let-7a suppresses liver fibrosis via TGFβ/SMAD signaling transduction pathway

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Abstract. Liver fibrosis is the most common pathological outcome and the most severe complication of chronic liver diseases. Accumulating evidence suggests that miRNAs are involved in cell proliferation, differentiation, apoptosis, as well as the occurrence and development of various diseases. In this study, we found that the expression of let-7a was markedly decreased in the liver tissue samples and blood samples from patients with liver fibrosis compared with healthy volunteers. Furthermore, let-7a was downregulated in the liver tissues and blood samples in mouse models of liver fibrosis. Further analysis indicated that let-7a suppresses the activation level of hepatic stellate cells (HSCs). In addition, overexpression of let-7a reduced cell viability and promoted apoptosis of HSCs. Western blot analysis showed that let-7a might inhibit HSCs through TGFβ/SMAD signaling pathway. The present study provides a potential accurate target and vital evidence to better understand the underlying pathogenesis for early diagnosis and treatment of liver fibrosis.

Introduction

Chronic liver diseases remain a major problem throughout the world and lead to considerable morbidity and mortality (1). The most common pathological outcome and the most severe complication of chronic liver diseases are liver fibrosis which are global health and economic burdens (2). Liver fibrosis characteristically show over-accumulation of extracellular matrix (ECM), mainly type I fibrillar collagen (Collagen I) in response to chronic liver damage (3). Various etiological factors and stimuli can result in development of liver fibrosis, including viral hepatitis (hepatitis C and B), environmental carcinogens, excessive ethanol consumption, activated hepatic stellate cells (HSCs) and non-alcoholic steatohepatitis (NASH) (4,5). Liver fibrosis is generally asymptomatic and is often overlooked by patients and their families. When patients are diagnosed with liver fibrosis, the strategies for treatment and therapeutic options are seriously limited (6). In addition, there are no vaccine or effective anti-fibrotic drugs so far, and the patients with liver fibrosis need to receive long-term and repeated drug treatments, which is difficult to accept (7). Therefore, it is necessary to search for a better therapy to directly prevent liver fibrosis, and a better understanding of pathogenesis underlying the development of liver fibrosis is urgently needed.

In recent years, therapy based on cells has been widely investigated in the area of tissue or organ protection (8). Therefore, we focus on the cells which are connected with the development of liver fibrosis. HSCs have been considered as a kind of lipocytes, which exhibit a key role in the process of liver fibrosis when the liver is damaged (9,10). HSCs are regarded as the potential aetiology because they are the key cell type which is in charge of ECM formation during hepatic fibrogenesis (11). Activation of HSCs is essential to the initiation and progression of liver fibrosis (12). When HSCs are activated, they become fibrogenic myofibroblasts and exhibit the function of fibrogenic myofibroblast-like cells, which can secrete α-smooth muscle actin (α-SMA), TIMP-1 and inhibit HSC cell apoptosis and collagen-I secretion (13-15). Excessive insoluble collagen I and matrix components in the intracellular and perisinusoidal spaces can bring severe compromise to the hepatic function (16,17).

MicroRNAs (miRNAs) have been found to be the most abundant small non-coding RNAs (ncRNAs) (18). miRNAs are a type of 18 to 25 nucleotides (nt) in length, single-stranded and evolutionary conserved endogenous RNAs (19). miRNAs can regulate the expression of post-transcriptional genes by incompletely binding with their target genes 3′-UTR. Accumulating evidence suggests that miRNAs are related to the cell proliferation, differentiation, apoptosis and development of various diseases (20). Recent studies have shown that
miRNAs take part in the process of liver diseases, such as liver injury, hepatic cirrhosis, hepatocellular carcinoma and liver fibrosis (21-24). Moreover, there is increasing evidence that miRNAs are involved in various fibrotic diseases, including kidney, cardiac, cystic and pulmonary fibrosis (25-28). miRNAs may serve as antifibrotic or profibrotic genes during liver fibrosis. However, the mechanism effects of miRNAs need further investigation.

let-7a has been found to be involved in the development of chronic liver diseases (29). However, let-7a has not been indicated to be related to liver fibrosis. Here, we investigated the expression level of let-7a in patients with liver fibrosis to reveal the molecular mechanism of let-7a regulating the pathogenesis of liver fibrosis. Our study aimed to provide an accurate target and vital evidence to better understand the underlying pathogenesis for early diagnosis and treatment for liver fibrosis.

