miR-378a-5p improved the prognosis and suppressed the progression of hepatocellular carcinoma by targeting the VEGF pathway

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Background: The malignant tumor hepatocellular carcinoma (HCC) has a poor prognosis and ineffective therapeutic options. miR-378a-5p is a micro-ribonucleic acid (miRNA) that is overexpressed in many cancers. However, its role in the progression of human HCC is unclear.

Methods: Quantitative real-time polymerase chain reaction (qRT-PCR) was used to measure miR-378a-5p levels in tissues from patients with HCC and from HCC cell lines. Following transfection, flow cytometry and cell viability assays were used to measure cell proliferation. HCC cell invasive and migration capacities were assessed using Transwell assays. Western blots were performed with HCC cells to identify the expression of vascular endothelial growth factor (VEGF).

Results: The HCC tissues and cells had significantly reduced miR-378a-5p expression compared with normal liver tissues and cells, while miR-378a-5p mimics suppressed the colony formation, viability, migration and invasive capacity of HCC cells. The HCC tissues and cell lines had upregulated VEGF expression. In HCC cells, miR-378a-5p expression was negatively correlated with VEGF expression, and miR-378a-5p targeted VEGF in HCC cells.

Conclusions: miR-378a-5p improved the HCC prognosis and suppressed HCC progression by targeting the VEGF pathway.

Keywords: Hepatocellular carcinoma (HCC); miR-378a-5p; vascular endothelial growth factor; migration; invasion; progression

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Introduction

Among malignant tumors, hepatocellular carcinoma (HCC) is the sixth most common in terms of occurrence and cancer-related death (1,2). Typically, HCC treatment involves surgery-based comprehensive therapies. However, the HCC prognosis remains poor, even following surgical resection, as there is a high frequency of recurrence and cancer cell migration into adjunct normal tissues (3,4). The 5-year postoperative recurrence rate of HCC is >60%, mainly due to recurrence and chemoresistance (5,6). Therefore, novel molecular mechanisms must be developed to treat tumor growth and migration.

Micro-ribonucleic acids (miRNAs) are 19–22 nucleotides long and noncoding, and induce mRNA degradation by binding of the target mRNA 3’-untranslated region (UTR) or inhibiting translation (7). Studies have found
that miRNAs have key functions in many physiological and pathological processes, and miRNAs act as oncogenes or tumor-suppressive genes to regulate cancer progression (8). Accumulating evidence shows that in liver cancer, miRNAs have possible applications as tissue-specific biomarkers. Fang et al. found that miR-188-5p directly targeted fibroblast growth factor 5 (FGF5) to suppress HCC tumor cell proliferation and metastasis (9). In human HCC, miR-1180 promotes resistance to apoptosis via nuclear factor (NF)-κB signaling pathway activation (10). Other miRNAs, including miR-345, miR-487a, and miR-613, are abnormally expressed and promote or suppress HCC incidence and development (11-13). Therefore, the unique expression features of miRNA may form a basis for novel markers for diagnostic and therapeutic targets in HCC.

miR-378a-5p is obtained from miR-378, which has oncogenic properties and is expressed in many cancers (14,15). Kooistra et al. found that miR-378a-5p negatively regulates oncogene-induced senescence (16). miR-378a-5p has been reported to suppress proliferation in some tumor cells, including renal cell carcinoma (17), triple-negative breast cancer (18), and colorectal cancer (19). Guo et al. found that high miR-378 levels decreased vascular endothelial growth factor (VEGF) production, which correlates with the most aggressive and poorly differentiated form of gastric cancer (19). Interestingly, studies found that the VEGF pathway is more highly expressed in HCC, which may promote HCC migration, invasion, and angiogenesis (20,21). Nevertheless, miR-378a-5p’s effects in regulating HCC biology by targeting the VEGF pathway remain unclear.

Here, we hypothesized that miR-378a-5p suppresses HCC progression by targeting VEGF, and we investigated the potential molecular mechanism of miR-378a-5p and its correlation with the VEGF pathway in HCC progression. We investigated miR-378a-5p expression in HCC cell lines and tissues from human patients, analyzed miR-378a-5p cellular functions, namely invasion, cell viability, colony formation, and migration, in HCC cells, and examined the fundamental mechanisms of miR-378a-5p functions in liver cancer.

