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ORIGINAL ARTICLE

Liver microRNA-21 is overexpressed in non-alcoholic steatohepatitis and contributes to the disease in experimental models by inhibiting PPAR α expression

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ABSTRACT

Objective Previous studies suggested that microRNA-21 may be upregulated in the liver in non-alcoholic steatohepatitis (NASH), but its role in the development of this disease remains unknown. This study aimed to determine the role of microRNA-21 in NASH.

Design We inhibited or suppressed microRNA-21 in different mouse models of NASH: (a) low-density lipoprotein receptor-deficient (*Ldlr*^{-/-}) mice fed a high-fat diet and treated with antagomir-21 or antagomir control; (b) microRNA-21-deficient and wild-type mice fed a methionine-choline-deficient (MCD) diet; (c) peroxisome proliferation-activator receptor α (PPAR α)-deficient mice fed an MCD diet and treated with antagomir-21 or antagomir control. We assessed features of NASH and determined liver microRNA-21 levels and cell localisation. MicroRNA-21 levels were also quantified in the liver of patients with NASH, bland steatosis or normal liver and localisation was determined.

Results Inhibiting or suppressing liver microRNA-21 expression reduced liver cell injury, inflammation and fibrogenesis without affecting liver lipid accumulation in *Ldlr*^{-/-} fed a high-fat diet and in wild-type mice fed an MCD diet. Liver microRNA-21 was overexpressed, primarily in biliary and inflammatory cells, in mouse models as well as in patients with NASH, but not in patients with bland steatosis. PPAR α , a known microRNA-21 target, implicated in NASH, was decreased in the liver of mice with NASH and restored following microRNA-21 inhibition or suppression. The effect of antagomir-21 was lost in PPAR α -deficient mice.

Conclusions MicroRNA-21 inhibition or suppression decreases liver injury, inflammation and fibrosis, by restoring PPAR α expression. Antagomir-21 might be a future therapeutic strategy for NASH.

INTRODUCTION

Non-alcoholic fatty liver disease (NAFLD) is a condition ranging from simple triglyceride accumulation in the liver (steatosis) to steatosis combined with lobular inflammation and hepatocyte injury (clarification and ballooning) with or without

Significance of this study

What is already known on this subject?

- MicroRNAs are attractive therapeutic targets since they can be modulated in patients.
- Microarray studies suggested that microRNA-21 could be overexpressed in the liver in patients with non-alcoholic steatohepatitis (NASH).
- Peroxisome proliferation-activator receptor α (PPAR α) is a target of microRNA-21 and is implicated in NASH.

What are the new findings?

- MicroRNA-21 contributes to key features of NASH, that is, cell injury, inflammation and fibrosis, through PPAR α inhibition.
- Using different mouse models of NASH, we demonstrate that microRNA-21 blockade prevents NASH development.
- MicroRNA-21 is overexpressed in the liver of patients with NASH, but not in those with bland steatosis.
- Liver microRNA-21 in NASH is primarily expressed in inflammatory and biliary cells.

How might it impact on clinical practice in the foreseeable future?

- MicroRNA-21 antagonism might be useful in the future in the treatment of NASH.

fibrosis. The latter is referred to as non-alcoholic steatohepatitis (NASH).¹ NAFLD and NASH are increasingly relevant public health issues owing to their close association with the worldwide epidemics of diabetes and obesity.² NAFLD/NASH is one of the most common chronic liver diseases. Indeed, in the USA, 39% of newly identified cases of chronic liver disease had NAFLD.³ NASH has potential for fibrosis, cirrhosis decompensation and hepatocellular carcinoma. Despite its prevalence and severity, therapeutic options for NASH are limited.¹ Therefore, understanding the mechanisms



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leading to hepatic inflammation and fibrosis in NAFLD is of great importance for the diagnosis and treatment of NASH.

MicroRNAs are a family of short non-coding RNAs (19–22 nucleotides in length) that regulate gene expression at the post-transcriptional level via messenger RNA degradation or translational inhibition.^{4–6} MicroRNA microarray analysis identified microRNA-21 (miR-21) as one of the most upregulated micro-RNAs in the liver of patients with NASH.⁷ This upregulation was subsequently confirmed.⁸ In two mouse models of NASH (choline-deficient and amino acid-defined diet and high-fat diet (HFD)), miR-21 was also overexpressed in the liver.^{8–9} However, these studies did not address whether or not liver miR-21 upregulation was a cause or a consequence of NASH. In addition, the cellular localisation of miR-21 has not yet been elucidated.

