 and AGS) and the normal gastric epithelium GES-1 cell line were acquired from the Shanghai Institute of Biochemistry and Cell Biology (Shanghai, China). All the cell lines were cultured in Dulbecco's modified Eagle's medium (DMEM) supplemented with 10% foetal bovine serum (FBS), 100 U/ml of penicillin and 100 mg/ml of streptomycin (Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA) in accordance with the manufacturer's protocol. The total RNA concentration was examined using a NanoDrop ND-100 Spectrophotometer (NanoDrop Technologies, Wilmington, DE, USA). For the detection of miR-454 expression, total RNA was reverse-transcribed into cDNA using a TaqMan® MicroRNA Reverse Transcription kit (Applied Biosystems; Thermo Fisher Scientific, Inc.), and endogenous U6 snRNA was used as a control. For the quantification of MAPK1 expression, cDNA was synthesised using PrimeScript™ RT reagent kit (Takara Biotechnology Co., Ltd., Dalian China), followed by qPCR with SYBR Premix Ex Taq Master Mix (Takara Biotechnology Co., Ltd.). GAPDH was used as an internal control for MAPK1 mRNA expression. All experiments were performed in triplicate, and data were analysed with the 2^(-ΔΔCt) method (27).

RNA extraction and reverse transcription-quantitative polymerase chain reaction (RT-qPCR). Total RNA was isolated from tissue samples or cells using TRIzol reagent (Invitrogen; Thermo Fisher Scientific, Inc.) according to the manufacturer's protocol. The total RNA concentration was examined using a NanoDrop ND-100 Spectrophotometer (NanoDrop Technologies, Wilmington, DE, USA). For the detection of miR-454 expression, total RNA was reverse-transcribed into cDNA using a TaqMan® MicroRNA Reverse Transcription kit (Applied Biosystems; Thermo Fisher Scientific, Inc.). Quantitative polymerase chain reaction (qPCR) was performed with TaqMan MicroRNA Assay kit (Applied Biosystems; Thermo Fisher Scientific, Inc.), and endogenous U6 snRNA was used as a control. For the quantification of MAPK1 expression, cDNA was synthesised using PrimeScript™ RT reagent kit (Takara Biotechnology Co., Ltd., Dalian China), followed by qPCR with SYBR Premix Ex Taq Master Mix (Takara Biotechnology Co., Ltd.). GAPDH was used as an internal control for MAPK1 mRNA expression. All experiments were performed in triplicate, and data were analysed with the 2^(-ΔΔCt) method (27).

Table I. Association between miR-454 expression and clinicopathological features of the gastric cancer patients (n=45).

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Total</th>
<th>Low</th>
<th>High</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;60</td>
<td>18</td>
<td>10</td>
<td>8</td>
<td>0.626</td>
</tr>
<tr>
<td>≥60</td>
<td>27</td>
<td>13</td>
<td>14</td>
<td></td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>35</td>
<td>17</td>
<td>18</td>
<td>0.524</td>
</tr>
<tr>
<td>Female</td>
<td>12</td>
<td>6</td>
<td>4</td>
<td></td>
</tr>
<tr>
<td>Tumor size (cm)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;5</td>
<td>24</td>
<td>12</td>
<td>12</td>
<td>0.873</td>
</tr>
<tr>
<td>≥5</td>
<td>21</td>
<td>11</td>
<td>10</td>
<td></td>
</tr>
<tr>
<td>Differentiation</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Well and moderate</td>
<td>20</td>
<td>9</td>
<td>11</td>
<td>0.463</td>
</tr>
<tr>
<td>Poor and signet</td>
<td>25</td>
<td>14</td>
<td>11</td>
<td></td>
</tr>
<tr>
<td>Lymph node metastasis</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>No</td>
<td>21</td>
<td>7</td>
<td>14</td>
<td>0.026</td>
</tr>
<tr>
<td>Yes</td>
<td>24</td>
<td>16</td>
<td>8</td>
<td></td>
</tr>
<tr>
<td>Invasive depth</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>T1+T2</td>
<td>19</td>
<td>4</td>
<td>15</td>
<td>0.001</td>
</tr>
<tr>
<td>T3+T4</td>
<td>26</td>
<td>19</td>
<td>7</td>
<td></td>
</tr>
<tr>
<td>TNM stage</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>I-II</td>
<td>19</td>
<td>6</td>
<td>13</td>
<td>0.025</td>
</tr>
<tr>
<td>III-IV</td>
<td>26</td>
<td>17</td>
<td>9</td>
<td></td>
</tr>
</tbody>
</table>
Cell Counting Kit-8 (CCK-8) assay. Transfected cells were collected at 24 h post-transfection and seeded into 96-well plates at a density of 3x10^3 cells per well. After culturing for 0, 24, 48 and 72 h, CCK-8 assay was performed according to the manufacturer's instructions. In brief, 10 µl of CCK-8 reagent (Beyotime, Shanghai, China) was added to each well. The cells were incubated at 37°C with 5% CO2 for 2 h. Absorbance was determined at a wavelength of 450 nm using an ELX808 absorbance reader (BioTek Instruments, Inc., Winooski, VT, USA). Each assay was performed in triplicate and repeated three times.