Patients and methods

**Human liver samples.** Liver samples were collected from patients and volunteers attending The Second Affiliated Hospital of Qiqihar Medical University (Qiqihar, China) from January, 2016 to March, 2017. Fibrotic liver samples were taken from 12 non-tumorous patients (6 males and 6 females; age range 20-40 years) who were diagnosed to have liver fibrosis. The tissues of patients were taken and after pathological examination the remaining tissues were treated with RNAKeeper (Vazyme Biotech Co., Ltd., Nanjing, China) and stored at -80°C (Thermo Fisher Scientific, Inc., Waltham, MA, USA). Normal liver tissues (n=12) were obtained from healthy liver transplant donors undergoing hepatic resection and regarded as controls in this study. All clinical samples were collected in The Second Affiliated Hospital of Qiqihar Medical University.

This study was approved by the Clinical Ethics Committee of The Second Affiliated Hospital of Qiqihar Medical University. All participants agreed with the research, participated in this study willingly and signed an informed consent before sample collection.

The control tissues and liver fibrosis tissues were stored at -80°C for further analysis. According to the pathological results of liver biopsy, the cases of the patients with liver fibrosis was 4 (S0), 4 (S1), 2 (S2), 1 (S3), 1 (S4). According to the different stage, the value of acoustic radiation force impulse (ARFI) of the patients with liver fibrosis was 0.98±0.10, 1.14±0.11, 1.38±0.40, 1.91±0.70 and 2.12±0.75 m/sec, respectively.

**Collection of human blood samples.** Blood samples were obtained from patients and volunteers in The Second Affiliated Hospital of Qiqihar Medical University from January, 2016 to October, 2017. Human blood samples were collected from 22 hospitalized patients who had liver fibrosis (11 males and 11 females; age range 21-40 years) and 22 age- and sex-matched healthy subjects according to a protocol approved by the Institutional Review Board. The healthy volunteers who suffered from known acute or chronic diseases, who took medications or alcohol within 24 h or were aged less than 21 years were excluded. All human experiments were conducted according to the clinical Ethics Committee of The Second Affiliated Hospital of Qiqihar Medical University.

**RNA extraction and cDNA synthesis.** The frozen liver samples were thawed and homogenized at 4°C prior to RNA isolation. TRIzol reagent (Thermo Fisher Scientific, Inc.) was used to isolate the total RNAs from the liver samples, blood samples and cells according to the protocols. After isolation, RNAs were quantified by Nanodrop machine (Thermo Fisher Scientific, Inc.) and RNAs were used in further experiments when the concentration was higher than 200 (ng/µl) and A260/280 ratio was 1.7-2.0. Single-stranded cDNAs were synthesized using reverse transcription kit (Vazyme Biotech Co., Ltd.). For cDNA synthesis, a 10 µl reaction volume composed of 500 ng RNAs, 4 µl reverse transcriptase and double-distilled water (ddH2O) were added into 200 µl EP tubes. The reactions were initially incubated at 50°C for 15 min and then at 85°C for 5 sec.

**Cell culture.** The HSC cell line LX-2 cells (cat.no.GD-C624092) were obtained from Changsha Yingrun Biotechnologies Co., Ltd. (Changsha, China). The cells were cultured in DMEM medium (HyClone; GE Healthcare Life Sciences, Logan, MT, USA) containing 10% fetal bovine serum (FBS; Thermo Fisher Scientific, Inc.), 100 µg/ml penicillin-streptomycin antibiotics (cat. no. C0222; Beyotime Institute of Biotechnology, Haimen, China) and 2 µM L-glutamine (Sigma-Aldrich; Merck KGaA, Darmstadt, Germany). The cells were seeded in 25 cm² flasks and grown in an incubator with 5% CO₂ and 37°C (Thermo Fisher Scientific, Inc.). The cells were plated at a density of 5x10⁴ cells/well and routinely subcultured using 0.25% trypsin when the confluence reached approximately 80%. The method of quantification was the 2–ΔΔCq method (30).