Therefore, the present study aimed to determine the role of miR-21 in the pathogenesis of NASH. For this purpose, we used complementary approaches: (a) a pharmacological inhibition of miR-21 using antagomir-21 in a genetic model of NASH (low-density lipoprotein receptor-deficient (*Ldlr*^{-/-}) mice fed an HFD); and (b) a genetic deletion of miR-21 (*miR-21*^{-/-} mice) in mice exposed to a dietary model of NASH (methionine-choline-deficient (MCD) diet). We also determined whether miR-21 effect implicates peroxisome proliferation-activator receptor α (PPAR α), a known miR-21 target, using PPAR α -deficient (*Ppara*^{-/-}) mice.

MATERIALS AND METHODS

Mouse models of NASH

All experiments were performed in accordance with the guidelines formulated by the European Community for experimental animal use (L358-86/609EEC) and were approved by the French Ministry of Agriculture (agreement no. A75-15-32). All animals received human care according to the criteria outlined in the *Guide for the Care and Use of Laboratory Animals* prepared by the National Academy of Sciences and published by the National Institutes of Health.

All mice were on a C57BL/6 background. Different NASH models were used: *Ldlr*^{-/-} mice fed an HFD, and *miR-21*^{-/-} mice fed an MCD diet.

Eight-week-old male *Ldlr*^{-/-} mice (Charles River, L'Isle d'Abresle, France) were put on an HFD containing 1.25% cholesterol, 15% cacao butter and 0% cholate (Safe Diets) for 14 weeks (figure 1A). When 12 and 17 weeks old, mice received three retroorbital intravenous injections (once a day for three consecutive days) of antagomir-21 (sequence 5'-UCAACAUCAG UCUGAUAAGCUA-3'; 16 mg/kg body weight, VBC Biotech, Vienna, Austria), or antagomir control (sequence 5'-AAGGC AAGCUGACCCUGAAGUU-3'; 16 mg/kg body weight; VBC Biotech Vienna, Austria) or phosphate buffer saline (PBS). This design was adapted from previous reports showing that antagomirs are able to induce a potent and prolonged miR inhibition.^{10–12} Preliminary experiments showed that miR-21 expression was more robustly decreased by antagomir-21 given at a dose of 16 than 8 mg/kg body weight (data not shown). C57BL/6 wild-type (WT) untreated mice were used as controls. All mice groups were euthanased at the age of 22 weeks.

Nine-week-old male WT or *miR-21*^{-/-} mice were put for 10 weeks on MCD containing 18.3% methionine-deficient amino acid mix, 10% sucrose, 10% lard, 5% cellulose, 1% choline-deficient vitamin mix, 3.5% mineral mix and 53% cornstarch (SSNIFF Diet).¹³ C57BL/6 WT mice under chow diet served as controls. All mice were euthanased at the age of 19 weeks.

Sixteen-week-old male WT or *Ppara*^{-/-} mice were put for 5 weeks on MCD diet. When 17 weeks old, mice received three retroorbital intravenous injections of antagomir-21 or antagomir control (16 mg/kg body weight). C57BL/6 (WT) mice under chow diet served as controls. All mice were euthanased at the age of 21 weeks.

On the day of the sacrifice, mice were sedated with 2.5% iso-fluorane and blood was collected from the inferior vena cava into a 25 gauge \times 1' needle pre-coated with 3.8% sodium citrate. Blood was spun at 1500 g for 15 min to prepare platelet poor plasma that was then stored at -80°C . The mice were then euthanased. Livers were isolated and snap frozen in liquid nitrogen and stored at -80°C or fixed in formalin and paraffin-embedded.

