Abstract. MicroRNAs (miRNAs) represent a class of small RNAs that participate in the regulation of tumor progression. However, the identification of functional miRNAs in tumors has not been thoroughly elucidated. In the present study we aim to investigate the impact of altered miR-2053 expression in hepatocellular carcinoma (HCC) cells. The results of the present study demonstrated that miR-2053 overexpression inhibited cell proliferation, migration and invasion, and promoted apoptosis in a HCC cell line, while miR-2053 knockdown induced the opposite cellular phenotypic changes. Mechanistically, it was found that overexpression of miR-2053 resulted in the downregulation of the phosphoinositide 3-kinase (PI3K) and Wnt/β-catenin signaling pathways, which are aberrantly expressed in HCC. Collectively, the results indicate that miR-2053 serves as a tumor suppressor with a crucial role in inhibiting the proliferation, migration and invasion of HCC via targeting the PI3K and Wnt/β-catenin signaling pathways. These data indicate a potential application of miR-2053 in cancer therapy.

Introduction

Hepatocellular carcinoma (HCC) accounts for up to 90% of all primary liver cancer worldwide, and has been reported to cause >662,000 mortality cases worldwide annually, particularly in developing countries (1,2). Despite improvements in the treatment of liver cancer, the survival rate after 5 years is <30% due to its high recurrence and metastasis rate (3,4). Although the treatment efficacy for HCC is improving, the lack of biomarkers for early diagnosis and effective therapeutic targets results in consistently poor curative rates. Consequently, a better understanding of the mechanisms and potential biomarkers are warranted to improve HCC patient treatment.

MicroRNAs (miRNAs) are small, single-stranded, non-coding functional RNA molecules of ~22 nucleotides that are widely present in eukaryotic cells (5,6). MiRNAs have been identified in various cancer types and are involved in cancer development and progression, acting as oncogenes or tumor suppressors (7,8). Previous studies show that miRNAs are involved in liver disease and progression of liver cancer. For example, miR-324-3p promotes HCC growth, however, miR-148b serves as a tumor suppressor in HCC by inhibiting proliferation and invasion (9). In addition, several miRNA signatures are related to chronic hepatic infection, cirrhosis, and steatosis (10,11). The specific expression of certain miRNAs has been found to have biological behavior in terms of tumor aggressiveness, metastatic potential and even responsiveness to treatment. Therefore, any abnormal expression of miRNA molecules is thought to be associated with hepatocarcinogenesis. Accumulating evidence suggests that multiple miRNAs are aberrantly expressed in and associated with various processes of cellular carcinoma development. miR-2053 is a miRNA molecule located at 8q23.3 in the human genome. It has been reported that miR-2053 may induce proliferation of adult cardiomyocytes (12). However, to the best of our knowledge, the function of miR-2053 in HCC has not been explored.

Hepatocarcinogenesis is a complex, multistep process in which several signaling cascades are altered, leading to the development of a heterogeneous biological tumor. The phosphoinositide 3-kinase (PI3K)/serine/threonine protein kinase (AKT)/Mammalian target of rapamycin (mTOR) and the Wnt/β-catenin pathways serve important roles in HCC proliferation and cell cycle progression (13,14). Studies have shown that PI3K signaling is activated in 30-50% of HCC cases, and the downstream ribosomal protein S6 (RPS6) is also activated in 50% of HCC patients (15). Previous studies have demonstrated that the Wnt/β-catenin signaling pathway...
plays a critical role in proliferation and cell cycle progression of HCC cells (16,17). miR-1247-5p functions as a tumor suppressor by activating Wnt3 (18). It was also demonstrated that miR-138 modulates prostate cancer cell invasion and migration via the Wnt/β-catenin pathway (19). In the present study, the association of miR-2053 with the PI3K/AKT/mTOR and Wnt/β-catenin pathways in HCC cells was explored.

The effects and mechanisms of miR-2053 on the progression of HCC were investigated. The data demonstrated that overexpression of miR-2053 inhibited the proliferation, migration and invasion of HCC cells and induced cell apoptosis, which may be through inhibiting the activation of the AKT/mTOR and Wnt/β-catenin pathways. These findings reveal the function and regulatory mechanisms of miR-2053 in human HCC.