**Cell transfection.** To overexpress or silence let-7a, let-7a mimics, let-7a inhibitor, miR-NC and antagonirm-NC were transfected into LX-2 cells. Transient transfections were carried out using transfection reagent X-treme (Vazyme Biotech Co., Ltd.) on the basis of previous studies. On the day prior to cell transfection, LX-2 cells were plated in 96-well or 6-well plates. Let-7a mimics, let-7a inhibitor, miR-NC and antagonirm-NC were designed and synthesized by Shanghai GenePharma Co., Ltd. (Shanghai, China). The concentration of let-7a-5p mimics and miR-NC was 100 nM. The concentration of let-7a-5p inhibitor and antagonirm-NC was 50 nM. After 24 h transfection the cells were used for further experiments.

**Cell viability analysis.** LX-2 cells were placed at a density of 2x10³ cells/well in 96-well plates (Corning Incorporated,
Corning, NY, USA). After culture of 24 h, 20 µl volume of MTT solution (5 mg/ml, BioSharp, Hefei, China) was added and the cells were incubated for 4 h at 37°C in an incubator. Then about 150 µl of dimethyl sulfoxide (DMSO) was added and the plates were gently shaken. The optical density values (λ=490 nm) were detected using the enzyme linked immunosorbent assay. All experiments were repeated at least three times.

**TUNEL staining.** To measure the apoptosis levels of LX-2 cells, 1x10^5 LX-2 cells were grown in a small dish. The cells transfected with let-7a mimics, let-7a inhibitor, miR-NC and antagonor-NC were used for TUNEL staining. The culture medium in the dish was discarded, and cells were fixed in 4% PFA for 30 min at room temperature after being washed with PBS. After the closure and penetration procedures, TUNEL staining solution Vial1 and Vial2 were mixed at a ratio of 1:9 and the cells were incubated with the TUNEL reagent for 1 h at room temperature in the dark. After being washed with PBS, cells were incubated with Hoechst staining solution for 20 min at room temperature, and kept in the dark. Finally, the cells were observed under a fluorescence microscope (Nikon Corporation, Tokyo, Japan). Ten images were randomly captured of every angle. The quantification of TUNEL staining was performed by measuring the percentage of TUNEL-positive cells.

**Western blotting.** Cells after transfection with let-7a were lysed in 500 µl volume of RIPA lysis buffer for 20 min in a 4°C freezer. After extraction, the concentration of total proteins was measured using BCA protein detection kit (Beyotime Institute of Biotechnology). After heat denaturation at 100°C for 10 min, equal amounts of extracted protein samples were separated by 8-10% SDS-PAGE and transferred onto PVDF membrane (EMD Millipore, Billerica, MA, USA). After using 5% bovine serum albumin (Sigma-Aldrich; Merck KGaA) to block non-specific protein binding at room temperature for 30 min, the membrane was incubated with anti-TGF-β or anti-SMAD antibody with a gentle shaking overnight at 4°C. Then, the membrane was incubated with secondary antibody and the signal was detected using an ECL system. The expression of GAPDH was used as loading control to quantify the expression of TGF-β and SMAD by using Image Studio software. For western blot analysis, anti-TGF-β (anti-rabbit; cat. no. CST3711; 1:1,000) anti-SMAD2 (anti-rabbit, cat. no. CST3339; 1:1,000), anti-SMAD3 (anti-rabbit, cat. no. 9513; 1:1,000) antibodies, the second antibody (anti-rabbit; cat. no. CST7074; 1:1,000), or the second antibody (anti-mouse; cat. no. 5571; 1:1,000) all from Cell Signaling Technology, Inc., (Danvers, MA, USA) and GAPDH (anti-mouse; cat. no. sc-47724; 1:500) from Santa Cruz Biotechnology, Inc., (Dallas, TX, USA).