RNA isolation and miR-21 quantification

Total RNA was extracted from snap frozen human or mouse liver samples with Trizol reagent according to the manufacturer's instructions (Invitrogen, Paris, France). MiR-21 levels were quantified using TaqMan assay (Tm 00397, Applied Biosystems) following the manufacturer's instructions. miR-21 expression was normalised to U6 snRNA (Tm 001973, Applied Biosystems) for mouse and human samples, together with RNU6B (Tm 001093; Applied Biosystems) for human samples and snoRNA234 (Tm001234; Applied Biosystems) for mouse samples as endogenous control. Relative expression was calculated using the 2-delta-delta CT method followed by geometric average, as recommended.¹⁴

In situ hybridisation

In situ hybridisation was adapted from a previously described method and is described in detail in the online supplementary methods.¹⁵ Liver samples from *miR-21*^{-/-} mice served as negative control and failing myocardium induced by a myocardial infarction as a positive control (online supplementary figure S1).¹⁶

Hepatic inflammatory cells sorting and analysis by flow cytometry

Mouse livers were perfused in vivo with PBS and then collected, weighted and minced with sterile scissors in PBS supplemented with 1% fetal calf serum (100 mg liver tissue in 1 mL PBS). Liver cell suspension was filtered through a 40 μm membrane. Polymorphonuclear cells were then isolated by density-gradient centrifugation over endotoxin-free Ficoll 1083 (Sigma-Aldrich) for 30 min at 500 g without brake at room temperature. Pellets and polymorphonuclear cells rings were collected separately, washed and counted. Cells from pellets were stained with anti-Epcam allophycocyanin antibodies (eBiosciences anti-CD326, #17-5791-82) for 30 min at 4°C . Cells from the rings were stained with anti-mouse CD11b-fluorescein-isothiocyanate (eBiosciences, #130-081-201), anti-NK-1.1-phycoerythrin (BD Biosciences, #553165), anti-Ly6G-phycoerythrin (BD Biosciences, #551461), anti-CD45-PerCP (BD Biosciences, #557235) and anti-CD3-PE-Alexa700 (eBiosciences, #17-0032-82) antibodies for 30 min at 4°C . Cells were then washed and resuspended in PBS supplemented with 1% fetal calf serum and extemporaneously with 1 $\mu\text{g}/\text{mL}$ 4',6-diamidino-2-phenylindole. Phenotyping of polymorphonuclear cells rings was performed on a LSR II flow cytometer (BD Biosciences, San Jose, California, USA). CD45+ subpopulations or Epcam+ cells were isolated for quantification of miR-21 expression using an ARIA II SORP cell sorter (BD Biosciences). Post-acquisition analysis was performed using FlowJo software.

