Mig-6 Promotes the Apoptosis and Inhibits of the Flux of Autophagy in HCC Cell Lines Through Modulating Mig-6/miR-193a-3p/TGF- β 2 Axis

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Abstract

Background: Mitogen-inducible gene 6 (*Mig-6*) is a tumor suppressor gene that plays an important role in many types of cancers by interacting with EGFR. Thus far, little is known about the molecular mechanism of *Mig-6* in hepatocellular carcinoma (HCC). Also, the relationship between *Mig-6* and miRNAs needs to be elucidated. Therefore, this study first aimed to find whether *Mig-6* could promote apoptosis and inhibition of the flux of autophagy by its downstream miRNA in HCC cell lines.

Methods: Two cell lines, HepG2 and HLE, were used in this study. *Mig-6* overexpression and knockdown models, miR-193a mimics and inhibitors models, were established. MiRNA microarray profiling was used to verify *Mig-6* regulated miRNA. Real-time PCR, Western blot analysis were carried out to detect RNA and protein expression. Evaluation of fluorescent LC3 puncta and cell flow cytometer assay were used to test the autophagy and apoptosis respectively.

Results: *Mig-6* induced the apoptosis and reduced the autophagy of HCC cell lines. MiR-193a-3p is a *Mig-6*-regulated miRNA in *Mig-6*

overexpression model, and miR-193a-3p affected the apoptosis and autophagy of HCC cells by regulating the expression of TGF- β 2. Additionally, the relationship between *Mig-6* and transforming growth factor TGF- β 2 was explored in depth for the first time.

Conclusion: These findings revealed an important regulatory axis *Mig-6*/miR-193a-3p/TGF- β 2 in the apoptosis and autophagy of HCC cells, providing a novel insight into the therapeutic target in HCC.

Background

Hepatocellular carcinoma (HCC) is the most prevalent type of primary liver cancer accounting for approximately 70–90% of liver cancer cases. In terms of morbidity, it ranks the second most dominating reason for cancer-related deaths globally [1, 2]. HCC is associated with high morbidity and mortality. Chronic hepatitis, especially infection induced by hepatitis, is closely related to the occurrence of HCC. Elucidating the pathogenesis of HCC can help in the early diagnosis and treatment of this disease. The mitogen-inducible gene 6 (*Mig-6*), also known as ERBB receptor feedback inhibitor 1, is a transcriptionally induced EGFR inhibitor that is also a tumor suppressor [3, 4]. *Mig-6* is a cytoplasmic protein that binds to the EGFR dimer interface, locking it into a catalytically inactive conformation, as well as inducing internalization and degradation of EGFR via a lysosomal mechanism [5]. The expression of *Mig-6* has been demonstrated to be suppressed in liver cancer tissues, leading to increased EGFR-AKT signaling and enhanced cell migration [6]. *Mig-6* has also been reported to regulate the proliferation of liver cancer cells through the classic EGFR pathway. However, some of the regulatory effects of *Mig-6* on HCC cells, such as the effect on HCC autophagy, cannot be explained by the classic EGFR pathway, forcing the exploration of new functions of *Mig-6*.

MicroRNAs (miRNAs) are small noncoding RNA molecules with 19–22 nucleotides that regulate the protein expression[7] They regulate the gene expression at the posttranscriptional level by decreasing the stability or inhibition of translation of messenger RNAs (mRNAs) through targeting the 3'-untranslated region of mRNAs[8].

Increasing or decreasing the levels of specific miRNAs can result in aberrant protein expression, leading to the critical change in cell functions; for example, proliferation, cell cycle, apoptosis, and invasion. Many studies showed that miRNAs participated in the biological behavior of HCC. For instance, miR-139-5p influenced HCC cell invasion and proliferation capacities by decreasing the expression of SLITRK4[9]. MiR-133b promotes cell proliferation and metastasis in HCC by regulating SF3B4[10]. The downregulated expression of miR-193a-3p was reported in many types of cancers, including non-small-cell lung cancer (NSCLC)[11],colorectal cancer[12]and breast cancer[13]. MiR-193a-3p was also related to the proliferation, migration, invasion, and apoptosis of liver cancer cells[14–16]. Therefore, the potential mechanism of miR-193a-3p in HCC needs to be explored.

