MicroRNA-411 inhibits malignant biological behaviours of colorectal cancer cells by directly targeting PIK3R3

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Abstract. Colorectal cancer (CRC) is the third most common cancer and the fourth leading cause of cancer-related mortality worldwide. Aberrant expression of miRNAs play important roles in the development and progression of various types of cancers by modulating oncogenic and tumour-suppressor pathways. Therefore, exploring the functions of microRNAs (miRNAs) that specifically contribute to CRC tumourigenesis and tumour development would greatly aid in obtaining more information on CRC and provide new targets for its diagnosis and treatment. miRNA-411 (miR-411) was previously observed to be aberrantly expressed in multiple human cancers. However, the expression pattern, function and underlying molecular mechanism of miR-411 in CRC remain unclear. Therefore, the present study was performed to detect miR-411 expression, investigate the biological roles of miR-411 and identify its mechanism of action in CRC cells. Here, miR-411 expression was significantly downregulated in human CRC tissues and cell lines, and low levels of miR-411 were correlated with lymph node metastasis, distant metastasis and TNM stage. Resumed expression of miR-411 suppressed cell proliferation and invasion but promoted apoptosis in CRC. Additionally, phosphoinositide-3-kinase regulatory subunit 3 (PIK3R3) was identified as a direct target of miR-411 in CRC. PIK3R3 was upregulated in the CRC tissues and inversely correlated with miR-411 expression. Downregulation of PIK3R3 had tumour-suppressive effects similar to those of miR-411 overexpression in CRC. Moreover, upregulation of PIK3R3 could rescue the tumour-suppressing effects of miR-411 overexpression in CRC cells. More importantly, miR-411 specifically suppressed the activation of the AKT/mTOR signalling pathway in CRC. Therefore, miR-411 functions as a tumour-suppressive miRNA by directly targeting PIK3R3 and indirectly regulating AKT/mTOR signalling pathway. miR-411 may serve as a new therapeutic target for patients with CRC.

Introduction

Colorectal cancer (CRC) is the third most common cancer and the fourth leading cause of cancer-related mortality worldwide (1,2). Approximately 1 million new cases and 600,000 deaths due to CRC are estimated to occur annually around the world (3). CRC is one of the most prevalent cancers in western populations (4) but has low incidence rates in Asia, Africa and South America (5). However, the frequency of CRC in China has rapidly increased such that this carcinoma has emerged as the fifth most common cancer and the fourth most common cause of cancer-related deaths in the country (6). Surgical resection followed by chemotherapy and/or radiotherapy is currently the effective modality for CRC patients (7). Prognosis of CRC patients remains unsatisfactory despite the remarkable developments in the diagnosis and treatments of CRC (8). Local recurrence and distant metastasis are the primary causes of the unfavourable prognosis of CRC (9). Therefore, understanding the molecular mechanisms underlying CRC progression is essential to identify effective biomarkers and novel therapeutic methods for CRC patients.

MicroRNAs (miRNAs) are an emerging group of single-strand non-coding small RNAs (~22 nucleotides) first discovered in the early 1990s in Caenorhabditis elegans (10). miRNAs negatively regulate gene expression by interacting with the 3'-untranslated regions (UTRs) of corresponding target messenger RNAs (mRNAs) in a base-pairing manner, leading to translational repression or degradation of target mRNAs (11). Computational estimations suggest ~1,000 miRNAs in the human genome, which regulate one-third of human protein-encoding genes (12). Multiple bodies of evidence have indicated that miRNAs are abnormally expressed in almost all human neoplasms (13,14). Previous studies have demonstrated that miRNA deregulation may be significantly correlated with diagnosis, treatment and prognosis in human cancer (15-17). miRNAs play important roles in tumourigenesis and tumour development by regulating many diverse biological processes, such as cell proliferation, cycle, apoptosis, invasion, migration, metastasis, angiogenesis and epithelial-mesenchymal transition (18-20). miRNAs can

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act as tumour suppressors or oncogenes, depending on their transcript targets (21,22). Therefore, miRNAs have been proposed to be potential indicators and therapeutic targets in various types of cancers (23,24).

miR-411, located on chromosome 14q32, was previously observed to be aberrantly expressed in multiple human cancers (25-28). However, the expression pattern, function and underlying molecular mechanism of miR-411 in CRC remain unclear. Therefore, the present study was performed to detect miR-411 expression, investigate the biological roles of miR-411 and identify its mechanism of action in CRC cells.