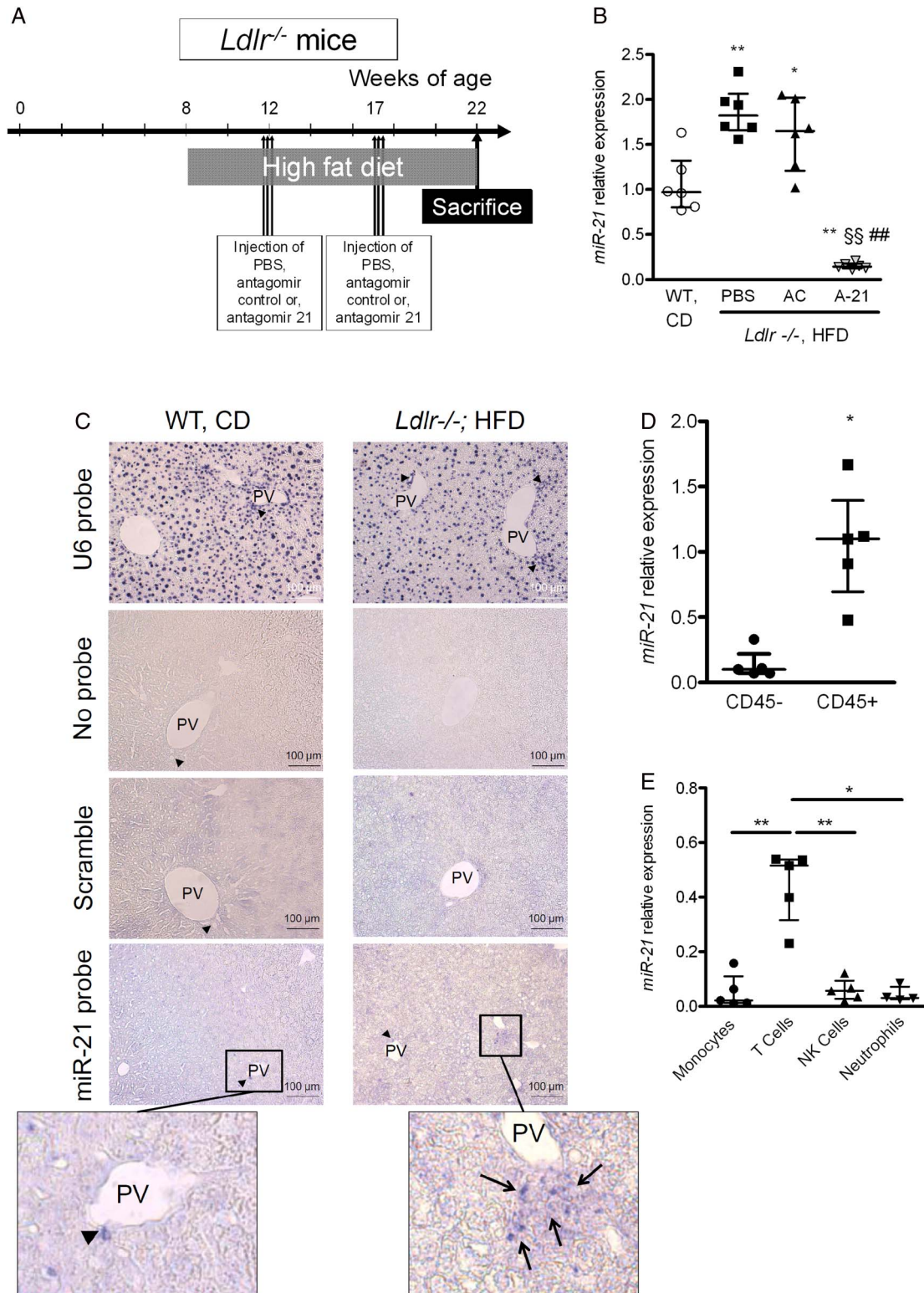


Figure 1 MicroRNA-21 (miR-21) was overexpressed in the liver of low-density lipoprotein receptor-deficient (*Ldlr*^{-/-}) mice fed a high-fat diet (HFD), primarily in inflammatory and biliary cells. (A) Mouse study design. (B) miR-21 expression in the liver of wild-type (WT) mice fed a chow diet (CD) and of *Ldlr*^{-/-} mice fed an HFD treated with the vehicle (phosphate buffer saline (PBS)), antagomir control (AC) or antagomir-21 (A-21). ***p*<0.01 versus WT mice fed a CD; §§*p*<0.01 versus *Ldlr*^{-/-} mice treated with PBS; ##, *p*<0.01 versus *Ldlr*^{-/-} mice treated with AC; (C) in situ hybridisation. miR-21 was detected in liver inflammatory (arrow) and biliary cells (arrow head) from *Ldlr*^{-/-} mice fed an HFD. Original magnification ×200. (D) miR-21 expression in sorted liver inflammatory (CD45⁺) and non-inflammatory (CD45⁻) cells from *Ldlr*^{-/-} mice fed an HFD. (E) miR-21 expression in subpopulations of liver inflammatory (CD45⁺) cells from *Ldlr*^{-/-} mice fed an HFD. **p*<0.05; ***p*<0.01. Data are given as median (horizontal bar) and IQR (error bar). PV, portal vein.

Metabolic analyses

Serum alanine aminotransferase (ALT) and aspartate aminotransferase (AST) levels, and fasting glucose, total cholesterol, high-density lipoprotein (HDL)-cholesterol and triglyceride levels were determined using commercial enzymatic kits on a DxC800 analyzer (Beckman-Coulter, Villepinte, France).