In this study, the enhanced expression of *Mig-6 in vitro* was found to not only weaken the flux of autophagy but also promote the apoptosis of HCC cells. The miRNA expression profiling chips were adequately applied, and three bioinformatics algorithms were used to screen out its potential downstream target miR-193a-3p and transforming growth factor (TGF)- β 2. Afterward, the impact of miR-193a-3p on the apoptosis and autophagy of HCC was studied and evaluated. Finally, the indirect control of *Mig-6* and TGF- β 2 was studied in depth for the first time, providing a novel insight into the *Mig-6*-based therapeutic approach to treat human HCC.

Materials And Methods

Cell lines

Human hepatocellular cancer cell lines HepG2, Hep3B, PLC/PRF/5, HLE, and HuH7were maintained in Dulbecco's modified Eagle's medium (Biological Industries) supplemented with 10% fetal calf serum (Biological Industries), 10 IU/mL penicillin (Sigma, St.Louis, MO), and 100 mg/mL streptomycin (Sigma).Cells were grown on sterile T25 cell culture flask (Corning) and were passaged every 2–3 d using 0.25% trypsin (Biological Industries).

Transfecton of expression plasmids,RNAi, miRNA mimics,and inhibitors

The cells were seeded in 6-well plates 24 h prior to the experiment. Plasmids control vector (pcDNA3) and Mig-6 overexpression vector (Takara Biotechnology, Dalian, China),(TGF- β 2over expression vector (OriGene Technologies Inc, Rockville, MD, USA),RNAi for *Mig-6* and non-targeting siRNA (GenePharma, Shanghai, China), miR-193a mimics, miR-193a inhibitors,mimic control,and inhibitor

control (RIBOBIO, Guangzhou, China) were all using Lipofectamine 3000 (Invitrogen, Carlsbad, CA, USA) according to the manufacturer's protocol. The mRNA and protein levels were assessed 48 h following transfection.

Quantitative real-time PCR

Assays to quantify miR-193a-3p were performed using TaqMan miRNA probes and TaqMan® MicroRNA Assays (Applied Biosystems, Foster City, CA, USA) according to the manufacturer's instructions. The reactions as follows: 95 °C for 5 min, 40 cycles of 95 °C for 15 s, 60 °C for 1 min. Quantitative the TGF-β2 mRNA levels was performed using SsoFast™ EvaGreen® Supermix (Bio-rad). The reactions as follows: 95 °C for 30 s, 40 cycles of 95 °C for 5 s, 60 °C for 20 s. Light Cycler® 480 (Roche) was used to perform the real-time PCR. The primer sequences are provided below:

TGF-β2 (forward) CATCCCGCCCACTTTCTAC;

TGF-β2 (reverse) TCCGTTGTT CAGGCACTCT;

Actin (forward) ATAGCACAGCCTGGATAGCAACGTAC;

Actin (reverse) CACCTTCTACAATGAGCTGCGTGTG.

Experiments were repeated in triplicate.

Western blot analysis.

Total proteins from cell lines were extracted in a lysis buffer (CAT.78510) (Thermo Fisher Scientific, Rockford, IL) and quantified using Bradford method (CAT.23226) (Thermo Fisher Scientific). Sixty micrograms of protein were separated by 12% SDS-PAGE (Bio-Rad, USA). After transferring by Trans-Blot® Turbo™ (Bio-Rad, USA), the polyvinylidene fluoride (PVDF) membranes (Millipore, Billerica, MA, USA) were incubated overnight at 4 °C with the following antibodies *Mig-6* (1: 1000, CAT.11630-1-AP) (protein tech), anti-P62 (1:1000, CAT.55274-1-AP) (protein tech), anti-LC3b (1:2000, CAT.NB100-2220) (Novus Biologicals USA), anti-TGF-β2 (1:1000, CAT.ab36495) (Abcam), β-actin (1:1000, CAT.20536-1-AP) (protein tech). After incubation with peroxidase-coupled anti-mouse or rabbit IgG (Zhongshan jinqiao, China) at 37 °C for 2 hours. An enhanced chemiluminescence (ECL) detection system (Bio-Rad, USA) was used to visualize signals following standard protocols. β-actin served as an endogenous protein control for normalization.