**Methods**

**Patients and samples**

Fifty primary liver cancer samples and their adjacent normal tissue samples from liver cancer patients who underwent surgical treatment at the Sichuan Academy of Medical Sciences and Sichuan Provincial People’s Hospital (Sichuan, China) between 2015 and 2017 were used in this study. The patients’ clinicopathological parameters, including sex, age, serum alpha-fetoprotein (AFP), lymph node metastasis, tumor size, tumor-node-metastasis (TNM) stage, vascular invasion, and distant metastasis, were recorded. The Sichuan Academy of Medical Sciences and Sichuan Provincial People’s Hospital Ethics Committee approved all study protocols.

**Cell cultures and transfection**

We purchased human HCC Hep3B and SNU-449 cells from American Type Culture Collection (Manassas, VA, USA). We authenticated the Hep3B and SNU-449 cells with short tandem repeat profiling. High-resolution screening and interspecies cross-contamination detection were performed using the AmpFLSTR™ Identifiler™ Plus PCR Amplification Kit. Dulbecco’s modified Eagle’s medium (DMEM; Invitrogen, Carlsbad, CA, USA) was used to culture the HCC cells. The culture medium was incubated at 37 °C in 5% CO₂ and humidified air and was supplemented with 10% fetal bovine serum and 1% streptomycin/penicillin (Gibco). The HCC cells were transfected with miR-378a-5p mimics and negative control (NC) mimics (ZoonBio Biotechnology Co., Ltd., Nanjing, Jiangsu, China) using Lipofectamine™ 2000 (Invitrogen, Carlsbad, CA, USA).

**Western blot analysis**

We prepared human HCC tissue samples and cell lysates in a buffer mixture containing 1 mL radioimmunoprecipitation assay (Beyotime, Shanghai, China). Sodium dodecyl sulfate-polyacrylamide gel was used to separate the total protein, which was shifted to polyvinylidene difluoride membranes (EMD Millipore, Billerica, MA, USA) at 320 mA at 4 °C for 2 h. Non-fat milk (10%) was used to block the membranes, which were next incubated with the primary antibodies (anti-VEGF-A: 1:1,000 dilutions, Abcam, Eugene, OR, USA) overnight. Horseradish peroxidase-conjugated secondary antibody [Peroxidase-AffiniPure Goat Anti-Rabbit IgG (H + L), 1:200 dilution, ZSGB-Bio, Nanjing, Jiangsu, China] was used to probe the membranes at room temperature for 2 h. The protein bands were visualized with electrochemiluminescence (Millipore, Billerica, MA, USA).
**Quantitative real-time polymerase chain reaction (qRT-PCR)**

We extracted HCC tissue samples and total RNA from HCC cells using TRIzol (Invitrogen). miR-378a-5p expression was measured with the TaqMan MicroRNA Reverse Transcription Kit (TâKaRa, Otsu, Shiga, Japan) and miRNA-specific TaqMan microRNA assays (Applied Biosystems, San Diego, CA, USA). The qRT-PCR assays for miR-378a-5p detection were conducted on a SLAN-96P RT-PCR instrument (Sansure Biotech, Changsha, Hunan, China). miR-378a-5p expression levels were expressed as fold differences relative to that of RNU6B (RNA, U6 small nuclear 6, pseudogene, an internal control) using the comparative threshold cycle ($2^{-\Delta\Delta Ct}$) method. The RNU6B- and miR-378a-5p-specific forward primers were 5’-ACGCAAATTCGTGAAGCGTT-3’ and 5’-CTGACTCCAGGTCCTGTGTA-3’, respectively. The corresponding reverse primers were Uni-miR qRT-PCR primers (TâKaRa, Otsu, Shiga, Japan).

**Cell proliferation assay**

HCC cell proliferative ability was determined using Cell Counting Kit-8 (Dojindo, Kyoto, Japan). Cells (3×10^5 per well) were seeded in 6-well plates and maintained in an incubator. After transfection for 24 h, 5×10^3 cells per well were incubated in Dulbecco's modified eagle medium in 96-well plates. The absorbance values of the experimental wells were read at 450 nm after 24, 48, 72, and 96 h of incubation using a microplate reader.