Histology

Human liver samples

Paraffin-embedded liver sections (3 μm thick) were stained with H&E (haematoxylin, RAL diagnostics; and eosin, RAL diagnostics) and Picro-sirius (Sirius red, RAL diagnostics; picric acid, VWR Prolabo). The steatosis score assessed the quantities of large or medium-sized lipid droplets, but not foamy microvesicles, from 0 to 3: S0: <5%; S1: 5–33%; S2: 34–66%; and S3: >67%. Ballooning was graded from 0 to 2: 0, normal hepatocytes with cuboidal shape and pink eosinophilic cytoplasm; 1, presence of clusters of hepatocytes with a rounded shape and pale cytoplasm usually reticulated; and 2, same as grade 1 with some enlarged hepatocytes, at least twofold that normal cells.¹⁷ Lobular inflammation was defined as a focus of two or more inflammatory cells within the lobule. Foci were counted at 20 \times magnification: 0, none; 1, ≤ 2 foci per 20 \times ; and 2, >2 foci per 20 \times field. Stage of fibrosis was assessed using the score described by NASH-CRN as follows: stage 0 (F0), none; stage 1 (F1), perisinusoidal zone 3 or portal fibrosis; stage 2 (F2), perisinusoidal and periportal fibrosis without bridging; stage 3 (F3), bridging fibrosis; and stage 4 (F4), cirrhosis.¹⁸

Mice liver samples

Paraffin-embedded liver sections (3 μm thick) were stained with H&E (haematoxylin, RAL diagnostics; and Eosin, RAL diagnostics), Picro-sirius (Sirius red, RAL diagnostics; picric acid, VWR Prolabo), rabbit anti-CD3 antibody (DAKO, #A0452) and a monoclonal rat anti-mouse CD68 antibody (AbD Serotec, MCA1957). Steatosis was defined as the percentage of hepatocytes containing large or medium-sized lipid droplets. Steatosis was scored on H&E stained sections by a single experienced pathologist (VP) who was blinded to mice groups. The following scale was used: 0 (<5%), 1 (5–33%), 2 (34–66%) and 3 (>67%). The Sirius red positive area was measured in at least 10 fields without vessels or capsule (original magnification, $\times 200$) and quantified using ImageJ software. Inflammatory cells (CD3⁺ or CD68⁺) were counted in ≥ 15 microscope views (original magnification, $\times 200$).

Frozen liver sections (4 μm) were stained with the neutral lipid marker Oil Red O (Sigma-Aldrich) to confirm steatosis.

Hepatic collagen analysis

Hepatic total collagen content was determined using the total collagen assay (Quickzyme Biosciences, UK) according to the manufacturer's instructions.

Gene expression analysis

cDNA synthesis was performed with QuantiTect Reverse Transcription Kit (Qiagen). PCR was performed on an ABI StepOne Plus with the use of Power SYBR Green PCR Master Mix (Eurogentec). Liver expression of all genes has been normalised using three housekeeping genes (*Gapdh*, *Hprt* and *Ppia*). The primers used are described in online supplementary table S1. Relative expression was calculated using the 2-delta-delta CT method followed by geometric average, as recommended.^{19 20}

Hepatic lipid analysis

Fifty micrograms of frozen liver tissue was homogenised by sonication in 1 mL of acetone. After overnight incubation at 4°C, samples were centrifuged at 4000 g for 10 min. Triglyceride (Diasys, Holzheim, Germany) and total cholesterol (Biomérieux, Craponne, France) contents were then measured in the supernatant using enzymatic kits according to manufacturers' instructions. Triglycerides and total cholesterol content were normalised to protein content determined by Dc Protein Assay (BioRad, Marnes la Coquette, France).

Liver cytokine and chemokine protein level

Cytokines were quantified on total liver protein extracts. Frozen liver tissue samples were homogenised in TNE buffer (10 mmol/L Tris-HCl (pH 7.5), 1% Nonidet P40, 150 mmol/L NaCl, 1 mmol/L EDTA) containing a cocktail of protease and phosphatase inhibitors (Roche). The suspension was centrifuged at 20 000 g for 20 min at 4°C. The supernatant was then stored at -80°C before use. Levels of tumour necrosis factor (TNF)- α and monocyte chemoattractant protein-1 (MCP-1) in liver extracts were measured using a cytometric bead array (Mouse Flowcytomix, eBiosciences). Cytokines levels were normalised to protein content determined by Dc Protein Assay (BioRad, Marnes-la-Coquette, France).