Cell apoptosis experiments.

Apoptosis was detected using an Annexin V-FITC/PI double staining kit (Dojindo, Gaithersburg, MD, USA). Cells that incubated in 12-well plates were collected by Accutase (Multi sciences, China) after 48 h transfection and washed twice with cold phosphate-buffered saline (PBS) by gentle shaking. Cells were then resuspended and added to binding buffer (1X); cell density was adjusted to 1 × 10⁶ cells /ml. In the dark, 5 μl Annexin V-FITC and 5 μl PI Solution were added to the cell suspension volume of 100 μl and incubated for 15 min at room temperature before the addition of 400 μl binding buffer (1X). Ten thousand

events per sample were acquired using a FACScan flow cytometer and the percentage of cell apoptosis was analyzed using BD Accuri™ software (both from Becton-Dickinson, San Jose, CA, USA).

Evaluation of fluorescent LC3 puncta

LC3 puncta were indicated by mRFP-GFP-LC3 adenovirus (Gene Chem, Shanghai, China) Briefly, cells were transfected with mRFP-GFP-LC3 for 48 h before receiving Transfection treatments. After 24-hours Transfection treatment, cells were observed under a fluorescence microscopy (Nikon). Ten views were examined per slide and the images were acquired.

miRNA microarray profiling and analysis.

Total RNA was extracted with the use of TRIzol reagent (Invitrogen, Carlsbad, CA, USA). Human miRNA Microarray(GeneChip miRNA 4.0,Affymetrix)was used for microRNA profiling. The quality of total RNA was evaluated by measuring the ratio of A260/A280 on an ultraviolet pectrophotometer (NanoDrop 2000 Thermo) and by Checking the integrity and purity with a nucleic acid electrophoresis analyzer (Agilent 2100 Bioanalyzer)Biotin-labeled complementary RNA (cRNA) generated by in vitrotranscription reactions(Affymetrix, USA) was fragmented and hybridized with GeneChip Hybridization Oven 645(Affymetrix, USA) at 45 °C for 16 h.After hybridization, the arrays were washed, stained with GeneChip™ Hybridization, Wash, and Stain Kit (Thermo Fisher Scientific) and scanned using an Affymetrix GeneChip Scanner 3000 .After normalization and background filtration, the differentially expressed genes were analyzed.Using TargetScan,microRNA.ORG and miRDB software for target gene prediction.

Result

Mig-6 induced the apoptosis and reduced the autophagy of HCC cell lines.

To validate the role of *Mig-6* in HCC cell lines,we examined the impact exerted by increasing *Mig-6* on HCC cell apoptosis and autophagy with *Mig-6* plasmid.HepG-2 and HLE cells were transfected with *Mig-6* plasmid, respectively. As expected, two cell lines transfected with *Mig-6* showed the enhanced percentage of apoptosis(Fig. 1a) .In the meanwhile western blotting were using for evaluated the expression levels of autophagy-associated protein (Fig. 1b). The expression levels of p62, which is mainly degraded by autophagy pathway, is greatly induced in the *Mig-6* plasmid groups compared with those of the PC groups. But microtubule associated protein 1 light chain 3β (LC3b), following transfection with *Mig-6* plasmid were significantly decreased. Autophagic flux experiment showed that in two cell lines, *Mig-6* plasmid weakened the flux of autophagy, with a less number of autophagosomes (yellow) and autolysosomes (red) in two celllines(Fig. 1c).

