Knockout of microRNA-21 reduces biliary hyperplasia and liver fibrosis in cholestatic bile duct ligated mice

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Abstract

Cholestasis is a condition that leads to chronic hepatobiliary inflammation, fibrosis, and eventually cirrhosis. Many microRNAs (miRs) are known to play a role in fibrosis progression; however, the role of miR-21 during cholestasis remains unknown. Therefore, the aim of this study was to elucidate the role of miR-21 during cholestasis-induced biliary hyperplasia and hepatic fibrosis. Wild-type (WT) and miR21−/− mice underwent sham or bile duct ligation (BDL) for 1 wk, before evaluating liver histology, biliary proliferation, hepatic stellate cell (HSC) activation, fibrotic response, and Smad-7 expression. In vitro, immortalized murine biliary cell lines (IMCL) and human hepatic stellate cell line (hHSC) were treated with either miR-21 inhibitor or control before analyzing proliferation, apoptosis, and fibrotic responses. In vivo, the levels of miR-21 were increased in total liver and cholangiocytes after BDL, and loss of miR-21 decreased the amount of BDL-induced biliary proliferation and intrahepatic biliary mass. Also, loss of miR-21 decreased BDL-induced HSC activation, collagen deposition, and expression of the fibrotic markers TGF-β1 and α-SMA. In vitro, IMCL and hHSCs treated with miR-21 inhibitor displayed decreased proliferation and expression of fibrotic markers and enhanced apoptosis when compared to control.
treated cells. Furthermore, mice lacking miR-21 show increased Smad-7 expression, which may be driving the decrease in biliary hyperplasia and hepatic fibrosis. During cholestatic injury miR-21 is increased and leads to increased biliary proliferation and hepatic fibrosis. Local modulation of miR-21 may be a therapeutic option for patients with cholestasis.

INTRODUCTION

Cholangiocytes are the target of cholangiopathies, such as primary biliary cirrhosis (PBC) and primary sclerosing cholangitis (PSC), which are associated with dysregulation of cholangiocyte proliferation/loss. Cholestatic injury occurs by the obstruction of intra- or extrahepatic bile ducts, and is characterized by bile ductular reaction and extensive fibrosis. The bile duct ligation (BDL) and multidrug resistance gene-2 knockout (Mdr2−/−) mouse models mimic some features of PSC including biliary damage and liver fibrosis. Normally, cholangiocytes are mitotically quiescent but following damage (such as cholestasis) they begin to proliferate to repair the biliary tree to compensate for damage and loss of functionality. Alongside this enhanced proliferative capacity, liver fibrogenesis occurs by the excessive accumulation of extracellular matrix proteins secreted by various cell types including activated hepatic stellate cells (HSCs). HSCs can be activated through a number of different factors secreted from cholangiocytes, including transforming growth factor (TGF)-β. Following TGF-β receptor activation, phosphorylation of the secondary messengers small mothers against decapentaplegic 2 and 3 (Smad-2/3) occurs which allows translocation to the nucleus where Smad-2/3 binds to the transcription factor Smad-4 and allows for increased cellular proliferation and fibrotic response. The effects of Smad-2/3 can be inhibited by the inhibitory Smad-7.

MicroRNAs (miRs) are conserved, small (20–25 nucleotide) non-coding RNAs that regulate RNA silencing and post-translational regulation of gene expression. Following cholestatic injury there is a wide range of miRNAs that are downregulated; however, only a few miRNAs like miR-199, miR-200, and miR-34 are upregulated. miR-21 is an ubiquitously expressed miRNA that is upregulated in many different cancer types. A previous study found that miR-21 increases fibrogenesis during an experimental model of non-alcoholic steatohepatitis via inhibition of Smad-7.

Previously, we have shown that miR-21 is upregulated in a model of alcoholic liver injury and decreases HSC apoptosis. However, this study did not delve into the role of miR-21 on cholangiocytes during injury or HSC-promoted fibrosis. The role of miR-21 during cholestatic injury is largely unknown; therefore, we aimed to uncover the role of miR-21 during cholestatic injury.