Materials and methods

Tumour specimens. The present study was approved by the Human Ethics Committee of the Cancer Hospital of China Medical University, Liaoning Cancer Hospital and Institute. Written informed consent was obtained from all patients. A total of 46 CRC tissues and corresponding adjacent normal tissues were obtained from patients who underwent surgical resection at the Department of Colorectal Surgery, Cancer Hospital of China Medical University, Liaoning Cancer Hospital and Institute between May 2014 and February 2016. None of these patients with CRC had been treated with chemotherapy, radiotherapy or any adjuvant therapy before surgery. All tissue specimens were immediately frozen in liquid nitrogen after resection and stored at -80˚C.

Cell lines and culture condition. CRC cell lines (SW480, SW620, HCT116, HT29, CaCo-2 and LoVo) and the 293T cell line were purchased from the American Type Culture Collection (ATCC; Manassas, DE, USA). For miR-411 detection, the total RNA was reverse-transcribed to cDNA using a TaqMan MicroRNA Reverse Transcription kit (Applied Biosystems, Carlsbad, CA, USA). The relative expression of miR-411 was examined by TaqMan MicroRNA PCR kit (Applied Biosystems). The miR-411 expression levels were normalised to those of U6 snRNA. PIK3R3 mRNA was quantified by synthesising cDNA using PrimeScript RT reagent kit (Takara Bio, Dalian, China). These cDNAs were used to detect the expression of PIK3R3 mRNA by quantitative PCR using a SYBR Premix Ex Taq™ kit (Takara Bio). The PIK3R3 mRNA expression was normalised to those of GAPDH. The primers were designed as follows: miR-411, 5’-GGGGTATGAGCCGTATGAG-3’ (forward) and 5’-TGGCCTGCTGGGATTTC-3’ (reverse); U6 snRNA, 5’-CTCGTTGCGAACA-3’ (forward) and 5’-TGGGTTCGTGAGGTGACG-3’ (reverse); PIK3R3, 5’-CTTGCCTCCTGTTGGAAGGCGAT-3’ (forward) and 5’-GACGTTGAGGAGTCGTTCTGAT-3’ (reverse); and GAPDH, 5’-GAAGGTGAAGGTCGGAATGTG-3’ (forward) and 5’-GAAGATGTTGATGGATTCTTC-3’ (reverse). Relative expression levels were calculated using the 2-ΔΔCt method (29).

Cell Counting Kit-8 (CCK-8) assay. Cell proliferation was determined using the CCK-8 assay. Briefly, transfected cells were harvested at 24 h post-transfection. A total of 3x10³ cells suspended in 200 µl of DMEM with 10% FBS were seeded into each well of a 96-well plate and cultured at 37˚C for 0, 1, 2 or 3 days. At each time-point, 10 µl of CCK-8 solution (Dojindo Molecular Technologies, Kumamoto, Japan) was added to each well. After an additional 2 h of incubation at 37˚C, the optical density of each well at a wavelength of 450 nm was measured with a microplate reader (Model 550; Bio-Rad Laboratories, Shanghai, China). Each assay was performed in quintuplicate and repeated at least thrice.

Matrigel invasion assay. Transwell chambers (Corning Inc., Cambridge, MA, USA) with 8-µm pore size filters covered with Matrigel (BD Biosciences, Franklin Lakes, NJ, USA) were used to perform the Matrigel invasion assay. Transfected cells were harvested at 48 h post-transfection. Transfected cells (5x10⁴) in FBS-free DMEM medium were seeded in the upper chambers, and DMEM containing 10% FBS was supplemented into the lower chambers to serve as a chemoattractant. After incubation at 37˚C under 5% CO₂ for 24 h, cells that did not invade through the pores were removed carefully by a cotton swab. Invasive cells were fixed using 4% paraformaldehyde, stained with 0.1% crystal violet solution, washed in phosphate-buffered saline (PBS) and dried in air. Images were captured, and cell number was counted in five randomly selected fields under an inverted microscope (x200 magnifications; X71; Olympus Corp., Tokyo, Japan).