Identification of Mig-6-regulated miRNAs

Several underlying mechanisms of *Mig-6* were hypothesized; the regulation of miRNAs by *Mig-6* was an area that had never been studied. *Mig-6*-regulated miRNAs were identified by transducing 293T with

lentiviruses encoding control virus or lentivirus-mediated *Mig-6* (Gene Chem, Shanghai, China). After transduction for 48 h, the cells were treated with puromycin for 48 h to remove the nontransduced cells. Successful transduction was examined by fluorescence microscopy of GFP-expressing cells (Fig. 2a), which was further confirmed by western blot analysis (Fig. 2b). The GFP-positive cell population was used immediately for analysis using the GeneChip assay. The results showed that a total of 44 *Mig-6*-regulated miRNAs were identified, of which 11 were upregulated and 33 downregulated (Fig. 2c). Among them the expression of 4 and 16 miRNAs in *Mig-6* overexpressed cells was increased or inhibited by more than 2.5-fold, respectively (Table 1). Among all the miRNAs, miR-193a-3p is the most potent, with downregulated expression reported in many types of cancers. The HepG-2 and HLE cells were transfected with *Mig-6* plasmid and siRNA to further verify the relationship between *Mig-6* and miR-193a-3p. The transfection efficiency was measured using Western blot analysis (Fig. 3a). When the expression of *Mig-6* was promoted, the miR-193a-3p level correspondingly increased in each cell line, and when the expression of *Mig-6* was suppressed, the decrease in the miR-193a-3p was observed (Fig. 3b).

Table 1
Expression changes of Mig6-regulated miRNAs

miRNA	Regulation	Control virus or LV-Mig-6 Fold change
hsa-miR-1269a	up	3.663214041
hsa-miR-193a-3p	up	2.960149418
hsa-miR-1244	up	2.512778203
hsa-miR-371b-5p	up	2.503430956
hsa-miR-5093	down	-5.363445158
hsa-miR-297	down	-4.571560354
hsa-miR-4793-3p	down	-4.556744037
hsa-miR-610	down	-4.539741026
hsa-miR-4440	down	-4.535851264
hsa-miR-195-3p	down	-4.458473988
hsa-miR-6877-3p	down	-4.173369522
hsa-miR-3148	down	-3.956889197
hsa-miR-8071	down	-3.647623099
hsa-miR-185-3p	down	-3.48629987
hsa-miR-3064-5p	down	-3.347459653
hsa-miR-595	down	-3.344313283
hsa-miR-3907	down	-3.319532324
hsa-miR-7844-5p	down	-2.856073948
hsa-miR-4455	down	-2.736743073
hsa-mir-297	down	-2.606974853

MiR-193a-3p induced the apoptosis and reduced the autophagy of HCC cell lines.

MiR-193a-3p is reported to relate to various pathological processes of liver cancer, including proliferation, migration, invasion, apoptosis, radioresistance, and chemotherapeutic tolerance. However, the relationship between miR-193a-3p and autophagy has not been reported to date. This study attempted to clarify the impact exerted by miR-193a-3p. First, the HepG-2 and HLE cells were transfected with miR-193a-3p mimics or miR-193a-3p inhibitors, and the transfection efficiency was measured by reverse transcription-polymerase chain reaction (Fig. 4a). Then, apoptosis was inspected in the HepG-2 and HLE

cells with flow cytometric analysis. A larger percentage of apoptotic cells were presented in the cells transfected with miR-193a-3p mimics; however, miR-193a-3p inhibitors decreased the apoptotic rate (Fig. 4b ,c). Finally, the effect of miR-193a-3p on the autophagy capacity of HepG-2 and HLE cells was verified using western blot analysis and autophagic flux experiment. As anticipated, miR-193a-3p mimics increased the expression of p62 but reduced the expression of LC3b, as well as weakened the flux of autophagy. The reverse consequence was also observed in the miR-193a-3p inhibitor group (Fig. 4d,e).

MiR-193a-3p affected the apoptosis and autophagy of HCC cells by regulating the expression of TGF- β 2.