MATERIALS AND METHODS

Materials

All reagents were obtained from Sigma-Aldrich, Co (St. Louis, MO) unless otherwise indicated. Cell culture reagents and media were obtained from Invitrogen Corporation (Carlsbad, CA). Antibodies for immunohistochemistry and immunofluorescence were
obtained from Abcam (Cambridge, MA) unless indicated otherwise. Total RNA was isolated from total liver tissues, purified cholangiocytes, and selected cell lines using the TRI Reagent from Sigma Life Science and reverse transcribed with the Reaction Ready First Strand cDNA synthesis kit (SABiosciences, Frederick, MD) as described. Total RNA was extracted from HSCs (isolated by Laser Capture Microdissection, LMD) using the Arcturus® PicoPure® RNA Isolation Kit (Applied Biosystems; Waltham, MA) according to the manufacturers protocol. The selected primers were purchased from Qiagen (Valencia, CA). The following primers were used: miR-21, mouse-MS00011487; RNU6-6P, mouse-MS00033740; Bax, mouse-PPM02917E-200; cleaved Caspase-3, mouse-PPM02922F-200; α-smooth muscle actin (α-SMA), mouse-PPM04483A-200; proliferating cell nuclear antigen (PCNA), mouse-PPM03456F-200; transforming growth factor-β1 (TGF-β1), mouse-PPM02991B-200; matrix metallopeptidase-9 (MMP-9), mouse-PPM03661C-200; small mothers against decapentaplegic 7 (Smad-7), mouse-PPM03073F-200; and glyceraldehyde 3-phosphate dehydrogenase (GAPDH), mouse-PPM02946E-200.

Animal Models

All animal procedures were performed according to protocols approved by the Baylor Scott & White Healthcare IACUC Committee. MicroRNA21 (miR-21) knockout (miR-21−/−) mice and background-matched miR-21+/+ (wild-type, WT), strain B6; 129) mice were purchased from Jackson Laboratory (Sacramento, CA); the breeding colony is established in our animal facility. No gross defects or phenotypical changes are noted in the miR-21−/− mice. Male FVB/NJ WT mice (control for Mdr2−/− mice) were purchased from Jackson Laboratory. The breeding colony for Mdr2−/− mice (purchased from Jackson Laboratory), a model of PSC, is established in our animal facility. Animals were maintained in microisolator cages in a temperature-controlled environment with 12:12-hr light-dark cycles; all animals were fed ad libitum a standard chow diet with free access to drinking water. Studies were performed in 12 wk-old male miR-21−/− and Mdr2−/− mice (25–30 gm) and the corresponding WT mice that were subjected to sham or bile duct ligation (BDL) for 1 wk. Liver tissue samples and blocks (paraffin and frozen), serum, cholangiocytes, and cholangiocyte supernatants (after incubation at 37°C for 4 hr) were collected as described.

Isolated Cholangiocytes and Hepatic Stellate Cells, and Cell Lines

Virtually pure cholangiocytes were obtained by immunoaffinity separation by using a monoclonal antibody, rat IgG2a (a gift from Dr. R. Faris, Brown University, Providence, RI), against an unidentified antigen expressed by all mouse cholangiocytes. Cholangiocytes were isolated from 3 different groups of animals (each group consisted of 4 animals for a total of 12 mice). For each cholangiocyte preparation, we used 4 mice due to the low yield obtained with one single animal. Following cholangiocyte isolation, approximately 10 million cholangiocytes were incubated at 37°C with 1 ml of HBS containing CaCl2 for 4 hrs before supernatants were collected.

Activated HSCs were isolated from frozen liver tissue samples by laser microdissection (LMD) using the Leica LMD7000 (Leica Biosystems; Buffalo Grove, IL) located at the Temple Health & Bioscience District (Temple, TX). OCT-frozen liver sections (10 μm) were
fixed to glass foiled poly ethylene naphthalate membrane slides and immunofluorescence for synaptophysin-9 (SYP-9, a maker of activated HSCs) \(^\text{30}\) was performed to visualize activated HSCs. Fluorescently labeled, activated HSCs were manually separated from unwanted cells using an ultraviolet laser. Microdissected HSCs were then collected into PCR tubes and RNA was isolated using the Arcturus\textsuperscript{®} PicoPure\textsuperscript{®} RNA Isolation Kit (Applied Biosystems; Waltham, MA) according to the manufacturers protocol.