Transwell invasion assay. Transwell invasion assay was performed using Transwell plates (8 µm pores; Corning Inc., Corning, NY, USA) coated with Matrigel (BD Biosciences, Franklin Lakes, NJ, USA). Briefly, 5x10^4 transfected cells in FBS-free DMEM were added into the upper compartment of the chamber. The lower chamber was filled with DMEM containing 20% FBS to serve as a chemoattractant. After being incubated at 37°C with 5% CO2 for 24 h, the cells, which remained on the upper surface of the membrane, were carefully removed with cotton swabs. The invasive cells were fixed in methanol for 10 min, stained with 0.5% crystal violet solution for 10 min, washed in PBS and air-dried. Finally, images of five randomly selected fields of the invasive cells were obtained and counted under an inverted microscope (IX83; Olympus Corporation, Tokyo, Japan) at the magnification of x200.

Flow cytometric assay. An Annexin V-fluorescein isothiocyanate (FITC)/propidium iodide (PI) staining kit (BD Biosciences) was utilised to determine the cell apoptotic rate. Cells were collected and washed in ice-cold PBS after 48 h of transfection. The cells were stained with 5 µl of Annexin V-FITC and 5 µl of PI for 15 min at room temperature in the dark after their resuspension in 1X binding buffer. Flow cytometry (Beckman Coulter, Inc., Brea, CA, USA) was then performed to determine the apoptosis rate. CellQuest® software (version 3.3; Becton Dickinson, Heidelberg, Germany) was used to analyse the data. The tests were performed in triplicate and repeated three times.

Target prediction. The potential targets of miR-454 were predicted using TargetScan (www.targetscan.org) and microRNA (www.microrna.org).

Luciferase reporter assay. Bioinformatic analysis predicted that MAPK1 is a candidate target of miR-454. Luciferase reporter assays were performed to determine whether MAPK1 is a direct target of miR-454. Luciferase reporter plasmids, psiCHECK2-MAPK1-3'UTR wild-type (Wt) and psiCHECK2-MAPK1-3'UTR mutant (Mut), were synthesised and confirmed by Shanghai GenePharma Co., Ltd. (Shanghai, China). The cells were seeded into 24-well plates at 50-60% confluence. After incubation overnight, the plasmid (psiCHECK2-MAPK1-3'UTR Wt or psiCHECK2-MAPK1-3'UTR Mut) together with miR-454 mimics or miR-NC were transfected into cells using Lipofectamine 2000 according to the manufacturer's protocol. After 48 h of incubation at 37°C with 5% CO2, the transfected cells were collected and analysed for luciferase activity using the Dual-Luciferase Reporter Assay System (Promega Corporation, Manheim, Germany). The firefly luciferase activity was normalised to Renilla luciferase activity.