The miR-193a-3p target predicted by all three target gene prediction software was regarded with high confidence (Table S1). Thus, 62 nominated target genes were hit by miR-193a-3p. Among them, the TGF- β 2 gene gained attention. Previous studies reported that TGF- β 2 was regulated by multiple miRNAs and associated with apoptosis and autophagy in certain cell lines. In this study, the expression of TGF- β 2 protein in both HepG-2 and HLE cells decreased after the transfection of miR-193a-3p mimics, whereas treatment with miR-193a-3p inhibitors increased the expression of TGF- β 2 (Fig. 5a). TGF- β 2 mRNA was also quantified after transfecting miR-193a-3p mimics or inhibitors into HCC cells, but no statistically significance difference was found at the mRNA level (Fig. 5b). Next, the apoptosis and autophagy were applied to assess the impact of miR-193a-3p and TGF- β 2 on HCC. The HepG-2 and HLE cells transfected with miR-193a-3p mimics showed an increased percentage of apoptotic cells and weakened flux of autophagy. However, TGF- β 2 plasmid could not only inhibit apoptosis and enhance the flux of autophagy but also weaken the effects exerted by miR-193a-3p mimics (Fig. 5c,d,e). These results substantiated that miR-193a-3p affected the apoptosis and autophagy of HCC cells by controlling the expression of TGF- β 2.

Mig-6 modulated apoptosis and autophagy of HCC cells through the miR-193a-3p/TGF- β 2 pathway.

The relationship between *Mig-6* and TGF- β 2 has not been reported to date. Therefore, the correlation between *Mig-6* and TGF- β 2 protein level was analyzed in five HCC cell lines. An inverse relationship was observed between the expression of *Mig-6* and TGF- β 2. High expression of *Mig-6* was found in Huh7 and PLC/PRF/5 cells; both of them showed low levels of TGF- β 2. On the contrary, the HepB3 and HLE cells with the low level expression of *Mig-6* had high expression of TGF- β 2 (Fig. 6a). A series of function experiments were performed in HepG-2 and HLE cells to assess further impacts of overexpression of *Mig-6*. As indicated by western blot analysis, the level of TGF- β 2 protein decreased after HCC cells were transfected with *Mig-6* plasmid (Fig. 6b,c), without any mRNA alteration (data not show). Meanwhile, miR-193a-3p reversed the decrease in TGF- β 2 induced by the *Mig-6* plasmid (Fig. 6b,c). As presented in Fig. 1A, a larger percentage of apoptotic cells were present in the cells transfected with *Mig-6* plasmid compared with the control group, whereas they were rescued by miR-193a-3p inhibitors (Fig. 6d). The autophagy of HCC cells was tested by western blot analysis and autophagic flux assay. Weakened autophagy due to the overexpression of *Mig-6* was reversed by miR-193a-3p inhibitors (Fig. 6b,c,e).

Discussion

Autophagy is a tightly regulated and highly conserved lysosomal degradation pathway for the degradation of long-lived proteins and cytoplasmic organelles. Recent accumulating data point to autophagy as a key player in various human diseases, such as cancer[17]. Apoptosis is the process by which a cell ceases to grow and divide, resulting in the controlled death of the cell. The initiation of apoptosis depends on the activation of a series of cysteine-aspartic proteases known as caspases [18]. Autophagy correlates with apoptosis. For example, the autophagy of damaged organelles can repair the injury to a small number of mitochondria; however, if a sufficiently large number of mitochondria are damaged, the intrinsic pathway of apoptosis is activated following the high levels of released cytochrome C[19]. This study demonstrated that *Mig-6* promoted the apoptosis of HCC cells, combined with the inhibition of the flux of autophagy. Therefore, *Mig-6* might exert its role by regulating the dynamic balance between apoptosis and autophagy in HCC cells.

As reported, *Mig-6* could be regulated by certain miRNAs[20–23], Especially miR-589-5p and miR-374a, regulated the proliferation of HCC cells by targeting *Mig-6*[24, 25]. These studies stimulated the interest in exploring whether *Mig-6* could affect the changes in miRNAs. The miRNA microarray profiling was used to identify miRNAs specifically regulated by *Mig-6*. A total of 44 *Mig-6*-regulated miRNAs were identified, of which 11 were upregulated and 33 downregulated (Figure S1). Among these miRNAs, four miRNAs were upregulated more than 2.5 times by *Mig-6*, and miR-193a-3p was one of them.