Flow cytometry. Cell apoptosis was analysed using the Annexin V-FITC apoptosis detection kit (Invitrogen). Transfected cells were harvested at 48 h post-transfection, washed with PBS and fixed using 80% ice-cold ethanol in PBS. Afterwards, the cells were resuspended in 300 µl of 1X binding buffer and stained with 5 µl of FITC-Annexin V and 5 µl of propidium iodide (PI) in the dark at room temperature for 20 min. BD FACSCalibur™ flow cytometer was used to detect the cell apoptosis.
miR-411 is downregulated in human CRC tissues and cell lines. RT-qPCR was performed on 46 paired CRC tissues and corresponding adjacent normal tissues to investigate the status of miR-411 in CRC. Compared with the corresponding adjacent normal tissues, the expression of miR-411 was significantly downregulated in the CRC tissues (Fig. 1A; P<0.05). miR-411 expression levels in six CRC cell lines (SW480, SW620, HCT116, HT29, CaCo-2 and LoVo) and normal FHC cells were also analysed by RT-qPCR. Fig. 1B shows that all tested CRC cells showed significantly lower expression levels of miR-411 compared with the normal FHC (P<0.05). These data suggest that miR-411 plays important roles in CRC formation and progression.

Low expression of miR-411 correlates with adverse clinicopathological characteristics of CRC patients. miR-411 expression and clinical characteristics of patients with CRC were associated to explore the clinical value of miR-411 in CRC. The CRC patients were subsequently divided into either the miR-411 low-expression group (n=23) or the miR-411 high-expression group (n=23). The median expression level of miR-411 in all samples was regarded as the cut-off. As shown in Table I, the low expression level of miR-411 was significantly correlated with lymph node metastasis (P=0.018), distant metastasis (P=0.007) and TNM stage (P=0.008). However, no correlation was observed between miR-411 expression and other clinicopathological characteristics, such as sex (P=0.555), age (P=0.548), tumour size (P=0.345) and differentiation (P=0.552). These results suggest that miR-411 is a possible prognostic biomarker for CRC patients.
both with relatively lower endogenous miR-411, to transfect miR-411 mimics to investigate the functional roles of miR-411 in CRC cells. After transfection, RT-qPCR analysis demonstrated that miR-411 was markedly upregulated in the HCT-116 and CaCo-2 cells after transfection with miR-411 mimics (50 nM; Fig. 2A, P<0.05). Subsequently, the effect of miR-411 overexpression on cell proliferation was assessed. CCK-8 assay indicated that the ectopic expression of miR-411 inhibited HCT-116 and CaCo-2 cell proliferation (Fig. 2B; P<0.05). Matrigel invasion assay was performed to examine the invasion abilities of HCT-116 and CaCo-2 cells after transfection with miR-411 mimics or miR-NC. The results revealed...
that restored expression of miR-411 decreased cell invasion abilities in HCT-116 and CaCo-2 cells (Fig. 2C; P<0.05). Then, we analysed the effect of miR-411 on cell apoptosis using flow cytometry. Fig. 2D shows that upregulation of miR-411 increased the apoptosis rate in the HCT-116 and CaCo-2 cells (P<0.05). Thus, these findings suggest that miR-411 plays a suppressive role in CRC progression.

PIK3R3 is a direct target of miR-411 in CRC. Potential targets of miR-411 were predicted using bioinformatic analysis to determine the molecular mechanisms responsible for the tumour-suppressing roles of miR-411. Among hundreds of potential candidates, PIK3R3 was selected for further target identification (Fig. 3A) because of its important roles in CRC tumourigenesis and tumour development (30). In addition, PIK3R3 was also validated as a direct target of other miRNAs in CRC. Hence, PIK3R3 was chosen for further confirmation. To verify this prediction, luciferase reporter assay was conducted in 293T cells cotransfected with miR-411 mimics or miR-NC and pMIR-PIK3R3-3’-UTR Wt or pMIR-PIK3R3-3’-UTR Mut. The results showed that miR-411 significantly repressed the relative luciferase activities with the wild-type PIK3R3 3’-UTR (Fig. 3B; P<0.05) but did not alter the activity of the mutant PIK3R3 3’-UTR. To further confirm the potential role of miR-411 in the regulation of PIK3R3, we detected PIK3R3 expression in HCT-116 and CaCo-2 cells following transfection with miR-411 mimics or miR-NC. RT-qPCR and western blot analysis revealed that miR-411 overexpression caused a significant downregulation in PIK3R3 expression at the mRNA (Fig. 3C; P<0.05) and protein (Fig. 3D; P<0.05) levels in the HCT-116 and CaCo-2