**Statistical analysis.** The data were analyzed from three or more independent experiments. Analysis was performed by using GraphPad Prism 5 software (GraphPad Software, Inc., La Jolla, CA, USA). Differences between the groups were analyzed by one-way ANOVA, followed by a Tukey’s test for multiple comparisons. The Mann-Whitney U test was used for comparison between groups. P<0.05, P<0.01 and P<0.001 was considered statistically significant.

**Results**

**Expression of let-7a in patients with liver fibrosis.** To determine whether let-7a is related to liver fibrosis, the expression of let-7a was detected in the tissues and blood samples from patients with liver fibrosis using RT-qPCR analysis. First, the analysis of supersonic inspection showed that the hepatic vein was not clear (Fig. 1A). The echo in the liver from patient with liver fibrosis become deeper and weaker with uneven distribution, and the analysis showed that the hepatic vein was not clear (Fig. 1A). The level of let-7a was differentially expressed in the liver fibrosis tissues compared to normal liver tissues as indicated in Fig. 1B. The results of RT-qPCR revealed that the expression of let-7a was lower in the liver tissues from patients with liver fibrosis compared with that in healthy volunteers, and the difference was statistically significant (Fig. 1B). In addition, let-7a was markedly downregulated in the blood samples of patients with

Table I. Primer sequences.

| Primers | Forward | Reverse |
|---------|---------|---------|
| U6      | CCTCGCCAGCACATATACT | TTCACGAATTTGCGTGTCAT |
| GAPDH   | TCCACTGGCGCTTTCACC  | GCCAGAGTATGACCCTTCTT |}

λ=490 nm) were detected using the enzyme linked immunosorbent assay. All experiments were repeated at least three times.

**TUNEL staining.** To measure the apoptosis levels of LX-2 cells, 1x10^5 LX-2 cells were grown in a small dish. The cells transfected with let-7a mimics, let-7a inhibitor, miR-NC and antagonor-NC were used for TUNEL staining. The culture medium in the dish was discarded, and cells were fixed in 4% PFA for 30 min at room temperature after being washed with PBS. After the closure and penetration procedures, TUNEL staining solution Vial1 and Vial2 were mixed at a ratio of 1:9 and the cells were incubated with the TUNEL reagent for 1 h at room temperature in the dark. After being washed with PBS, cells were incubated with Hoechst staining solution for 20 min at room temperature, and kept in the dark. Finally, the cells were observed under a fluorescence microscope (Nikon Corporation, Tokyo, Japan). Ten images were randomly captured of every angle. The quantification of TUNEL staining was performed by measuring the percentage of TUNEL-positive cells.

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**Construction model of liver fibrosis.** To generate the model of liver fibrosis, 20 female C57BL/6 mice (approximately 18-22 g) were purchased. Mice were kept at 25°C and raised for a week before experiments. Mice were intraperitoneally injected with olive oil vehicle (Sigma-Aldrich; Merck KGaA) and they served as the control. In addition, mouse model with liver fibrosis was intraperitoneally injected with CCl4 (0.5 µl/g, Sigma-Aldrich; Merck KGaA) which was normally used as hepatotoxic chemical twice weekly. After injections of 10 weeks, the mice were sacrificed and the liver tissues and blood samples were harvested.

**Statistical analysis.** The data were analyzed from three or more independent experiments. Analysis was performed by using GraphPad Prism 5 software (GraphPad Software, Inc., La Jolla, CA, USA). Differences between the groups were analyzed by one-way ANOVA, followed by a Tukey’s test for multiple comparisons. The Mann-Whitney U test was used for comparison between groups. P<0.05, P<0.01 and P<0.001 was considered statistically significant.

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Validation of let-7a expression in mice with liver fibrosis. To identify the level of let-7a in the mice with liver fibrosis, a model of liver fibrosis was generated. Briefly, mice (n=10) were intraperitoneally injected CCl4 twice weekly for 10 weeks, while the control group (n=10) were treated with olive oil. The decreased expression of let-7a in the liver tissues from mice of liver fibrosis was revealed by RT-qPCR (Fig. 2A). Similarly, according to the RT-qPCR analysis, the expression of let-7a was observably decreased in the blood samples of mice with liver fibrosis compared with control group, and the difference was statistically significant (Fig. 2B). Therefore, we concluded that let-7a was reduced in the liver tissues and blood samples of mice with liver fibrosis.