The \textit{in vitro} experiments were performed in our immortalized murine biliary line (IMCL) \(^\text{31}\) as well as our human hepatic stellate cell line (hHSC) that were purchased from ScienCell Research Laboratories (Carlsbad, CA). IMCL and hHSC were maintained at standard conditions. IMCL and hHSC were treated with either 75 nM of mirVana\textsuperscript{™} miR-21 inhibitor or negative control (Thermo Fisher Scientific, Austin, TX) for 48 hr according to the manufacturers protocol before cell pellets were collected. The miR-21 inhibitor contains a sequence that is 100\% complementary to the sequence of active miR-21; therefore, binding between the inhibitor and miR-21 should be highly specific. As well, we have previously shown that use of this inhibitor \textit{in vitro} is able to increase hepatocyte and HSC expression of DR5 and Fas ligand, known targets of miR-21, by approximately two fold \(^\text{26}\). Unfortunately, since the inhibitor only binds to activated forms of miR-21 we are unable to accurately detect miR-21 expression levels since you may still get signals from the gene encoding miR-21, pri-miR-21, and pre-miR-21. For this reason, we based the specificity of our inhibitor on the expression of Bcl-2, a downstream target, in IMCL and hHSC (see Supplementary Figure 1).

**Evaluation of miR-21 Expression**

Isolation of miRNAs was performed in total liver and isolated cholangiocytes using the Ambion mirVana\textsuperscript{™} miRNA Isolation Kit from Life Technologies (Thermo Fischer Scientific; Waltham, MA). Single stranded cDNA was synthesized from 1 μg of RNA from the aforementioned samples using the TaqMan microRNA reverse transcription kit (Applied Biosystems; Waltham, MA) and was amplified by quantitative PCR (qPCR) using sequence specific primers from the TaqMan microRNA Assays on an Applied Biosystems Viia7 Real-Time PCR System (Thermo Fisher Scientific; Waltham, MA) according to the manufacturers protocol. The threshold cycle (\(C_T\)) is defined as the fractional cycle number at which the fluorescence passes the fixed threshold.

Control and late stage PSC patient samples (n=1 each) were obtained from Dr. Invernizzi under a protocol approved by the ethics committee by the Humanitas Research Hospital (Rozzano, Italy); the protocol was reviewed by the Veterans’ Administration IRB and R&D committee. The protocol was approved by the Texas A&M HSC Institutional Review Board. Total RNA was extracted from formalin-fixed, paraffin-embedded sections from samples obtained from 1 control and 1 PSC patients using the RNeasy FFPE kit (Qiagen; Valencia, CA). From these samples, miR-21 expression was evaluated as described above.
Assessment of Liver Morphology, Serum Chemistry, Intrahepatic Bile Duct Mass, and Biliary Proliferation

Hematoxylin and eosin (H&E) staining was performed in paraffin-embedded liver sections (4–5 μm, 10 different fields analyzed from each sample from 3 different animals). H&E stained liver sections were evaluated by a board-certified pathologist to determine the degree of lobular damage, hepatic necrosis, and portal inflammation.

The serum levels of alanine transaminase (ALT) and alkaline phosphatase (ALP) were measured in Sham WT, BDL WT, Sham miR-21−/−, and BDL miR-21−/− mice by a Dimension RxL Max Integrated Chemistry system (Dade Behring; Deerfield, IL) by the Chemistry Department, Baylor Scott & White Healthcare.

Intrahepatic bile duct mass (IBDM) was evaluated by semi-quantitative immunohistochemistry for cytokeratin-19 (CK-19, a cholangiocyte specific marker) 32. Biliary proliferation was evaluated in formalin-fixed, paraffin-embedded liver sections (4–5 μm, 10 different fields analyzed from each sample from 3 different animals) by semi-quantitative immunohistochemistry for Ki-67 33. We used qPCR to analyze PCNA, Bax, and cleaved Caspase-3 gene expression in RNA isolated from total liver, hHSC, and IMCL.

qPCR was performed using RT2 SYBR Green/ROX quantitative PCR master mix for the Applied Biosystems ViiA7 qPCR system (Life Technologies; Carlsbad, CA) according to the manufacturer’s protocol. The comparative C_T method (ΔΔC_T) was used for quantification of gene expression.

The proliferative rate of IMCL and hHSC was measured using the CellTiter 96 aqueous assay kit (Promega, Madison, WI). Transfected cells (10,000/well) were plated in 96-well plates (BD Biosciences) and incubated at 37°C, and cell proliferation was assessed after 48 hr as described 7.