**Colony formation assays**

Hep3B or SNU-449 cell lines were transfected for 48 h. Next, we seeded and scattered the Hep3B or SNU-449 cells equally in 6-well tissue culture plates and cultured the cells in complete medium (2 mL). The culture medium was refreshed every 4 days with 1× phosphate buffer solution. At the end of the experiment, methanol was used to fix the surviving colonies, and crystal violet (1.25%) was used to stain them. We counted colonies with at least 50 cells under a light microscope.

**Wound-healing assays**

We seeded HCC cells (3×10^5 per well, 12-well plates) and incubated them for 24 h to form a monolayer of approximately 80% confluence. A scratch wound was gently made in each well using a pipette tip (100 μL). Phase-contrast images of the same field were obtained at the 0 and 48 h under 100x magnification.

**Transwell assays**

We used a Transwell assay to measure the HCC cell migration and invasive ability using 24-well Transwell chambers. After 24 h of transfection, the upper chamber was filled with the cell suspensions, and the bottom chamber was filled with 10% FBS-containing medium. Next, we cultured the cells in a humidified incubator (24 h, 5% CO₂, 37 °C). Subsequently, we fixed, stained by crystal violet (1.25%), photographed, and counted the cells that had migrated or invaded and that had adhered to the surface on the bottom. Finally, the number of stained cells was determined under a microscope at 200x.

**Dual-luciferase reporter gene analysis**

The has-miR-378a-5p target was identified by TargetScan online software (http://www.targetscan.org). VEGF oligonucleotides (positions 2088–2094) in which miR-378a-5p binding sites were present (WT) or deleted (MUT) were inserted in the pGL3-control vector to create recombinant plasmids according to the protocols. Luciferase activity was detected using Hep3B or SNU-449 cells. After 48 h of incubation, the cells were lysed in cell culture luciferase lysis buffer, and the luciferase activity was analyzed using the Dual-Luciferase Reporter Assay System (Promega, USA). The relative activity of the luciferase was determined by the activity ratio of firefly luciferase to Renilla luciferase.

**Statistical analysis**

The data obtained from three separate experiments are expressed as the mean ± standard error (SE). We analyzed the data with repeated-measures analysis of variance (ANOVA). Comparison between two groups was made using Student’s t-tests. Correlations between two variables were analyzed with Pearson correlation analysis. The data were analyzed using GraphPad Prism 5 (GraphPad Software, San Diego, CA, USA) and SPSS 22.0 (SPSS Inc., Chicago, IL, USA).
Table 1 Association between miR-378a-5p expression and clinicopathological characteristics

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>No. of patients</th>
<th>X ± SD</th>
<th>t</th>
<th>P</th>
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<tr>
<td>Sex</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Male</td>
<td>20</td>
<td>0.38±0.13</td>
<td>0.413</td>
<td>0.682</td>
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<tr>
<td>Female</td>
<td>30</td>
<td>0.36±0.14</td>
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<td>Age, years</td>
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<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;50</td>
<td>24</td>
<td>0.39±0.12</td>
<td>1.307</td>
<td>0.198</td>
</tr>
<tr>
<td>≥50</td>
<td>26</td>
<td>0.35±0.14</td>
<td></td>
<td></td>
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<tr>
<td>Serum AFP, ng/mL</td>
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<td></td>
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<td></td>
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<tr>
<td>≤20</td>
<td>24</td>
<td>0.38±0.15</td>
<td>0.800</td>
<td>0.427</td>
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<tr>
<td>&gt;20</td>
<td>26</td>
<td>0.35±0.11</td>
<td></td>
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<tr>
<td>Serum HBsAg</td>
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<td></td>
<td></td>
<td></td>
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<tr>
<td>Negative</td>
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<td>0.37±0.14</td>
<td>0.314</td>
<td>0.755</td>
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<tr>
<td>Positive</td>
<td>23</td>
<td>0.36±0.12</td>
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<tr>
<td>Tumor size, mm</td>
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<tr>
<td>≤10</td>
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<td>0.42±0.14</td>
<td>3.249</td>
<td>0.002</td>
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<tr>
<td>&gt;10</td>
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<td>TNM stage</td>
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<tr>
<td>0 &amp; I &amp; II</td>
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<td>3.563</td>
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<tr>
<td>III &amp; IV</td>
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<tr>
<td>Distant metastasis</td>
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<tr>
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<tr>
<td>Positive</td>
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<td>Vascular invasion</td>
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<tr>
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<td>32</td>
<td>0.41±0.13</td>
<td>4.160</td>
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<tr>
<td>Positive</td>
<td>18</td>
<td>0.27±0.07</td>
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</table>

Results

Clinicopathological characteristics of HCC patients

Expression of miR-378a-5p in HCC patient tissue was related to tumor size (in mm; P<0.05), vascular invasion (P<0.001), distant metastasis (P<0.05), and TNM stage (P<0.05). The relative miR-378a-5p expression was unassociated with age (P>0.05), sex (P>0.05), serum AFP (P>0.05), or serum HBsAg (P>0.05; Table 1).