Western blot analysis. Total protein was extracted from the tissue samples or cells using radioimmunoprecipitation assay lysis buffer (Beyotime Institute of Biotechnology, Haimen, China). Subsequent to the quantification of total protein concentration with a BCA assay kit (Beyotime Institute of Biotechnology), equal quantities of protein were separated by 10% SDS-PAGE and transferred to polyvinylidene fluoride membranes (EMD Millipore, Billerica, MA, USA). The membranes were blocked with 5% skimmed dry milk in TBST at room temperature for 2 h and incubated overnight at 4°C with primary antibodies: mouse anti-human monoclonal MAPK1 (sc-136288; 1:1,000 dilution; Santa Cruz Biotechnology, Inc., Santa Cruz, CA, USA) and mouse anti-human monoclonal GAPDH (sc-32233; 1:1,000 dilution; Santa Cruz Biotechnology, Inc.). The membranes were washed three times with TBST before they were incubated with goat anti-mouse horseradish-peroxidase-conjugated secondary antibody (sc-2005; 1:5,000 dilution; Santa Cruz Biotechnology, Inc.) at
Figure 2. miR-454 inhibits GC cell proliferation, invasion and increases apoptosis in vitro. (A) The expression of miR-454 was determined by RT-qPCR in MGC-803 and AGS cells 48 h after transfection with miR-454 mimics or miR-NC. *P<0.05 compared with miR-NC. (B) CCK-8 assay was performed in MGC-803 and AGS cells after transfection with miR-454 mimics or miR-NC. *P<0.05 compared with miR-NC. (C) Transwell invasion assay was utilised to evaluate the effect of miR-454 overexpression on the invasion ability in the MGC-803 and AGS cells. *P<0.05 compared with miR-NC. (D) The effect of miR-454 on the apoptosis of MGC-803 and AGS cells was examined with flow cytometry analysis. *P<0.05 compared with miR-NC. GC, gastric cancer; RT-qPCR, reverse transcription-quantitative polymerase chain reaction; CCK-8, Cell Counting Kit-8.
room temperature for 1 h. Protein bands were visualised using an enhanced chemiluminescence solution (Pierce, Rockford, IL, USA). Densitometry was quantified using Image J software version 1.41 (National Institutes of Health, Bethesda, MD, USA). GAPDH served as an internal control.

Statistical analysis. Data are presented as mean ± standard errors. All statistical analyses were performed with Student's t-tests or one-way analysis of variance plus multiple comparisons on SPSS 18.0 (SPSS, Inc., Chicago, IL, USA). Student-Newman-Keuls test was used as a post-hoc test following ANOVA. Spearman's correlation analysis was used to analyze the association between miR-454 and MAPK1 mRNA levels in GC tissues. The categorical data in this research were miR-454 group vs. miR-NC group; MAPK1 siRNA vs. NC siRNA group; miR-NC, miR-454 mimics+pcDNA3.1, miR-454 mimics+pcDNA3.1-MAPK1 groups. P<0.05 was considered to be statistically significant.

Results

miR-454 is downregulated in both GC tissues and cell lines.

In order to evaluate the role of miR-454 in GC development, miR-454 expression was determined in 45 paired GC tissues and matched adjacent non-tumourous tissues by RT-qPCR. The results showed that the GC tissues had a significantly lower expression level of miR-454 compared with adjacent non-tumourous tissues (Fig. 1A; P<0.05). To clarify the clinical significance of miR-454 in GC, all patients were divided into an miR-454 low expression group (n=23) and miR-454 high expression group (n=22) on the basis of the median miR-454 expression. Table I shows that a low miR-454 expression level was positively associated with lymph node metastasis (P=0.026), invasive depth (P=0.001) and TNM stage (P=0.025). However, miR-454 expression was not significantly associated with age (P=0.626), sex (P=0.524), differentiation (P=0.463) and tumour size (P=0.873).

We detected miR-454 expression in five GC cell lines (MGC-803, MKN-1, SGC-7901, BGC-823 and AGS) and in normal gastric epithelium GES-1 cell line to validate whether aberrant downregulation of miR-454 also occurs in GC cell lines. RT-qPCR analysis data revealed that miR-454 was aberrantly downregulated in all GC cell lines compared with its expression in normal GES-1 cell line (Fig. 1B, P<0.05). These results suggest that miR-454 plays important roles in GC occurrence and development.

miR-454 inhibits GC cell proliferation and invasion and promotes apoptosis in vitro.