### Table I. Correlation between the miR-411 expression and clinicopathological features of CRC cases.

<table>
<thead>
<tr>
<th>Features</th>
<th>Cases</th>
<th>Low</th>
<th>High</th>
<th>P-value</th>
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<tr>
<td>Sex</td>
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<td>0.555</td>
</tr>
<tr>
<td>Male</td>
<td>24</td>
<td>13</td>
<td>11</td>
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</tr>
<tr>
<td>Female</td>
<td>22</td>
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<td>12</td>
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<tr>
<td>Age (years)</td>
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<tr>
<td>≥60</td>
<td>18</td>
<td>8</td>
<td>10</td>
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<td>Tumour size (cm)</td>
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<tr>
<td>≥5</td>
<td>31</td>
<td>14</td>
<td>17</td>
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<tr>
<td>Differentiation</td>
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<td>12</td>
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<tr>
<td>Poor</td>
<td>20</td>
<td>9</td>
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<tr>
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<tr>
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<td>0.007*</td>
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<tr>
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<tr>
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<td>19</td>
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<td>15</td>
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<tr>
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CRC, colorectal cancer; miR-411, microRNA-411; TNM, tumour-node-metastasis. *Statistically significant.

![Graph A](image1.png)

![Graph B](image2.png)

![Graph C](image3.png)

![Graph D](image4.png)

Figure 3. PIK3R3 is a direct target of miR-411 in CRC. (A) The miR-411 target sequences in the 3’-UTR of PIK3R3 and its mutant containing altered nucleotides in the 3’-UTR were predicted. PIK3R3 was upregulated in CRC and plays important roles in CRC tumourigenesis and tumour development. In addition, PIK3R3 was also validated as a direct target of other miRNAs in CRC. Hence, PIK3R3 was chosen for further confirmation. (B) Luciferase reporter assay was performed using 293T cells cotransfected with Wt or Mut PIK3R3 3’-UTR reporter plasmid, miR-411 mimics or miR-NC. Luciferase activity was determined by Dual-Luciferase reporter system. (C and D) PIK3R3 mRNA and protein expression levels in HCT-116 and CaCo-2 cells transfected with miR-411 mimics or miR-NC were determined by RT-qPCR and western blot analysis.
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These results showed that PIK3R3 is a direct target of miR-411 in CRC. Similar findings were observed in CRC tissues. We measured PIK3R3 expression in 46 paired CRC tissues and corresponding adjacent normal tissues to further examine the relationship between miR-411 and PIK3R3 mRNA expression levels in CRC tissues. Spearman's correlation analysis showed an inverse relationship between miR-411 and PIK3R3 mRNA expression levels in CRC tissues ($r = -0.6753, P < 0.0001$).

miR-411 level is negatively correlated with PIK3R3 level in CRC tissues. We measured PIK3R3 expression in 46 paired CRC tissues and corresponding adjacent normal tissues to further examine the relationship between miR-411 and PIK3R3. Fig. 4A shows that PIK3R3 mRNA was upregulated in CRC tissues compared with that in adjacent normal tissues ($P < 0.05$). Western blot results showed that PIK3R3 protein was highly expressed in CRC tissues (Fig. 4B; $P < 0.05$). Furthermore, an inverse association between miR-411 and PIK3R3 mRNA level in CRC tissues was observed using Spearman's correlation analysis (Fig. 4C, $r = -0.6753, P < 0.0001$).