Materials and methods

Cell lines and cell culture. The human HCC cell line Huh7 was obtained from Cell Bank of Chinese Academy of Sciences (Shanghai, China) and routinely maintained in high-glucose Dulbecco's Modified Eagle's Media (DMEM) supplemented with 10% fetal bovine serum (FBS; Gibco; Thermo Fisher Scientific, Inc., Waltham, MA, USA), 100 U/ml penicillin, and 100 mg/ml streptomycin. All cell lines were cultured at 37°C in a humidified atmosphere containing 5% CO₂.

Plasmid transfection. Cells were seeded in a 24-well plate and after the cells reached 80% confluence, transient transfection of miRNA plasmids was carried out using Lipofectamine® 2000 (Invitrogen; Thermo Fisher Scientific, Inc.), according to the manufacturer's protocol. Briefly, the miR-2053 or siRNA-miR-2053 sequences were cloned into the pCMV-MIR vectors (Guangzhou Ribobio Co., Ltd., Guangzhou, China) (20). Lipofectamine 2000 (10 µl) was added to 250 µl DMEM without serum and incubated for 5 min at room temperature. The plasmids [pCMV-MIR-miR-2053 (miR-2053) or pCMV-MIR-si-miR-2053 (si-miR-2053), 50 nM] were added without serum and incubated for 5 min at room temperature. The cells were prepared by removing medium and washing twice with PBS followed by addition of 0.5 ml of medium without serum. The complexes (500 µl) were added to each well, and the plate was mixed gently by rocking back and forth, and incubated at 37°C in a humidified atmosphere containing 5% CO₂ for 24 h prior to evaluating gene expression. The medium was changed 6 h after transfection. Cells transfected with an empty pCMV-MIR vector were used as a negative control group (pNC) for the miR-2053 upregulation group, cells transfected with pCMV-MIR-siRNA (non-coding siRNA) vector was performed as a negative control (siNC) for si-miR-2053, and cells without any treatment was used as blank control group (Con). The mature sequence of miR-2053 is 5'-GUGUUAUAUAACCUCUAUUAC-3'; the siRNA targeting miR-2053 is 5'-GUAAAUAGAGGUUAUAUAAAC-3';

RNA isolation and reverse transcription-quantitative polymerase chain reaction (RT-qPCR). Total RNA was extracted using an RNA extraction kit (CoWin Bioscience Co., Ltd., Beijing, China) and total RNA was converted to cDNA using a miRNA cDNA Synthesis Kit (CoWin Bioscience Co., Ltd.). miRNA levels were examined using a miRNA qPCR detection kit (CoWin Bioscience Co., Ltd.). The thermocycling conditions were set at 95°C for 5 min, followed by 40 cycles of 95°C for 10 sec, 60°C for 30 sec, and 72°C for 1 sec. Each reaction was performed in triplicate. The expression of miR-2053 was calculated using the 2^{-ΔΔCq} method (21). Three independent experiments were performed to analyze the relative gene expression. The primers were as follows: MiR-2053, forward primer, 5'-CTGCGTTCGCGAGCA-3'; reverse primer, was obtained from the miRNA qPCR detection kit; U6, forward primer, 5'-GUA AAU UUU AC-3'; the siRNA targeting miR-2053 is 5' -GUA AAU UUU AC-3'; the siRNA targeting miR-2053 is 5' -GUA AAU UUU AC-3'; reverse primer, 5'-AAC GCT TCA CGA ATT TGC GT-3'.