Low expression of miR-378a-5p and VEGF in HCC tissues

The HCC tissues had significantly lower miR-378a-5p expression than the matched benign tissue samples (t=9.776, P<0.001; Figure 1A). The HCC tissue samples had significantly increased VEGF protein levels compared with the normal tissue samples (t=18.339, P<0.001; Figure 1B). Interestingly, Pearson correlation analysis showed that miR-378a-5p expression correlated negatively with VEGF protein levels (r^2=0.6568, P<0.001; Figure 1C).

miR-378a-5p mimics inhibited proliferation in HCC cell lines

The HCC cell lines in the miR-378a-5p mimics group had higher miR-378a-5p levels than that in the NC mimics.
Figure 1 miR-378a-5p overexpression and VEGF downregulation in HCC tissues. (A) miR-378a-5p expression in HCC tissues was determined by RT-PCR; (B) VEGF expression in HCC tissues was detected via western blot assays; (C) correlation between miR-378a-5p levels and VEGF in HCC tissues. Data represent the mean ± SE. **, P<0.01 was considered significant. VEGF, vascular endothelial growth factor; HCC, hepatocellular carcinoma.

Figure 2 miR-378a-5p overexpression inhibited Hep3B and SNU-449 cell proliferation. (A) miR-378a-5p expression in Hep3B and SNU-449 cells was determined via RT-PCR; (B,C) cell viability of Hep3B and SNU-449 cells was evaluated via CCK-8 assays; (D,E) clonogenic capacity of Hep3B and SNU-449 cells was assessed via cell colony formation assays. Representative images of the Hep3B and SNU-449 cells by photomicrographs (200×). Data represent the mean ± SE. **, P<0.01 was considered significant. RT-PCR, real-time polymerase chain reaction.
miR-378a-5p inhibited HCC cell migration and invasion

In the Transwell cell migration assays, miR-378a-5p overexpression in Hep3B (t=6.086, P<0.001; Figure 3A) and SNU-449 (t=5.626, P<0.001; Figure 3B) cell lines contributed to significantly decreased cell invasion compared with that in the NC mimics. In the wound-healing assays, downregulated miR-378a-5p promoted both Hep3B (t=5.394, P<0.001; Figure 3C) and SNU-449 (t=9.311, P<0.001; Figure 3D) cell migration, while the number of invaded cells was markedly lower when miR-378a-5p mimics were transfected into the cells.

miR-378a-5p reduced VEGF expression in HCC cell lines

The miR-378a-5p mimics had significantly decreased VEGF protein expression compared with the NC mimics (Hep3B: t=11.490, P<0.001; Figure 4A and B; SNU-449: t=13.876, P<0.001; Figure 4C and D). The cell proliferation assay showed that the NC mimics had higher cell proliferative capacity than did the miR-378a-5p mimics in Hep3B (F=138.776, P<0.001; Figure 2B) and SNU-449 (F=156.889, P<0.001; Figure 2C) cells. The colony formation assays showed that the miR-378a-5p mimics had reduced cell colony formation ability compared with the NC mimics (Hep3B: t=16.537, P<0.001; Figure 2D; SNU-449: t=21.874, P<0.001; Figure 2E).
miR-378a-5p reduced VEGF expression in HCC cell lines. (A,B,C) VEGF expression in Hep3B and SNU-449 cells was examined via western blot assays; (D,E) correlation between miR-378a-5p levels and VEGF in Hep3B and SNU-449 cell lines; (F) wild-type miR-378a-5p target sequences of VEGF mRNA 3′-UTR; (G,H) miR-378a-5p repressed the activity of pGL3-VEGF-WT but not pGL3-VEGF-MUT in Hep3B and SNU-449 cells. Data represent the mean ± SE. **, P<0.01 was considered significant. VEGF, vascular endothelial growth factor; HCC, hepatocellular carcinoma.