Determination of Fibrosis, HSC Activation and Smad7 Expression

The fibrotic reaction was evaluated by qPCR for α-SMA, TGF-β1, and Collagen-1a expression in total RNA extracted from total liver tissue, LMD-isolated HSCs, and hHSC as described above. Collagen deposition was visualized in liver sections using Sirius Red staining as described 7. Activation of HSCs was visualized in liver sections by immunofluorescent staining for either SYP-9 or α-SMA, which was co-stained with CK-19 (only expressed by cholangiocytes) to visualize intrahepatic bile ducts. The role of cholangiocyte-derived factors on HSC activation was determined by incubating hHSCs in vitro with cholangiocyte supernatants from Sham WT, Sham miR-21−/−, BDL WT, or BDL miR-21−/− mice for 48 hr. Fibrosis gene expression was determined by qPCR for FN1 and Collagen-1a.

Since miR-21 has been shown to bind Smad-7 during an experimental model of non-alcoholic steatohepatitis 25, we measured in total liver and isolated cholangiocytes the expression of Smad-7 by qPCR.
Statistical Analysis

Data are expressed as mean ± SEM. Differences between groups were analyzed by the Student unpaired t test when 2 groups were analyzed, and by two-way ANOVA when more than 2 groups were analyzed.

RESULTS

miR-21 Expression is Increased During Cholestasis

Following BDL, there was a significant increase in miR-21 expression in total liver and isolated cholangiocytes when compared to Sham WT mice (Figure 1A–B). miR-21 expression was also increased in cholangiocytes from Mdr2−/− mice when compared to the corresponding WT mice, as well as in total liver extracted from a late stage PSC patient when compared to a normal liver sample. These findings suggest that during cholestatic injury, such as BDL or PSC, miR-21 levels are increased and may contribute to the progression of biliary injury and hepatic fibrosis.

Loss of miR-21 Ameliorates Liver Damage

Following BDL, there was moderate portal inflammation, ductal proliferation, and multifocal areas of necrosis when compared to Sham WT mice; however, the amount of damage observed in BDL miR-21−/− mice was greatly decreased when compared to BDL WT mice suggesting that miR-21 promotes liver damage during cholestatic injury (Figure 2A). No significant changes were noted in Sham miR-21−/− mice compared to Sham WT (Figure 2A). Similarly, the levels of the liver enzymes alanine aminotransferase (ALT) and alkaline phosphatase (ALP) were increased in serum from BDL WT compared to Sham WT mice; however, ALT and ALP serum levels were decreased in BDL miR-21−/− mice compared to BDL WT mice (Figure 2B). These studies indicate that the loss of miR-21 during BDL-induced cholestatic injury ameliorates liver damage associated with this injury.

Knockout of miR-21 Decreases BDL-Induced IBDM and Biliary Proliferation

Consistent with previous findings, following BDL there was a significant increase in IBDM when compared to Sham WT mice, which was reduced in BDL miR-21−/− compared to BDL WT mice (Figure 3A). No significant changes in IBDM were noted in Sham miR-21−/− mice when compared to Sham WT mice (Figure 3A). As shown by qPCR in total liver samples, BDL WT mice have increased PCNA expression when compared to Sham WT, however, there was decreased PCNA expression in Sham miR-21−/− and BDL miR-21−/− mice when compared to Sham WT and BDL WT, respectively (Figure 3B). Furthermore, the biliary expression of Ki-67 was increased in BDL WT mice compared to Sham WT, which was significantly decreased in BDL miR-21−/− mice when compared to BDL WT mice (Figure 3C). No significant changes were noted in the biliary expression of Ki-67 in Sham miR-21−/− mice when compared to Sham WT mice (Figure 3C). These findings demonstrate that during cholestasis the loss of miR-21 reduces IBDM and biliary proliferation associated with this injury.
Inhibition of miR-21 Decreases Cholangiocyte Proliferation In Vitro

*In vitro*, IMCL treated with miR-21 inhibitor had decreased proliferative capacity when compared to IMCL treated with control (Figure 4A). Consequently, IMCL that were treated with miR-21 inhibitor had increased expression of the apoptotic factors Bax and cleaved Caspase-3 (Figure 4B and 4C) when compared to control treated cells. These data indicate that inhibition of miR-21 in cholangiocytes leads to decreased proliferation and increased apoptosis *in vitro*.