The role of let-7a in the activation of HSCs. To further confirm the role of let-7a in the activation of HSCs, LX-2 cells were transfected with let-7a mimics, let-7a inhibitor, miR-NC and antagonim-NC for 24 h. The expression of let-7a was detected by PCR analysis. The expression of let-7a was significantly increased after transfection of let-7a mimics, but decreased in the presence of let-7a inhibitor (Fig. 3A). After transfection, the expression of markers of HSC activation, including α-SMA, Col1a1 and Col1a4 was analyzed. We found that overexpression of let-7a significantly decreased the expression level of α-SMA, while the decrease in let-7a elevated the level of α-SMA (Fig. 3B). RT-qPCR assay indicated that LX-2 cells transfected with let-7a mimics decreased level of Col1a1, and the expression of Col1a1 was increased by transfection of let-7a inhibitor (Fig. 3C). In addition, the expression of Col1a4 was inhibited by let-7a mimics in comparison with miR-NC (Fig. 3D). Furthermore, the expression level of Col1a4 was higher in cells treated with let-7a inhibitor than that in group antagonim-NC (Fig. 3D). The above results confirmed that let-7a led to a marked decrease of HSC activation.

The effect of let-7a on cell viability and apoptosis of HSCs. To further study the role of let-7a in the cell viability and

Figure 1. Expression of let-7a in patients with liver fibrosis. We collected the liver tissues and blood samples from patients with liver fibrosis and healthy controls, respectively. (A) The results of supersonic inspection from patient with liver fibrosis. (B) The expression levels of let-7a in the normal liver tissue and liver fibrosis tissue were measured by RT-qPCR. (C) The expression levels of let-7a in the blood samples from patients with liver fibrosis and healthy volunteers were measured by RT-qPCR. n=3 per group. **P<0.01; ***P<0.001 compared with controls.

Figure 2. The expression of let-7a in model of liver fibrosis. Models of liver fibrosis and controls were constructed to validate whether expression of let-7a was dysregulated in the two groups. (A) RT-qPCR analysis validated the expression levels of let-7a in the normal liver tissue and liver fibrosis tissue from mice. (B) The expression levels of let-7a in the blood samples from controls and model of liver fibrosis were compared. **P<0.01 compared with controls.
apoptosis of HSCs, MTT assay and TUNEL staining were performed. As shown in Fig. 4A, let-7a was overexpressed, and the cell viability was inhibited. Compared with antagomir-NC, the cell viability of HSCs transfected with let-7a inhibitor was enhanced (Fig. 4A). Besides, TUNEL staining displayed that let-7a promoted apoptosis of HSCs, while let-7a inhibitor decreased the apoptotic cells (Fig. 4B). These observations implied that let-7a inhibited the cell viability but promoted apoptosis in HSCs.

**Let-7a suppresses liver fibrosis through TGF-β/SMAD signaling pathway.** To explore the underlying mechanism of let-7a regulating liver fibrosis, let-7a mimics, let-7a inhibitor, miR-NC and antagomir-NC were transiently transfected into HSCs. The protein expression levels of TGF-β, SMAD2 and SMAD3 were measured by western blotting. The results showed that the expression levels of TGF-β, SMAD2 and SMAD3 were decreased in HSCs transfected with let-7a mimics compared with miR-NC (Fig. 5A). Conversely, the protein levels of TGF-β, SMAD2 and SMAD3 were significantly elevated in cells after transfection with let-7a inhibitor (Fig. 5B). Overall, our results indicate that let-7a may regulate the activation, cell viability and apoptosis of HSCs through TGF-β/SMAD signaling pathway.