Western blot

About 50 mg of frozen mouse liver tissue samples were homogenised in 650 μL of TNE buffer (10 mmol/L Tris-HCl (pH 7.5), 1% Nonidet P40, 150 mmol/L NaCl, 1 mmol/L EDTA) containing a cocktail of protease and phosphatase inhibitors (Roche). Lysates were incubated at 4°C for 45 min and then centrifuged at 12 000 g for 5 min. Protein content was quantified in the supernatant using BCA Protein Assay (Thermo Fisher). Lysates were mixed with the reducing sample buffer (Bio-Rad) for electrophoresis and subsequently transferred onto polyvinylidene fluoride membranes (Thermo Fisher). Equal loading (50 μg) was verified using Ponceau red solution. Membranes were incubated with anti-PPAR α (1:1000, Ab24509, Abcam) antibody. After secondary antibody incubation (anti-rabbit, 1:3000; Amersham, GE Healthcare, Courtaboeuf, France), immunodetection proceeded using an enhanced chemiluminescence kit (Clarity Western ECL Substrate, BioRad, France) and bands were revealed using the Las-4000 Imaging System and Image MultiGauge software (Fujifilm, Tokyo, Japan). After initial immunodetection, membranes were stripped of antibodies and reprobed with mouse anti-GAPDH antibody (1:20 000; Millipore) and secondary antibody (anti-mouse, 1:3000; Amersham GE healthcare).

Measurement of hepatic total ketone bodies

Ketones were measured by the Williamson method.²¹ Briefly, frozen liver samples (200 mg) were deproteinised using perchloric acid 40%. Neutralised supernatants were used to measure acetoacetate and beta-hydroxybutyrate by a colorimetric method using beta-hydroxybutyrate deshydrogenase (Roche). The generation or the consumption of reduced nicotinamide adenine dinucleotide was measured by changes in absorbance at 340 nm. Results were expressed as micromole of ketone bodies per gram of liver tissue.

Patients

Twenty-five patients who underwent a percutaneous liver biopsy between 2008 and 2013 were selected from a pathological

database: 6 had no or mild abnormalities at liver histological examination, 8 had bland steatosis and 11 had NASH as defined using the SAF (for steatosis, activity and fibrosis) score.¹⁷ None of the patients had detectable anti-hepatitis C virus antibodies or HBs antigen. Alcohol consumption was below 21 drinks on average per week in men and 14 drinks on average per week in women.¹ None of the patients had a history of TIPS or of liver transplantation. Liver biopsy specimens were reviewed by a single experienced pathologist (VP) who was blinded to the clinical and laboratory data.

Statistics

Quantitative variables were expressed as median (interquartile range, IQR). Comparisons between groups of independent quantitative variables were performed using Mann–Whitney test. Spearman’s correlation coefficient was used to test the correlation between liver miR-21 expression levels and pathological features. All tests were two-sided and used a significance level of 0.05. Data analysis was performed with SPSS V.17.0 (SPSS, Chicago, Illinois, USA) and figures were designed using GraphPad Prism 5 software.

RESULTS

Expression of miR-21 is increased in biliary and inflammatory cells in the liver of *Ldlr*^{-/-} mice under HFD

MiR-21 expression was increased in the liver of *Ldlr*^{-/-} mice fed an HFD compared with WT mice under chow diet (figure 1B). Using in situ hybridisation, we observed that miR-21 was

primarily expressed in biliary and inflammatory cells in the liver of *Ldlr*^{-/-} mice under HFD (figure 1C). In accordance with real-time PCR results, the staining frequency and intensity of miR-21 was higher in *Ldlr*^{-/-} mice under HFD than in controls. In these controls, miR-21 staining was faint and restricted to biliary cells. To ascertain these observations, miR-21 expression was determined in liver inflammatory (CD45⁺) versus non-inflammatory (CD45⁻) cells from *Ldlr*^{-/-} mice under HFD as well as in subpopulations of inflammatory cells. MiR-21 expression was overexpressed in CD45⁺ cells and particularly in T cells (figure 1D, E).

In order to determine the potential role of miR-21 in NASH, we used an antagomir strategy. As shown in figure 1B, antagomir-21 strongly decreased liver miR-21 expression in *Ldlr*^{-/-} mice under HFD. When compared with antagomir control, antagomir-21 had no effect on serum fasting glucose, total cholesterol, HDL and triglyceride levels (see online supplementary table S2). These results thus establish the use of antagomir-21 in *Ldlr*^{-/-} mice under HFD as a valid strategy to investigate the role miR-21 in NASH.