Loss of miR-21 Decreases BDL-Induced HSC Activation and Fibrotic Reaction

Since the activation of HSCs is known to be a key player in the development of fibrosis, we analyzed HSC activation *in vivo* by staining for SYP-9 or α-SMA (markers of activated HSCs) and CK-19 (to visualize bile ducts) in liver sections by immunofluorescence. The expression of SYP-9 and CK-19 seemed unchanged between Sham WT and Sham miR-21−/− mice; however, the number of SYP-9 positive cells was increased in BDL WT, which was accompanied with increased IBDM. In BDL miR21−/− mice the number of SYP-9 positive cells was decreased compared to BDL WT, and this was also accompanied by decreased IBDM (Figure 5A). Staining for α-SMA and CK-19 showed the same trend as the SYP-9 staining (Figure 5B). Recently, studies have indicated that miR-21 plays a pro-fibrotic role during models of drug-induced cholestasis, and that inhibition of miR-21 can ameliorate this damage. In HSCs that were isolated from BDL WT we see increased expression of Collagen-1α when compared to Sham WT (Figure 5C). In contrast, in HSCs that were isolated from BDL miR-21−/− these parameters are decreased when compared to BDL WT (Figure 5C). No significant differences were noted between HSCs isolated from Sham WT and Sham miR-21−/−. These data further imply that increased miR-21 levels can contribute to increased fibrosis via HSC activation and fibrotic reaction.

Knockout of miR-21 Reduces the Amount of BDL-Associated Fibrosis

Next, we aimed to explore the role of miR-21 during BDL-induced fibrosis. In total liver samples, we found that the expression of α-SMA and TGF-β1 was significantly increased in BDL WT compared to Sham WT mice, but was significantly decreased in the livers of BDL miR-21−/− compared to BDL WT mice (Figure 6A and 6B). Furthermore, collagen deposition was increased in the livers of BDL WT mice compared to Sham WT mice, but was significantly decreased in the livers of BDL miR-21−/− mice when compared to BDL WT mice (Figure 6C). There was no significant difference between the expression of α-SMA and TGF-β1, and collagen deposition in the livers of Sham WT and Sham miR-21−/− mice. These findings, along with the above data indicating that miR-21 promotes HSC activation, suggest that the loss of miR-21 during cholestatic injury decreases the associated fibrotic reaction.

Inhibition of miR-21 Decreases HSC Proliferation and Fibrotic Reaction In Vitro

As demonstrated above, miR-21 seems to have a role in HSC activation and proliferation following bile duct ligation *in vivo* (Figure 5). To further validate these findings, we took hHSCs and treated them with either miR-21 inhibitor or control. Following treatment, proliferative capacity was decreased in hHSC treated with a miR-21 inhibitor when
compared to hHSC treated with control as demonstrated by MTS assay and PCNA expression (Figure 7A and 7B). Concomitantly, hHSCs treated with miR-21 inhibitor have increased Bax expression when compared to control treated cells (Figure 7C) indicating that the loss of miR-21 decreases HSC proliferation. We wanted to look further at the role of miR-21 during HSC activation since we previously noted that BDL miR-21−/− mice have decreased HSC activation when compared to BDL WT mice (Figure 5). By qPCR, we found that hHSCs treated with miR-21 inhibitor had decreased expression of α-SMA and MMP-9 when compared to control (Figure 7D). These findings suggest that the loss of miR-21 decreases the degree of HSC proliferation and fibrotic reaction, which could be an important target to halt cholestasis-induced hepatic fibrosis progression.

Loss of miR-21 Increases Smad-7 Expression In Vivo

Smad-7 has been reported as a target of miR-21, and since Smad signaling is involved in the promotion of hepatic fibrosis in both cholangiocytes and HSCs during models of cholestasis, we evaluated the expression of Smad-7 in our model. In BDL WT mice there was decreased expression of Smad-7 in both total liver and isolated cholangiocytes when compared to Sham WT (Figure 8A and 8B); however, BDL miR-21−/− mice showed increased Smad-7 when compared to BDL WT (Figure 8A and 8B). No significant changes in these factors were noted in Sham miR-21−/− when compared to Sham WT. This data demonstrates that miR-21 can regulate Smad-7 within cholangiocytes during cholestasis.