MGC-803 and AGS cell lines, which expressed the lowest levels of miR-454 among the five...
GC cell lines tested, were selected for further experiments. Both cell lines were transfected with miR-454 mimics or miR-NC to investigate the biological functions of miR-454 in GC. We examined the expression levels of miR-454 in the MGC-803 and AGS cells by RT-qPCR 48 h after transfection to assess the efficiency of the miR-454 transfection. Fig. 2A shows that miR-454 was markedly upregulated in the MGC-803 and AGS cells after transfection with miR-454 mimics (Fig. 2A; P<0.05). CCK-8 assays were then performed on the MGC-803 and AGS cells, which were transfected with miR-454 mimics or miR-NC, to evaluate the effect of miR-454 in GC cell proliferation. The results indicated that the upregulation of miR-454 decreased the proliferation of MGC-803 and AGS cells when compared with the miR-NC group (Fig. 2B; P<0.05). Transwell invasion assays were performed to explore the effect of miR-454 on GC cell metastasis. Fig. 2C shows that restoration expression of miR-454 decreased the cell invasion abilities of the MGC-803 and AGS cells (P<0.05). Moreover, we studied whether miR-454 has any effect on GC cell apoptosis. The results of flow cytometry assay indicated that miR-454 overexpression promoted MGC-803 and AGS cell apoptosis (Fig. 2D; P<0.05). These results suggested that miR-454 may play tumour-suppressing roles in GC progression.

MAPK1 is a direct target of miR-454 in GC. To determine the molecular mechanisms underlying the biological roles of miR-454 in GC, bioinformatics analysis was conducted to identify the potential targets of miR-454. MAPK1 was predicted as a candidate of miR-454 and thus selected for further confirmation because it was overexpressed in GC (28) and acted as an oncogene in GC formation and progression (29-33). Fig. 3A shows putative target sites of miR-454 in 3'-UTR of MAPK1. For the confirmation of this hypothesis, luciferase reporter assays were carried out on the MGC-803 and AGS cells transfected with luciferase plasmid harbouring a wild-type or mutant-type seed region in the 3'-UTR of MAPK1 and miR-454 mimics or miR-NC. As shown in Fig. 3B, the cotransfection of miR-454 mimics and wild-type MAPK1 3'-UTR significantly reduced luciferase activities (P<0.05). However, the cotransfection of the mutant MAPK1 3'-UTR and miR-454 mimics did not affect the luciferase activities in both MGC-803 and AGS cells. To further investigate whether miR-454 is capable of regulating MAPK1 expression in GC, RT-qPCR and western blot analysis were utilized to detect MAPK1 expression in MGC-803 and AGS cells after transfection with miR-454 mimics or miR-NC. The present results demonstrated that enforced expression of miR-454 significantly decreased MAPK1 expression in MGC-803 and AGS cells at mRNA level (Fig. 3C; P<0.05) and protein level (Fig. 3D; P<0.05). Therefore, MAPK1 is a direct target of miR-454 in GC.

MAPK1 is upregulated in GC tissues and negatively correlated with miR-454 expression levels. To elucidate the clinical relevance of miR-454 and MAPK1 in GC, we detected MAPK1 mRNA expression in 45 paired GC tissues and adjacent non-tumourous tissues by RT-qPCR. The results showed that the MAPK1 mRNA was significantly higher in GC tissues than in matched adjacent non-tumourous tissues (Fig. 4A; P<0.05). Western blot analysis results also indicated that the protein expression level of MAPK1 was highly expressed in GC tissues (Fig. 4B). Furthermore, Spearman's correlation analysis indicated an inverse association between miR-454 and MAPK1 mRNA levels in GC tissues (Fig. 4C; r= -0.6191, P<0.0001). These results further confirmed that MAPK1 expression was negatively regulated by miR-454 in GC, suggesting that the inhibitory effect of miR-454 on the MAPK1 is clinically relevant in GC.