**Discussion**

Fibrosis is a complicated process involving HSC activation, deposition of extracellular matrix and genetic and epigenetic changes including miRNA. Liver fibrosis is caused by viral hepatitis and alcoholic steatohepatitis, which leads to cirrhosis and hepatocellular carcinoma (31,32). Emerging studies show that HSC is pivotal event in liver fibrosis (33,34). Therefore, in this study, our purpose is to investigate the role of let-7a in hepatic stellate cells and whether let-7a is dysregulated in the normal liver tissues and liver fibrosis.

MicroRNAs (miRNAs) are a type of endogenous small non-coding RNAs, and microRNAs are related to the development of various diseases (35,36). miRNAs have been found to play a pro-fibrotic or anti-fibrotic role in HSC activation and be in connection with liver fibrosis. For example, miR-30a was downregulated in fibrotic liver tissues and isolated HSCs, and miR-30a suppresses HSC activation including cell proliferation, expression of α-SMA and collagen by targeting Snai1 protein (37). A study has shown that miR-185 suppressed fibrogenic activation of hepatic stellate cells by suppressing HSC activation through targeting RHEB and RICTOR. miR-212 promotes liver fibrosis via activating HSCs and TGF-β pathway through targeting SMAD7, which highlights that miR-212 can be regarded as a key biomarker or therapeutic target for liver fibrosis (38).

In this study, the results indicated that let-7a was markedly reduced in the liver tissues and blood samples from patients with liver fibrosis compared with healthy volunteers in clinic. In addition, the level of let-7a was decreased in liver tissues and blood samples in mice with liver fibrosis which were constructed by intraperitoneal injected with CCl4. Further analysis revealed that overexpression of let-7a decreased the mRNA level of markers of HSC activation, such as α-SMA, ColIa1 and ColIa4, suggesting that let-7a inhibited the activation level of HSCs. let-7a inhibitor increased the expression levels of these key genes, which were measured by RT-qPCR.
Figure 4. let-7a regulates cell viability and apoptosis of HSCs. To study the effect of let-7a on cell viability and apoptosis of HSCs, the cells were transfected with let-7a mimics, let-7a inhibitor, miR-NC and antagomir-NC for 24 h. After 24 h, MTT and TUNEL assay were performed. (A) The effect of let-7a on cell viability of HSCs was determined by MTT assay. (B) The role of let-7a in apoptosis of HSCs was tested by TUNEL staining. *P<0.05; **P<0.01 compared with controls.

Figure 5. let-7a inhibits the protein expression of TGF-β/SMAD. To identify the mechanism of let-7a in HSCs, the protein expression levels of TGF-β, SMAD2 and SMAD3 were detected in the cells after transfection with let-7a. (A) Western blot analysis indicated that the expression levels of TGF-β, SMAD2 and SMAD3 were downregulated by let-7a mimics. (B) The expression levels of TGF-β, SMAD2 and SMAD3 were upregulated by let-7a inhibitor transfection. *P<0.05; **P<0.01; ***P<0.001 compared with controls.
analysis. Besides, transfection of let-7a reduced cell viability and promoted apoptosis of HSCs. In contrast, let-7a inhibitor increased cell viability and inhibited apoptosis of HSCs, which were in accordance with our hypothesis. Moreover, western blot analysis showed that let-7a might inhibit HSCs through TGFβ/SMAD signaling pathway. The present study provides new insights into the mechanisms behind the antifibrotic effect of let-7a. This study provided a potentially accurate target and vital evidence to better understand the underlying pathogenesis for early diagnosis and treatment of liver fibrosis.

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Availability of data and materials

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

Authors' contributions

YinghuiZ, YingboZ, JG,YL and KJ drafted the manuscript. YinghuiZ collected human blood samples. JG and YL helped with RNA extraction and cDNA synthesis. YinghuiZ and KJ performed PCR. YingboZ contributed to cell viability analysis. YinghuiZ and YingboZ wrote the manuscript. All authors read and approved the final manuscript.

Ethics approval and consent to participate

This study was approved by the Clinical Ethics Committee of The Second Affiliated Hospital of Qiqihar Medical University (Qiqihar, China). All participants agreed with the research, participated in this study willingly and signed an informed consent before sample collection.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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