PIK3R3 knockdown inhibits proliferation and invasion but increases apoptosis in CRC cells. We used PIK3R3-specific siRNA to genetically knockdown endogenous PIK3R3 expression in HCT-116 and CaCo-2 cells to verify whether PIK3R3 knockdown would simulate the miR-411-mediated effects. Western blot analysis was used to determine the transfection efficiency and confirmed that PIK3R3 was significantly downregulated in the PIK3R3 siRNA-transfected HCT-116 and CaCo-2 cells (Fig. 5A; $P < 0.05$). Next, CCK-8 assay, Matrigel invasion assay and flow cytometric analysis demonstrated that down-regulation of PIK3R3 suppressed cell proliferation (Fig. 5B; $P < 0.05$) and invasion (Fig. 5C; $P < 0.05$) but increased apoptosis (Fig. 5D; $P < 0.05$) in the HCT-116 and CaCo-2 cells. Thus, our data confirmed that PIK3R3 downregulation had similar tumour-suppressive effects as miR-411 overexpression in CRC. These results further suggest that PIK3R3 is a direct downstream target of miR-411 in CRC.

Overexpression of PIK3R3 abrogates the tumour-suppressing effects of miR-411 in CRC cells. We confirmed that PIK3R3 is a direct target of miR-411. Thus, rescue experiments were employed to disclose whether PIK3R3 overexpression could abolish the tumour-suppressing effects induced by miR-411 in CRC cells. HCT-116 and CaCo-2 cells were transfected with miR-411 mimics with or without PIK3R3 overexpression plasmid pcDNA3.1-PIK3R3. After transfection, western blot results showed that the reduced PIK3R3 protein expression caused by miR-411 mimics was markedly restored by transfection of the pcDNA3.1-PIK3R3 in the HCT-116 and CaCo-2 cells (Fig. 6A; $P < 0.05$). In addition, the effects of miR-411 on CRC cell proliferation (Fig. 6B; $P < 0.05$), invasion (Fig. 6C; $P < 0.05$) and apoptosis (Fig. 6D; $P < 0.05$) were significantly reversed by PIK3R3 overexpression. These results indicated that miR-411 served as a tumour suppressor in CRC, at least in part, by directly regulating PIK3R3.

miR-411 regulates the AKT/mTOR signalling pathways in CRC. PIK3R3 plays crucial roles in tumourigenesis and tumour development by regulating the AKT/mTOR signalling pathway (32, 33). We detected p-AKT, AKT, p-mTOR and mTOR expression in HCT-116 and CaCo-2 cells transfected with miR-411 mimics or miR-NC to determine whether AKT/mTOR signalling pathway was involved in the regulatory effects of miR-411 on CRC. The results showed that upregulation of miR-411 significantly reduced the p-AKT and p-mTOR expression in HCT-116 and CaCo-2 cells, whereas transfection with miR-411 mimics did not affect the total AKT and mTOR protein levels (Fig. 7; $P < 0.05$). Thus, miR-411 suppressed the activation of the AKT/mTOR signalling pathway by targeting PIK3R3 to inhibit CRC progression.

Discussion

Aberrant expression of miRNAs plays important roles in the development and progression of various cancers by modulating oncogenic and tumour-suppressor pathways (22, 34). Therefore, exploring the functions of miRNAs that specifically contribute to CRC tumourigenesis and tumour development would greatly aid in obtaining more information on CRC and provide new targets for its diagnosis and treatment. In the present study, miR-411 was significantly downregulated in the CRC tissues and cell lines. In addition, the low expression of miR-411 was significantly correlated with lymph node metastasis, distant metastasis and TNM stage of CRC. Furthermore, restored expression of miR-411 suppressed cell proliferation...
and invasion but promoted apoptosis in CRC. Moreover, PIK3R3 was identified as a novel direct target of miR-411 in CRC, and miR-411 could diminish the AKT/mTOR signalling pathway in CRC. These results suggested that miR-411 may be used to design novel prognostic biomarker and therapeutic strategies for CRC patients.

miR-411 is aberrantly expressed in several types of cancers. For example, miR-411 was observed to be highly