MAPK1 knockdown exhibits similar effects to that of miR-454 overexpression in GC cells. MAPK1-specific siRNA was used to knockdown MAPK1 expression in the
Figure 5. Downregulation of MAPK1 inhibits GC cell proliferation, invasion and promotes apoptosis in vitro. (A) MGC-803 and AGS cells were transfected with MAPK1 siRNA or NC siRNA. After transfection, western blot analysis was performed to detect MAPK1 protein expression. *P<0.05 compared with NC siRNA. (B-D) The effects of MAPK1 underexpression on MGC-803 and AGS cell proliferation, invasion and apoptosis were explored using CCK-8 assay, Transwell invasion assay and flow cytometry analysis, respectively. ^P<0.05 compared with NC siRNA. GC, gastric cancer; CCK-8, Cell Counting Kit-8.
Figure 6. MAPK1 upregulation reverses the tumour-suppressive effects of miR-454 overexpression on GC cells. (A) MAPK1 protein level was determined in MGC-803 and AGS cells transfected with miR-454 mimics with or without pcDNA3.1-MAPK1. *P<0.05 compared with miR-NC. #P<0.05 compared with miR-454 mimics+pcDNA3.1-MAPK1. (B-D) Cell proliferation, invasion and apoptosis were assessed in MGC-803 and AGS cells transfected with miR-454 mimics with or without pcDNA3.1-MAPK1 by CCK8 assay, Transwell invasion assay and flow cytometry analysis. *P<0.05 compared with miR-NC. #P<0.05 compared with miR-454 mimics+pcDNA3.1-MAPK1. GC, gastric cancer; CCK-8, Cell Counting Kit-8.
MGC-803 and AGS cells to evaluate the effects of MAPK1 underexpression in GC cells after transfection with MAPK1 siRNA. Following transfection with MAPK1 siRNA, western blot analysis confirmed that MAPK1 was significantly down-regulated in both MGC-803 and AGS cells (Fig. 5A; P<0.05). Functional experiments revealed that MAPK1 knockdown inhibited cell proliferation (Fig. 5B; P<0.05) and invasion (Fig. 5C; P<0.05) and induced apoptosis (Fig. 5D; P<0.05) of MGC-803 and AGS cells, which resembled the suppressive effects of miR-454 overexpression in GC. These results further demonstrated that MAPK1 is a direct downstream target of miR-454 in GC.

Upregulation of MAPK1 reverses the tumour-suppressing effects of miR-454 in GC. Rescue experiments were performed to further determine whether MAPK1 is responsible for the functional effects of miR-454 in GC cells. First, we transduced miR-454 mimics in MGC-803 and AGS cells, with or without MAPK1 overexpression plasmid. Western blot analysis revealed that ectopic expression of miR-454 suppressed MAPK1 protein expression, whereas co-transfection of pcDNA3.1-MAPK1 can recover the MAPK1 expression in MGC-803 and AGS cells (Fig. 6A; P<0.05). In addition, the co-transfection of MAPK1 overexpression rescued the effects of miR-454 overexpression on the proliferation (Fig. 6B; P<0.05), invasion (Fig. 6C; P<0.05) and apoptosis (Fig. 6D; P<0.05) in MGC-803 and AGS cells. These results clearly demonstrated that miR-454 inhibited cell proliferation and invasion and promoted apoptosis in GC cells, at least in part by the downregulation of MAPK1.

Discussion

Recently, numerous miRNAs have been determined to contribute to gastric cancer (GC) initiation and progression, suggesting that miRNAs may be developed as effective diagnostic and prognostic molecular biomarkers and can be investigated as therapeutic targets for the patients with this disease (34-36). Therefore, further investigation of the miRNAs involved in GC development represents an opportunity to improve the prognosis of GC patients. In the present study, miR-454 was observed to be downregulated in GC tissues and cell lines. To clarify the clinical significance of miR-454 in GC, the median value (37,38) was used as the reference value to define the low/high expression of miR-454 expression in tumour tissues. The analysis showed that low miR-454 expression level in GC was correlated with lymph node metastasis, invasive depth and TNM stage. To explore the functions of miR-454 in GC, MGC-803 and AGS cell lines expressed the lowest levels of miR-454 among the five GC cell lines tested and were selected for further experiments. The functional experiments indicated that upregulation of miR-454 inhibited GC cell proliferation and invasion and induced apoptosis in vitro. Notably, MAPK1 was identified as a direct target gene of miR-454 in GC. MAPK1 was highly expressed in the GC tissues and was inversely associated with miR-454 expression level. Moreover, MAPK1 knockdown exhibited similar effects to that of miR-454 overexpression in GC cells. MAPK1 overexpression rescued the tumour-suppressing effects of miR-454 in GC.