MiR-193a-3p is reported as a tumor suppressor in many types of cancers. It inhibits the migration and invasiveness of NSCLC[26]. In colorectal cancer, miR-193a-3p inhibited cell proliferation and promoted the apoptosis[27]. MiR-193a-3p suppressed ovarian cancer cell growth and migratory/invasive capacities[28]. MiR-193a-3p had also been proven to be related to the migration and invasion[14], proliferation and apoptosis [15, 16], radioresistance, and chemotherapeutic tolerance in HCC[29, 30]. In addition, the expression of miR-193a-3p was identified as a predictor of HCC progression-free survival[31]. These results demonstrated the important role of miR-193a-3p in HCC. In this study, miR-193a-3p mimics induced a high level of apoptosis in HCC cell lines; however, miR-193a-3p inhibitors decreased the apoptotic rate, which was consistent with the previous findings. Also, the autophagy capacity of HCC cell lines was found to be weakened by miR-193a-3p mimics, whereas miR-193a-3p inhibitor could enhance the flux of autophagy. All these findings confirmed that miR-193a-3p had an anti-cancer effect on HCC by influencing cell autophagy and apoptosis *in vitro*.

Numerous targets of miR-193a-3p have been reported[26, 28, 32, 33]. In this study, the TargetScan, miRNA.ORG, and miRDB software were used for the prediction of target genes. Finally, TGF- β 2 was selected, which might play an important role in HCC as the target of miR-193a-3p. TGF- β has three different isoforms: TGF- β 1, TGF- β 2, and TGF- β 3, with similar but not identical biologic activities[34]. The role of TGF- β signaling in HCC progression is not completely understood. Immunosuppression induced by TGF- β 2 was assumed to be the main mechanism through which tumor cells could escape from immune surveillance[35]. TGF- β 2 was upregulated in liver disease, suggesting it to be a promising therapeutic target for tackling fibrosis and HCC[36]. Exogenous TGF- β 2 resulted in a significant elevation of the epithelial to mesenchymal transition by enhancing autophagic flux in HCC[37]. This study eventually

concluded that the upregulation of TGF- β 2 could inhibit apoptosis and promote the autophagic flow of HCC cells. It also confirmed the inverse correlation between *Mig-6* and TGF- β 2.

This study had certain limitations. First, the clinical significance of the interaction among *Mig-6*, miR-193a-3p, and TGF- β 2 in HCC needed to be analyzed, which could increase the value of *Mig-6*, miR-193a-3p, and TGF- β 2 in the diagnosis. Second, the overexpression of *Mig-6* or *Mig-6* gene knockout mice could be applied to verify the *Mig-6*/miR-193a-3p/TGF- β 2 axis *in vivo*. Third, many miRNAs regulated by *Mig-6* were also found in this study, but their mechanism was not studied. Subsequent experiments need to explore the new functions of *Mig-6*-regulated miRNAs in HCC and provide new perspectives on the treatment of HCC.

Conclusion

Taken together, our data verified that *Mig-6* and miR-193a-3p as the tumor suppressors in HCC, which can promote the apoptosis and inhibition of the flux of autophagy by targeting TGF- β 2. The *Mig-6* regulated miRNAs and the correlation between *Mig-6* and TGF- β 2 were also confirmed in HCC for the first time. This article provided clues for an important *Mig-6*/miR-193a-3p/TGF- β 2 regulatory axis, providing more perspectives on the treatment of HCC.

Declarations

Authors' contributions

ZXL, YLT and HSZ designed the study;ZXL, YLT, DH and FW performed the experiments and prepared the figures;ZXL and HSZ contributed to drafting the manuscript. All authors read and approved the final manuscript.

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Competing interests

The authors declare that they have no competing interests.

Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Consent for publication

Not applicable.

Ethics approval and consent to participate

The study was approved by the Ethics Review Committee of The First Affiliated Hospital of China Medical University (Shenyang, China).

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Figures

Figure 1

Effect of Mig-6 plasmid on the apoptosis and autophagy of HCC cell lines. a The apoptosis assay after 48h transfected with plasmid control and Mig-6 plasmid in HepG-2 and HLE cells, the percentage of apoptosis cells was quantified. b Western blotting and quantitative analysis of Mig-6, LC3b and p62 protein levels in HepG-2 and HLE cells. c Each group was transfected with a tandem mRFP-GFP-LC3 adenovirus for 24 h. Autophagosomes and autolysosomes were, respectively, visualized as yellow- and red-only punctas under a fluorescence microscope. (*P<0.05, **P < 0.01, ***P < 0.001).