Figure 5. Downregulation of PIK3R3 attenuates CRC cell proliferation and invasion but induces cell apoptosis. (A) The PIK3R3 protein expression in HCT-116 and CaCo-2 cells after transfection with PIK3R3 siRNA or NC siRNA was analysed using western blot analysis. *P<0.05 compared with NC siRNA. (B) CCK-8 assay was used to determine cell proliferation in HCT-116 and CaCo-2 cells following transfection with PIK3R3 siRNA or NC siRNA. *P<0.05 compared with NC siRNA. (C) Cell invasion capacities of HCT-116 and CaCo-2 cells transfected with PIK3R3 siRNA or NC siRNA were analysed using Matrigel invasion assay. *P<0.05 compared with NC siRNA. (D) Flow cytometric analysis was performed to assess the apoptosis of HCT-116 and CaCo-2 cells transfected with PIK3R3 siRNA or NC siRNA. *P<0.05 compared with NC siRNA.
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expressed in hepatocellular carcinoma (25). miR-411 was also found to be upregulated in lung cancer (26). However, contradictory reports have demonstrated the downregulated miR-411 expression in breast cancer tissues and cell lines (27,28).

Downregulated miR-411 expression was correlated with lymph node metastasis and histological grade of breast cancer (27). These contradictory studies indicated that miR-411 expression may be subject to tissue-specific regulatory processes in

Figure 6. Restoration of the expression of PIK3R3 abrogates the effects of miR-411 overexpression on CRC cells. (A) The protein expression levels of PIK3R3 were detected by western blot analysis in HCT-116 and CaCo-2 cells transfected with miR-NC, miR-411 mimics or miR-411 mimics+pcDNA3.1-PIK3R3. *P<0.05 compared with miR-NC or miR-411 mimics+pcDNA3.1-PIK3R3. (B) Cell proliferation, (C) invasion and (D) apoptosis in HCT-116 and CaCo-2 cells transfected with miR-NC, miR-411 mimics or miR-411 mimics+pcDNA3.1-PIK3R3 were examined using CCK-8 assay, Matrigel invasion assay and flow cytometric analysis, respectively. *P<0.05 compared with miR-NC or miR-411 mimics+pcDNA3.1-PIK3R3.
induce mRNA degradation when the binding complementarily is perfect. However, miRNAs regulate gene expression at post-transcriptional level when the binding at the 3’UTR of target genes is only partially complementary (37). PIK3R3 was upregulated in CRC tissues and inversely correlated with miR-411 expression levels. Downregulation of PIK3R3 had similar tumour-suppressive effects as miR-411 overexpression in CRC. Moreover, upregulation of miR-411 could rescue the tumour-suppressing effects of miR-411 overexpression on CRC cells. Thus, PIK3R3 is a novel direct and functional downstream target of miR-411 in CRC.

PIK3R3 is a member of the phosphatidylinositol 3-kinase (PI3K) family and was found to be abnormally upregulated in various types of human cancers, such as glioma (38), ovarian (39), gastric (40), hepatocellular carcinoma (32), lung (41) and breast cancer (42). The oncogenic roles of PIK3R3 on cancer initiation and progression has been described in several types of cancers, such as Ewing sarcoma (43), gastric (40), ovarian (39) and breast cancer (42). In CRC, PIK3R3 was upregulated in clinical specimens and cell lines. In addition, upregulated PIK3R3 expression has been positively correlated with CRC metastasis. Functional experiments revealed that PIK3R3 overexpression increased tumour migration, invasion and epithelial-to-mesenchymal transition in vitro and promoted metastasis in vivo (30). The present study revealed that miR-411 inhibited CRC progression by directly targeting PIK3R3 and indirectly regulating AKT/mTOR signalling pathway. Thus, miR-411/PIK3R3/AKT/mTOR pathway may serve as a potential target to treat CRC patients.

In conclusion, the present study provided first evidence that miR-411 exhibited tumour suppressive roles against CRC by directly targeting PIK3R3. These results suggested that miR-411 could be a novel therapeutic target for patients with CRC. In this study, we did not explore the effect of miR-411 on the normal FHC cells.

In the following experiments, we will examine the effects of miR-411 underexpression on FHC cells. The small sample size is a limitation of the study, and we will collect more CRC tissues and the relationship between the pathologies in the top and bottom percentiles and miR-411 expression. Despite we found that miR-411 increased apoptosis of CRC cells, the effect of miR-411 overexpression on the expression levels apoptotic markers did not analyzed. In our further research we should examine the effects of miR-411 on apoptotic markers in CRC cells. In this study, luciferase reporter assays were performed in 293T cells. The luciferase reporter assay will be repeated with the CRC cell lines used in the following experiments.

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References