HSCs Treated with Supernatants from Cholangiocytes Lacking miR-21 Have Decreased Fibrotic Reaction In Vitro

We next evaluated the effect of cholangiocyte supernatant from Sham and BDL WT and miR-21−/− mice (that do not contain miR-21) on HSC activation. We incubated hHSCs with supernatants from cholangiocytes isolated from Sham WT, BDL WT, Sham miR-21−/−, and BDL miR-21−/− animals. There was increased FN-1 and Collagen-1α expression in hHSCs treated with cholangiocyte supernatants from BDL WT when compared to supernatant from Sham WT mice; however, these factors were decreased in hHSCs treated with cholangiocyte supernatants from BDL miR-21−/− animals compared to supernatant from BDL WT mice (Figure 9A and 9B). These data imply that cholangiocyte signaling strongly impacts HSC activation during cholestatic injury, and these pro-fibrotic factors may be regulated by miR-21.

DISCUSSION

To summarize these findings, in the BDL and Mdr2−/− cholestatic models miR-21 is upregulated in total liver and cholangiocytes. We have also shown that human PSC sample has increased miR-21 levels when compared to normal liver. Within cholangiocytes and HSCs, miR-21 promotes proliferation and fibrosis, and decreases apoptosis. Once activated, HSCs are able to secrete fibrosis-promoting factors that lead to increased hepatic damage. We demonstrated that miR-21 regulates HSC activation and Smad signaling, which in turn stimulates fibrogenesis. Loss of miR-21 leads to decreased biliary hyperplasia, HSC activation, and fibrosis that are associated with BDL.
Studies have shown that the expression of miR-21 (that is ubiquitously expressed throughout the body) is increased in cholangiocarcinoma (CCA) tissue when compared to normal biliary epithelium. We found that hepatic miR-21 levels are significantly increased in PSC when compared to control samples. Since chronic cholestasis can be a risk factor for developing CCA it is important to understand the role of miR-21 during liver injury.

Previous studies have shown that miR-21 is upregulated during models of chronic liver injury, inhibits apoptosis, and regulates cell survival. Increased miR-21 levels are commonly noted during cell proliferation and stress; however, it remains controversial as to which liver cell type is targeted by miR-21. Here we show that miR-21 expression is upregulated in cholangiocytes following BDL and in the Mdr2−/− model of PSC. Furthermore, the loss of miR-21 reduced the BDL-induced cholangiocyte proliferation compared to BDL WT mice. This is further highlighted where we show that treatment of IMCL with a miR-21 inhibitor leads to decreased proliferative activity, with a concomitant increase in apoptosis.

With regard to the possible target cells of the miR-21 inhibitor in our in vivo models, we propose that cholangiocytes and HSCs are two important target cells of the miR-21 inhibitor. In support of this, in the in vitro studies we determined that cholangiocytes and HSCs contain miR-21, and treatment with the miR-21 inhibitor impacted growth and activation in these cell types. However, in vivo we utilized a miR-21−/− mouse model, resulting in complete loss of miR-21 in all cells. Further in vivo studies are needed to pinpoint the target cells of the miR-21 inhibitor in normal and cholestatic models.

Currently, there is limited data regarding the role of miR-21 in HSCs during hepatic fibrosis, and no current literature exists regarding its role in cholangiocytes during cholestasis. One study found that HSCs containing miR-21 promote their own activation during injury; however, existing publications do not consistently report on miR-21 levels in activated HSCs. Our findings indicate that mice lacking miR-21 have decreased HSC activation when compared to BDL WT mice. On the basis of our findings, we propose that HSC activation is regulated by miR-21 during cholestatic injury, and activation can be further maintained by protecting the activated HSCs from apoptosis.

Previous work has shown that Smad signaling can influence HSC activation and cholangiocyte proliferation during models of cholestatic injury. Here we show that loss of miR-21 increases Smad-7 expression in both total liver and isolated cholangiocytes. These findings provide evidence that miR-21 increases fibrogenesis during hepatic injury by inhibiting the inhibitory Smad-7.