Western blot analysis. After transfection for 48 h, total cellular protein was extracted using radioimmunoprecipitation (RIPA) lysis buffer (CoWin Biosciences Co., Ltd.). The lysates containing equal amounts of protein (20 µg) were loaded onto 10% SDS-PAGE gels (Beyo) and transferred onto polyvinylidene fluoride membranes (EMD Millipore, Billerica, MA, USA). The membrane was blocked with 5% skimmed milk for 1 h and incubated with primary antibodies overnight at 4°C, followed by 1 h of incubation with horseradish peroxidase-conjugated secondary antibodies (anti-rabbit IgG, cat. no. SA00001-2; anti-mouse IgG, cat. no. SA00001-1; both 1:5,000; both Proteintech Group, Inc., Chicago, IL, USA). Finally, the proteins were detected by enhanced chemiluminescence using the ECL Plus kit (ProteinTech Group, Inc.). Band densities were analyzed using the QUANTITY ONE 1-D software (version 4.6.7; Bio-Rad Laboratories, Inc., Hercules, CA, USA). The following primary antibodies were used for analysis: Anti-caspase-9 (cat. no. ab219590; 1:1,000), anti-AKT (cat. no. ab32505; 1:1,000), anti-phosphorylated AKT (cat. no. ab81283; 1:1,000), anti-mTOR (cat. no. ab2732; 1:1,000), anti-phosphorylated mTOR (cat. no. ab3154; 1:100), anti-Wnt3 (cat. no. ab32249; 1:10), anti-β-catenin (cat. no. ab32572; 1:1,000, all Abcam, Cambridge, UK), anti-E-cadherin (cat. no. 3195; 1:1,000; Cell Signaling Technology, Danvers, MA, USA), anti-Cyclin D1 (cat. no. 60186-1-lg; 1,1000), anti-p70 (cat. no. 14485-1-AP; 1,1000), anti-B-cell lymphoma 2 (BCL2; cat. no. 60178-1-lg; 1,1000), anti-BCL-2-associated X protein (BAX; cat. no. 60267-1-lg; 1,1000) and anti-cleaved caspase-3 (cat. no. 25546-1-AP; 1,1000) and GAPDH (cat. no. 60004-1-lg; 1,5000; all Proteintech Group, Inc.), which served as the loading control.

Cell proliferation assay. After 24 h of transfection, the cells (1x10⁴ cells/ml) were plated onto 96-well plates in 100 µl of complete medium. Cell Counting Kit-8 (CCK-8; Solarbio Science & Technology Co., Ltd., Beijing, China) was used according to the manufacturer's protocol. The plates were incubated at 37°C for 1.5 h, and the absorbance at 450 nm was measured. The proliferation rates were determined at 0, 24, 48 and 72 h post-transfection.

Clone formation assay. After 24 h of transfection, cells (~500) were planted in a 6-cm dish in 5 ml of medium and cultured.
until visible clones formed. After staining with 0.1% crystal violet for 30 min at room temperature, the colonies were counted under a microscope (magnification x4; Nikon Corporation, Tokyo, Japan) and imaged with a camera.

Cell migration and invasion assay. Cell invasion and migration assays were performed 24 h after transfection using a Transwell system (EMD Millipore, Billerica, MA, USA). Migration was assessed using uncoated transwells, and invasion was investigated using transwells coated with Matrigel (BD Biosciences, San Jose, CA, USA). A total of 1x10^5 cells transfected for 24 h in serum-free medium were added to the top chamber of the transwell. The bottom chamber was filled with medium containing 10% FBS. After 24 h incubating at 37°C, the cells on the upper surface of the membrane were gently removed with a cotton swab, and the membranes were washed three times with PBS, fixed and stained with 0.1% crystal violet for 5 min at room temperature. The cells were imaged and counted in 5 random fields using a light microscope (Nikon Corporation; magnification x100).

Flow cytometry analysis. Cell apoptosis was measured using the fluorescein isothiocyanate (FITC) Annexin V Apoptosis Detection Kit (4A Biotech Co., Ltd., Beijing, China), according to the manufacturer's protocol. Briefly, after transfection of the cells for 24 h, the medium was removed and the cells were incubated with serum-free medium for 24 h. The cells were digested using trypsin, washed with ice-cold PBS, centrifuged at 200 x g for 5 min, washed twice with ice-cold PBS and then stained with FITC-Annexin V and propidium iodide (PI). Apoptotic cells were detected using a flow cytometer (EPICS, XL-4; Beckman Coulter, Inc., Brea, CA, USA) and analyzed using FlowJo software (v7.6.1; FlowJo LLC, Ashland, OR, USA).

Statistical analysis. All statistical analyses were performed using SPSS 18.0 software. (SPSS, Inc., Chicago, IL, USA) The data were expressed as mean ± standard deviation. The Student's t-test was used to determine statistically significant differences between the two groups, and one-way analysis of variance was used to determine the statistical significance across multiple groups followed by Least Significant Difference post hoc comparison. P<0.05 was considered to indicate a statistically significant difference. All experiments were performed in triplicate.