t=10.134, P<0.001; Figure 4A and C). The miR-378a-5p expression and VEGF protein levels were negatively correlated (Hep3B: r²=0.6482, P<0.001; Figure 4D; SNU-449: r²=0.6014, P<0.001; Figure 4E). TargetScan analysis showed that miR-378a-5p targeted VEGF (Figure 4F), and luciferase assays revealed that miR-378a-5p repressed the activity of pGL3-VEGF-WT but not pGL3-VEGF-MUT in Hep3B (t=10.134, P<0.001; Figure 4G) and SNU-449 (t=10.134, P<0.001; Figure 4H) cells.

Discussion

Here, we report that miR-378a-5p expression was commonly downregulated in HCC tissues and correlated with HCC progression. Upregulation of miR-378a-5p expression reduced Hep3B and SNU-449 cell viability, proliferation, invasion, and migration. We also found that miR-378a-5p targeted and suppressed VEGF signaling, and the two were negatively correlated. The present results show that miR-378a-5p has a tumor-suppressive role in HCC progression by targeting VEGF signaling, which has previously been rarely reported.

Evidence has shown that miR-378a-5p is abnormally regulated in renal cell carcinoma (17) and stage II colon cancer (15). Here, miR-378a-5p expression was low in the HCC tissues and was related to TNM stage, distant
metastasis, and vascular invasion. Our findings show that miR-378a-5p acts as an anti-tumor factor and might play a key role in HCC prognosis, metastasis, and progression.

Studies have shown the crucial roles of miR-378a-5p in impeding cell viability, invasion, proliferation, and migration (17,22). Our results showed that transfection of miR-378a-5p mimics markedly decreased cell vitality and colony formation. Furthermore, miR-378a-5p expression led to reduced HCC cell invasion and migration, indicating that miR-378a-5p negatively regulated HCC cell progression. miR-378a-5p’s antitumor effect in this study is consistent with that in Pan et al.’s study, where, in renal cell carcinoma, miR-378a-5p was described as playing a tumor-suppressive role and was associated with a good prognosis (17). Another study showed that miR-378a-5p overexpression promoted vascular smooth muscle cell migration and proliferation (23). Excess miR-378a-5p has also been correlated with breast cancer tumorigenesis in vivo (18). Thus, whether the antitumor effects induced by miR-378a-5p are bidirectional and concentration-dependent require further study.

Cheng et al. found that the VEGF signaling pathway promoted HCC cell invasion, migration, and angiogenesis in vitro (24). Here, HCC tissues and cell lines had significantly increased VEGF protein levels, and VEGF expression in the Hep3B and SNU-449 cells was decreased by miR-378a-5p overexpression. Expression of miR-378a-5p was negatively associated with Hep3B and SNU-449 HCC cells. Moreover, we confirmed VEGF as the target of miR-378a-5p in HCC cells. These data demonstrated that miR-378a-5p produced antitumor effects by inhibiting VEGF. Research shows that the main mechanism of VEGF in tumor invasive and metastatic progression is that VEGF directly stimulates vascular endothelial cell differentiation, proliferation and migration and degrades vascular basement membranes, which promotes shedding of cancer cells into blood vessels, then transfers them to the adjacent fibrin (25,26). VEGF promotes formation of new lymphatic vessels around and inside tumors as well as original lymphangitic hyperplasia and increased vessel diameter. VEGF also facilitates tumor cells to lymphatic metastasis (27). However, cyclin-dependent kinase 6 is also a target signaling pathway of the tumor suppressor functions of miR-378 (19), which will be explored in future research.

Conclusions

In summary, miR-378a-5p inhibited tumors in HCC and in two HCC cell lines, and low expression of miR-378a-5p is a potential poor prognostic marker in patients with HCC. miR-378a-5p affects the migration, invasion, and proliferation of HCC cells by targeting the VEGF signaling pathway. With biological and clinical implications in HCC, miR-378a-5p has potential value in HCC therapy and may be a good prognostic and confirmatory diagnostic marker in HCC tumorigenesis.

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Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The Sichuan Academy of Medical Sciences and Sichuan Provincial People’s Hospital Ethics Committee approved all study protocols.

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