Figure 2

Prediction and confirmation of Mig-6 regulated miRNAs. a Visualization of GFP expression in Control virus and LV-Mig-6 group(100×) after 72h. b Western blotting and quantitative analysis of transduction efficiency of LV-Mig-6 in 293T cell lines. c Hierarchical cluster analysis of Mig-6-regulated miRNAs. The horizontal axis shows comparison groups of expression change of miRNAs. The left vertical axis shows clusters of Mig-6-regulated miRNAs whose fold change is listed in the right-hand table. Red indicates upregulated and green downregulated miRNAs.

Figure 3

Confirmation of Mig-6 regulated miRNAs. a Western blotting and quantitative analysis of transfection efficiency of Mig-6 plasmid and siMig-6 in HepG-2 and HLE cell lines. b Quantitative RT-PCR analysis of the expression levels of miR-193a-3p in Mig-6 up-regulate and down-regulate groups. (** $P < 0.01$, *** $P < 0.001$).

Figure 4

Effect of miR-193a-3p on the apoptosis and autophagy of HCC cell line. a Quantitative RT-PCR analysis of the expression levels of miR-193a-3p in HepG-2 and HLE cells transfected with miR-193a-3p mimics, inhibitors, and negative control. b, c The apoptosis assay 48h after four groups of HepG-2 and HLE cells transfected with equal doses of mimic control, miR-193a-3p mimics, inhibitor control and miR-193a-3p inhibitors, the percentage of apoptosis cells was quantified. d Western blotting and quantitative analysis for Mig-6, LC3b and p62 expression 48 h after aforementioned four groups' transfection in HepG-2 and HLE cells. e Each group was transfected with a tandem mRFP-GFP-LC3 adenovirus for 24 h.

Autophagosomes and autolysosomes were, respectively, visualized as yellow- and red-only punctas under a fluorescence microscope. (* $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$).

Figure 5

Effect of miR-193a-3p and TGF- β 2 plasmid on the HCC cell lines. a Western blotting and the quantitative analysis of TGF- β 2 protein levels in HepG-2 and HLE cells after transfected with equal doses of the miR-193a-3p mimics, inhibitors, and negative control. b Quantitative RT-PCR analysis of the expression levels

of TGF- β 2 mRNA in HepG-2 and HLE cells transfected with miR-193a-3p mimics, inhibitors, and negative control. c The apoptosis assay 48h after four groups of HepG-2 and HLE cells transfected with equal doses of mimic control plus plasmid control, miR-193a-3p mimics plus plasmid control, mimic control plus TGF- β 2 plasmid and miR-193a-3p mimics plus TGF- β 2 plasmid , the percentage of apoptosis cells was quantified. d Western blotting and quantitative analysis for TGF- β 2, LC3b and p62 expression 48 h after aforementioned four groups' transfection in HepG-2 and HLE cells. e Each group was transfected with a tandem mRFP-GFP-LC3 adenovirus for 24 h. Autophagosomes and autolysosomes were, respectively, visualized as yellow- and red-only punctas under a fluorescence microscope. (* P <0.05,** P < 0.01, *** P < 0.001).

Figure 6

Relationship between miR-193a-3p and TGF- β 2 , and the effect of Mig-6 plasmid and miR-193a-3p inhibitors on TGF- β 2. a The correlation of Mig-6 and TGF- β 2 in HCC cell lines. b,c, The apoptosis assay 48h after forementioned four groups' transfection in HLE cells. d Western blotting and quantitative analysis for Mig-6,TGF- β 2, LC3b and p62 expression 48h after four groups of HLE cells transfected with equal doses of inhibitor control plus plasmid control, miR-193a-3p inhibitors plus plasmid control, inhibitor control plus Mig-6 plasmid , and miR-193a-3p inhibitors plus Mig-6 plasmid. e Each group was transfected with a tandem mRFP-GFP-LC3 adenovirus for 24 h. Autophagosomes and autolysosomes were, respectively, visualized as yellow- and red-only punctas under a fluorescence microscope. (*P<0.05,**P < 0.01, ***P < 0.001).

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