Results

miR-2053 expression in Huh7-transfected cells. To increase or decrease miR-2053 expression, pCMV-MIR-miR-2053 and pCMV-MIR-si-miR-2053 plasmids were transfected into Huh7 cells, a human HCC cell line. RT-qPCR analysis revealed that miR-2053 expression exhibited no significant difference in control, pNC and siNC groups. miR-2053 expression was significantly increased in cells transfected with pCMV-MIR-miR-2053 plasmid (miR-2053 group) compared with that in the pNC group, and decreased by siRNA-miR-2053 (si-miR-2053 group) at 24 and 48 h after transfection compared with that in the siNC group (Fig. 1A and B; P<0.05).

miR-2053 inhibits proliferation of Huh7 cells. Proliferation serves an important role in the development of a tumor (22). CCK-8 assay was used to study the role of miR-2053 in the proliferation of Huh7 cells. The proliferation rate was measured at 72 h. Following transfection for 72 h, proliferation of Huh7 cells was significantly inhibited by miR-2053 overexpression in the miR-2053 group compared with that in the pNC group, and was promoted in the si-miR-2053 group compared with that in the siNC group (Fig. 2A). Additionally, a plate colony assay showed that cells transfected with miR-2053 formed fewer and smaller colonies compared with that in the pNC group in Huh7 cells, and si-miR-2053 increased the colony number compared with the siNC group (Fig. 2B and C). These results demonstrate that overexpression of miR-2053 inhibited the proliferative potential, and miR-2053 knockdown promoted cell proliferation of this HCC cell line.

miR-2053 inhibits Huh7 cell migration and invasion. Tumor invasion refers to the process of tumor cells destroying the surrounding normal tissue structure and detaching from the primary tumor (23). It was found that the number of cells successfully invading through a Matrigel-coated membrane was inhibited significantly in the miR-2053 group compared with that in the pNC group. Additionally, the number of invaded cells increased in the si-miR-2053 group compared...
Figure 2. miR-2053 inhibits the proliferation of Huh7 cells. (A) Huh7 cell proliferation measured by Cell Counting Kit-8 assay at 0, 24, 48 and 72 h after transfection. (B) Representative images and (C) quantification of the number of colonies resulting from the colony formation assay, 24 h following transfection. *P<0.05, miR-2053 group vs. pNC group, si-miR-2053 group vs. siNC group; n=3. Con, untransfected group; pNC, cells transfected with pCMV-MIR empty vector; miR-2053, cells transfected with pCMV-MIR- miR-2053 vector, miR-2053 overexpression group; siNC, cells transfected with pCMV-MIR-siRNA (non-coding siRNA) vector; si-miR-2053, cells transfected with pCMV-MIR-si-miR-2053, miR-2053 knockdown group.

Figure 3. miR-2053 inhibits Huh7 cell migration and invasion in vitro. After 24 h of transfection, Transwell and Matrigel assays were performed to assess cell migration and invasion (A) Magnification, x100. Quantification of the number of (B) invasive and (C) migratory cells derived from 5 randomly chosen fields. *P<0.05, miR-2053 group vs. pNC group, si-miR-2053 group vs. siNC group; n=3. Con, untransfected group; pNC, cells transfected with pCMV-MIR empty vector; miR-2053, cells transfected with pCMV-MIR- miR-2053 vector, miR-2053 overexpression group; siNC, cells transfected with pCMV-MIR-siRNA (non-coding siRNA) vector; si-miR-2053, cells transfected with pCMV-MIR-si-miR-2053, miR-2053 knockdown group.
with that in the siNC group (Fig. 3A and B; P<0.05). The number of cells migrating through non-coated membranes also decreased markedly in miR-2053-overexpressing Huh7 cells, and increased significantly in si-miR-2053 group (Fig. 3A and C; P<0.05). These results demonstrate that miR-2053 markedly suppresses the migration and invasion of Huh7 cells.