It is known that many hepatic cell types can interact and signal to one another via paracrine mediators to influence disease progression. Specifically, it has been noted that cholangiocytes can signal to HSCs to regulate their activation. Our data indicate that hHSCs treated with supernatant isolated from cholangiocytes lacking miR-21 have decreased fibrotic reaction, further verifying that cholangiocyte-secreted factors can influence HSC response during cholestatic injury. Previously, it has been shown that the lumen of intrahepatic bile ducts of cystic liver contains exosome-like vesicles, and these...
Exosomes are cholangiocyte-derived. As well, rats that undergo partial hepatectomy show increased serum levels of exosome-like vesicles containing miR-21 when compared to normal rats. Based on this work, we hypothesize that during cholestatic injury cholangiocytes may secrete exosomes containing miR-21 that can increase HSC activation.

Our data shed further light on the impact that miR-21 has on hepatic fibrosis, specifically in the realm of cholestatic injury. It has been proposed that miR-21 is strongly upregulated in HSCs during thioacetamide- and carbon tetrachloride-induced hepatic fibrosis. These studies along with our findings promote the idea that miR-21 regulates hepatic fibrosis through the modulation of HSC activation and proliferation. Aside from hepatic fibrosis, we provide evidence that miR-21 (i) is upregulated in cholangiocytes following injury, (ii) enhances cholangiocyte proliferation, and (iii) hinders cholangiocyte apoptotic processes. Increased HSC activation and cholangiocyte proliferation may be driven by miR-21 regulation of Smad-7. In this regard, modulation of miR-21 during fibrosis-related hepatic injury may be useful. Further elucidation of the pathways that are modulated by miR-21 during these processes is essential to uncovering therapeutic targets.

**Supplementary Material**

Refer to Web version on PubMed Central for supplementary material.

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**Abbreviations**

- **ALP** alkaline phosphatase
- **ALT** alanine transaminase
- **α-SMA** alpha-smooth muscle actin
- **BDL** bile duct ligation
- **CK-19** cytokeratin-19
- **Collagen-1α** collagen, type I, alpha 1
- **FN-1** fibronectin-1
- **GAPDH** glyceraldehyde-3-phosphate dehydrogenase
hHSC  human hepatic stellate cell line
IBDM  intrahepatic bile duct mass
IMCL  immortalized murine biliary cell line
miR-21  microRNA-21
PCNA  proliferating cell nuclear antigen
real-time PCR  qPCR
Smad-7  small mothers against decapentaplegic 7
SYP-9  synaptophysin-9
TGF-β1  transforming growth factor-β1
WT  wild-type