Dysregulation of miR-454 has been reported in multiple types of human cancer. For example, miR-454 was upregulated in breast cancer. High expression level of miR-454 indicated worse disease-free survival. miR-454 expression was positively correlated with worse clinical outcome for triple-negative breast cancer subtype (39). miR-454 was also found to be highly expressed in non-small cell lung cancer tissues and cell lines. High miR-454 expression is strongly associated with lymph node metastasis, advanced TNM stage and short overall survival. Multivariate regression analysis identified miR-454 overexpression as an independent unfavourable prognostic factor for patients with non-small cell lung cancer tissues (40). In hepatocellular carcinoma, miR-454 expression level was increased in the tumour tissues. Patients with hepatocellular carcinoma and high expression miR-454 level have lower 5-year overall survival and decreased disease-free survival than patients with low miR-454 levels (41). High expression level of miR-454 was also observed in uveal melanoma (23) and colorectal cancer (24). However, miR-454 was demonstrated to be downregulated in glioblastoma (25), osteosarcoma (26) and pancreatic ductal adenocarcinoma (42,43). These findings suggest that miR-454 expression exhibits tissue specificity and may be a diagnostic and prognostic biomarker for various types of cancer.

Abnormally expressed miR-454 plays oncogenic roles in the occurrence and development of several types of human cancers. For instance, Zhu et al reported that miR-454 underexpression inhibited cell proliferation, migration and invasion and promotes apoptosis in non-small cell lung cancer (40). Yu et al found that the downregulation of miR-454 suppressed hepatocellular carcinoma cell metastasis and epithelial-mesenchymal transition (44). Sun et al revealed that the upregulation of miR-454 promoted cell proliferation, colony formation and invasion in uveal melanoma (23). Liang et al showed that restoration of miR-454 expression increased the proliferation and anchorage-independent growth of colorectal cancer cells (24). Nevertheless, miR-454 was validated as a tumour suppressor in glioblastoma. Ectopic expression of miR-454 reduced glioblastoma cell proliferation and induced cell cycle arrest at G0/G1 phase (25). Niu et al indicated that resumption of miR-454 expression attenuated cell growth and invasion of osteosarcoma (26). Fan et al showed that restored miR-454 expression decreased cell growth, metastasis and angiogenesis in pancreatic ductal adenocarcinoma (42,43). These conflicting findings demonstrated that the biological functions of miR-454 have tissue specificity, and may be explained by the ‘imperfect complementarity’ of the interactions between miRNAs and their target genes (45). These findings also suggested that miR-454 may be a novel therapeutic target for the development of antineoplastic agents.

Several targets of miR-454 were identified, including PTEN (40) in non-small cell lung cancer, CHD5 (44) in hepatocellular carcinoma, CYLD (24) in colorectal cancer, PDK1 (25) in glioblastoma, c-Met (26) in osteosarcoma, LRP6 (42) and SDF-1 (43) in pancreatic ductal adenocarcinoma. In this study, MAPK1 was validated as a direct and functional downstream target of miR-454. MAPK1, which was known as ERK2, is a critical component of the MAPK signalling pathway (46). Previous studies reported that MAPK1 is upregulated in several types of human cancer, such as...
cervical cancer (47), ovarian cancer (48), non-small cell lung cancer (49), glioblastoma multiforme (50) and myeloma (51). Liang et al reported that MAPK1 is highly expressed in GC tissues and is correlated with TNM stage, serosa invasion and lymph node involvement (28). Functional assays revealed that MAPK1 participates in regulating GC cell proliferation, apoptosis, migration, invasion and metastasis (29-33). In combination with the present findings, the miR-454/MAPK1 pathway shows a potential to be investigated as a therapeutic method for patients with GC.

In conclusion, miR-454 is significantly downregulated in GC, and low expression level of this miRNA is associated with lymph node metastasis, invasive depth and TNM stage. In vitro studies demonstrated that miR-454 inhibited GC cell proliferation and invasion and increased apoptosis. Mechanistically, MAPK1 was identified as a direct target gene of miR-454 in GC. miR-454 may provide novel therapeutic targets for the treatment of GC. However, more experiments should be made in order to confirm that cell apoptosis is indeed affected. In addition, we will investigate the expression level of miR-454 in the serum of GC patients, and evaluate whether it can be used as a biomarker of GC prognosis.

Acknowledgements

This study was supported by Young Scholars Fund of Bethune Medical Scientific Research Program (no. 2013206042), and Special Health Projects of Jilin Province.

References