**miR-2053 promotes apoptosis in Huh7 cells.** Apoptosis plays an important role in the process of tumor growth (24). The effect of miR-2053 overexpression or knockdown on apoptosis was investigated in Huh7 cells. As shown in Fig. 4, 16.69% of cells overexpressing miR-2053 were positive for Annexin V and considered apoptotic. Of the cells transfected with si-miR-2053, 8.00% were apoptotic, while the siNC group contained 11.00% apoptotic cells (Fig. 4A and B). The mechanisms of miR-2053-induced apoptosis was further determined by analyzing the expression of 4 apoptosis-related proteins including BCL2, BAX, caspase-9 and activated caspase-3 (caspase-3-p17). BCL2 is an anti-apoptotic protein and BAX is a pro-apoptotic protein, while caspase-9 and activated caspase-3 also play key roles in the process of apoptosis (25,26). The results showed that overexpression of miR-2053 downregulated the level of BCL2 and upregulated BAX, caspase-9 and caspase-3-p17.

**miR-2053 inhibits the PI3K signaling pathway.** As shown in Fig. 5, the protein expression levels of total AKT, total mTOR, cyclin D1 and p70 decreased significantly in miR-2053-overexpressing Huh7 cells, and increased significantly in si-miR-2053 group (Fig. 5A and B; P<0.05). These results suggest that miR-2053 may inhibit the PI3K signaling pathway, which is involved in cell proliferation and survival.

**miR-2053 inhibits the Wnt/β-catenin signaling pathway.** As shown in Fig. 6, the protein expression levels of Wnt3, β-catenin and epithelial-cadherin decreased markedly in miR-2053-overexpressing Huh7 cells, and increased significantly in si-miR-2053 group (Fig. 6A and B; P<0.05). These results suggest that miR-2053 may inhibit the Wnt/β-catenin signaling pathway, which is involved in cell proliferation, migration and invasion.

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Figure 4. miR-2053 promotes apoptosis in Huh7 cells. (A) After 24 h of transfection, cell apoptosis was measured by flow cytometry after incubation with FITC-Annexin V and PI solution. (B) Percentage of cells positive for Annexin V and PI. (C) Protein expression levels of BCL2, BAX, caspase-9 and cleaved caspase-3. *P<0.05, miR-2053 group vs. pNC group, si-miR-2053 group vs. siNC group; n=3. Con, untransfected group; pNC, cells transfected with pCMV-MIR empty vector; miR-2053, cells transfected with pCMV-MIR-miR-2053 vector, miR-2053 overexpression group; siNC, cells transfected with pCMV-MIR-siRNA (non-coding siRNA) vector; si-miR-2053, cells transfected with pCMV-MIR-si-miR-2053, miR-2053 knockdown group; FITC, fluorescein isothiocyanate; PI, propidium iodide; BCL2, B-cell lymphoma 2, BAX, BCL-2-associated X protein.

Figure 5. miR-2053 inhibits the PI3K signaling pathway. (A) Protein expression levels of total and p-AKT, total and p-mTOR, cyclin D1 and p70. (B) Ratios of p-AKT/AKT and p-mTOR/mTOR, and the expression of cyclin D1 and p70. n=3, *P<0.05 vs. pNC control group. pNC, cells transfected with pCMV-MIR empty vector; miR-2053, cells transfected with pCMV-MIR-miR-2053 vector, miR-2053 overexpression group; p, phosphorylated; AKT, serine/threonine protein kinase; mTOR, Mammalian target of rapamycin.

Figure 6. miR-2053 inhibits the Wnt/β-catenin signaling pathway. (A) Protein expression levels of Wnt3, β-catenin and epithelial-cadherin. (B) Quantitative analysis of western blot results. N=3, *P<0.05 vs. pNC group. pNC, cells transfected with pCMV-MIR empty vector; miR-2053, cells transfected with pCMV-MIR-miR-2053 vector, miR-2053 overexpression group; E-cad, E-cadherin.
miR-2053 overexpression inhibits activation of the PI3K/AKT signaling pathway in Huh7 cells. The PI3K signaling pathway serves a central role in the progression of HCC, participating in proliferation and metastasis (27). According to our previous results, it was demonstrated that miR-2053 overexpression decreased cell proliferation, migration and invasion. This effect may be mediated through PI3K signaling. Therefore, the impact of miR-2053 overexpression on the PI3K/AKT signaling pathway was assessed. It was found that the ratio of phosphorylated (p) AKT to total AKT, the ratio of p-mTOR to total mTOR, and the levels of p70 and Cyclin D1 were downregulated in the miR-2053 group (Fig. 5). These results highlight an association between miR-2053 and the PI3K/AKT signaling pathway in Huh7 cells.