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Figure 1.
Evaluation of miR-21 expression following BDL. Following BDL, miR-21 levels are increased in total liver and isolated cholangiocytes (A, B). In Mdr2−/− mice cholangiocytes have increased miR-21 expression (C). Total liver samples from PSC patients have increased miR-21 levels compared to normal patient samples (D). Data are expressed as means ± SEM. n = 10 reactions in total RNA collected from total liver and cholangiocytes from 8 animals per group for qPCR. n = 3 reactions per sample in total RNA collected from total liver from 1 human control and 1 human PSC sample. *p<0.05 versus Sham WT, WT, Human Control.
Figure 2.
Assessment of liver injury. Loss of miR-21 ameliorates BDL-induced necrosis, lobular damage, and portal inflammation as indicated by H&E staining (A). Yellow dashed line indicates areas of necrosis while the yellow arrows point to bile ducts (A). BDL WT mice show increased serum levels of ALT and ALP when compared to BDL WT; however, these parameters are decreased in BDL miR-21−/− mice when compared to BDL WT mice (B). Data are expressed as means ± SEM. n = 3 reactions in serum collected from 8 animals each per animal group for serum chemistry. *p<0.05 versus Sham WT; #p<0.05 versus BDL WT. Representative images are shown for H&E. Original magnification, 20X.
Figure 3.
Measurement of IBDM and biliary proliferation. Following BDL, there is increased IBDM compared to Sham WT (A). The loss of miR-21 reduces the degree BDL-induced IBDM compared to BDL WT (A). Following BDL, mice have increased levels of PCNA, which is significantly decreased in BDL miR-21−/− mice compared to BDL WT (B). Immunohistochemistry and semi-quantitative analysis are shown for the proliferative marker Ki-67 (C). There is increased biliary Ki-67 expression in BDL WT compared to Sham WT, but the loss of miR-21 decreases BDL-induced biliary Ki-67 expression (C). Data are expressed as means ± SEM. n = 7 reactions in total RNA collected from 8 animals per group for qPCR, n = 10 pictures from 6 animals per group for immunohistochemistry. *p<0.05 versus Sham WT; #p<0.05 versus BDL WT. Representative images are shown for CK-19 and Ki-67 immunohistochemistry. Original magnification, 20X.
Figure 4.
Evaluation of IMCL proliferation and apoptosis. Following treatment with miR-21 inhibitor, IMCL have decreased proliferation, as shown by MTS assay, and increased Bax and cleaved Caspase-3 gene expression when compared to control treated (A, B, C). Data are expressed as mean ± SEM. n = 6 reactions in total RNA from 12 sets of cells for qPCR, n = 10 experiments per group for MTS assay. *p<0.05 versus IMCL control.
Figure 5.
Evaluation of HSC activation and fibrotic reaction. As shown by immunofluorescence co-stain for SYP-9 (green) and CK-19 (red) there is increased HSC activation and IBDM in BDL WT compared to Sham WT, but this is decreased in BDL miR-21$^{-/-}$ mice when compared to BDL WT, respectively (A). Immunofluorescent co-stain for α-SMA (green) and CK-19 (red) show increased HSC activation and IBDM in BDL WT compared to Sham WT, but this is decreased in BDL miR-21$^{-/-}$ when compared to BDL WT (B). In HSCs that are isolated from BDL WT there is a significant increase in Collagen-1α expression when compared to HSCs isolated from Sham WT; however, Collagen-1α expression is significantly decreased in HSCs isolated from BDL miR-21$^{-/-}$ when compared to HSCs isolated from BDL WT (C). Data are expressed as means ± SEM. n=6 reactions in total RNA from 6 animals per group for qPCR. *p<0.05 vs. Sham WT; #p<0.05 vs. BDL WT. Representative images are shown for SYP-9 and CK-19 immunofluorescence. Original magnification, 20X.
Figure 6.
Determination of fibrotic reaction. In BDL WT there is a significant increase in α-SMA and TGF-β1 gene expression compared to Sham WT; however, this is significantly decreased in BDL miR-21−/− when compared to BDL WT (A, B). Collagen deposition is significantly increased following BDL when compared to Sham WT; however, the loss of miR-21 blunts this effect (C). Data are expressed as means ± SEM. n = 6 reactions in total RNA from 8 animals per group for qPCR, n = 20 pictures from 6 animals used for Sirius Red staining. *p<0.05 versus WT; #p<0.05 versus BDL WT. Representative images are shown for Sirius Red staining. Original magnification, 20X.
Figure 7.
Evaluation of hHSC miR-21 proliferation, apoptosis and fibrotic reaction, *in vitro*. hHSCs treated with miR-21 inhibitor have decreased proliferation, increased Bax, and decreased α-SMA and MMP-9 gene expression when compared to control treated (B, C, D, E). Data are expressed as means ± SEM. n = 6 reactions from 12 sets of cells for qPCR, n = 10 experiments for MTS assay. *p<0.05 versus hHSC control.
Figure 8.
Evaluation of Smad-7 expression. In total liver and isolated cholangiocytes from BDL WT mice there is increased/ decreased Smad-7, as shown by qPCR, when compared to Sham WT (A, B). However, total liver and isolated cholangiocytes from BDL miR-21−/− mice show increased Smad-7 when compared to BDL WT (A, B). n = 9 reactions from total RNA from 6 animals for qPCR. *p<0.05 vs. Sham WT; #p<0.05 vs. BDL WT.
Figure 9.
Determination of hHSC fibrotic reaction, in vitro. hHSCs treated with supernatants extracted from cholangiocytes isolated from BDL WT mice show increased FN-1, and Collagen-1a expression when compared to hHSCs treated with supernatants from Sham WT mice (A, B); however, these parameters were decreased in hHSCs treated with cholangiocyte supernatants from BDL miR-21$^{-/-}$ mice when compared to BDL WT mice (A, B). Data are expressed as means ± SEM. n = 6 reactions from 12 sets of cells for qPCR. *p<0.05 versus hHSCs + WT cholangiocyte supernatants; #p<0.05 versus hHSCs + BDL WT cholangiocyte supernatants.