miR-2053 inhibits activation of the Wnt/β-catenin signaling pathway in Huh7 cells. Previous studies have found that miRNAs promote HCC invasion and metastasis via the Wnt/β-catenin signaling pathway (28,29). The results of the present study demonstrated that overexpression of miR-2053 resulted in a reduction in Wnt3 protein levels. It was also revealed that β-catenin was decreased and E-cadherin was increased in miR-2053-overexpressing Huh7 cells compared with that in the pNC group (Fig. 6). These results indicate that miR-2053 may impact HCC progression through the Wnt/β-catenin signaling pathway.

Discussion

Hepatocellular carcinoma is the third leading cause of cancer-associated mortality and the incidence of viral infections is increasing worldwide (30). Research has shown that miRNAs are abnormally expressed in HCC, and are involved in the regulation of tumor development and malignant changes (31). Certain miRNAs serve as promoters in hepatic tumorigenesis. Conversely, several miRNAs function as tumor suppressors. For example, miR-221 blockage prompts HCC survival (32,33), whereas miR-148b suppresses cell proliferation and invasion in HCC by targeting the Wnt/β-catenin pathway (34). Studies have found that miR-132 inhibits cell proliferation, invasion and migration of HCC (35). These findings indicate that miRNAs are intimately involved in the progression of HCC malignancy.

In the present study, it was shown that the overexpression of miR-2053 significantly inhibited proliferation, colony formation, cell migration and invasion, and induced cell apoptosis in a human HCC cell line. Knockdown of miR-2053 induced the opposite changes, increasing proliferation and migration, and reducing apoptosis. These results suggest that miR-2053 may function as a tumor suppressor in HCC. The molecular mechanisms underlying the role of miR-2053 in HCC cells remains unclear.

Previous studies have shown that the PI3K/AKT pathway is one of the most altered oncogenic pathways in tumor development, including HCC (36,37). The downstream effector, mTOR serves an important role in hepatitis and HCC development. Previous studies have demonstrated that AKT/mTOR serves an important role in the cell cycle of tumor cells (38,39). Therefore, the phosphorylation of AKT and mTOR, as well as expression of cell cycle regulators Cyclin D1 and p70, was examined in the present study. The results demonstrated that miR-2053 overexpression significantly reduced the levels of phosphorylated AKT and mTOR, and the levels of Cyclin D1 and p70 in Huh7 cells, suggesting that miR-2053 may inhibit HCC proliferation through regulating the AKT/mTOR signaling pathway.

Wnt3 belongs to the Wnt1 class of ligands and stimulates the canonical Wnt/β-catenin pathway (18,40). Wnt/β-catenin is involved in HCC progression. TRIM37 overexpression promotes cell migration and metastasis in HCC by activating Wnt/β-catenin signaling (41). It has also been found that the downregulation of miR-200a induced epithelial-mesenchymal transition phenotypes and influenced proliferation, migration and invasion of SGC790 and U251 cells through targeting the β-catenin pathway (42). Here, miR-2503 overexpression inhibited the expression of Wnt3 and β-catenin, and increased E-cadherin levels, downstream molecules in the Wnt signaling pathway. Taken together, these data demonstrated that miR-2053 may function as a tumor suppressor in HCC development by regulating cell migration and invasion.

In conclusion, it was demonstrated that miR-2053 overexpression inhibited Huh7 cell proliferation, colony formation, cell migration and invasion, and induced cell apoptosis. Additionally, the PI3K/AKT/ and Wnt/β-catenin signaling pathways were regulated by miR-2053 in this HCC cell line. Therefore, miR-2053 may be a potential target for novel invasive HCC treatment.

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Availability of data and materials

All data generated or analyzed during this study are included in this published article.

Authors’ contributions

TS, KM, CZ, JY, JL participated in the experiments, data analysis, manuscript design and writing. All authors have read and approved this manuscript.

Ethics approval and consent to participate

Not applicable.

Patient consent for publication

Not applicable.
Competing interests

The authors declare that they have no competing interests.

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