Antagomir-21 reduces liver injury, liver inflammation and liver fibrosis but not liver lipid accumulation in *Ldlr*^{-/-} mice under HFD

As expected in this mouse model of NASH, markers of liver injury (serum AST and ALT; figure 2A, B), of liver inflammation (liver MCP-1 and TNF-α gene and protein expression, liver CD3 and CD68 cell staining; figure 2C–J), of liver fibrosis

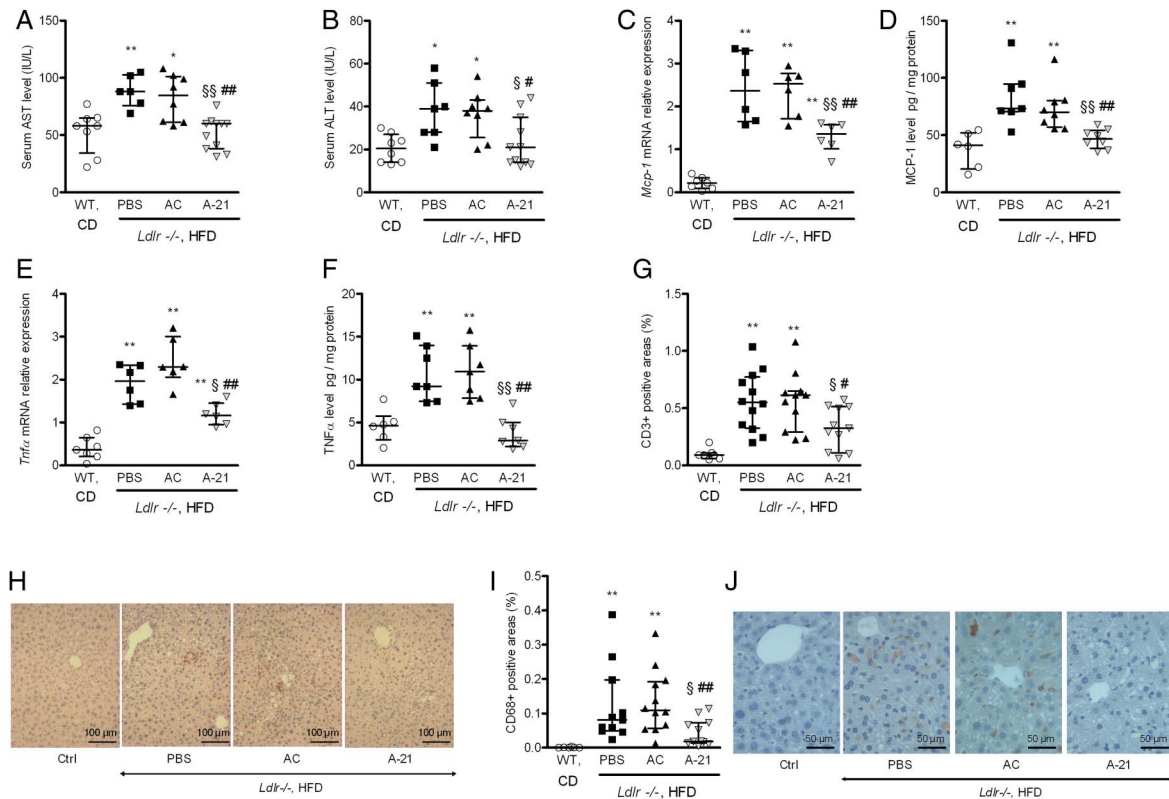


Figure 2 Antagomir-21 reduced liver injury and liver inflammation in the liver of low-density lipoprotein receptor-deficient (*Ldlr*^{-/-}) mice fed a high-fat diet (HFD). Antagomir-21 (A-21) reduced serum aspartate aminotransferase (AST) (A) and alanine aminotransferase (ALT) (B) levels, liver monocyte chemoattractant protein-1 (MCP1) (C and D) and tumour necrosis factor (TNF)-α (E and F) gene and protein expressions, as well as CD3⁺ (G, quantification; H, representative images, original magnification ×200) and CD68⁺ (I, quantification; J, representative images, original magnification ×400) cell infiltrates. *p<0.05, **p<0.01 versus wild-type (WT) mice fed a chow diet (CD); §p<0.05, §§p<0.01 versus *Ldlr*^{-/-} mice fed an HFD treated with phosphate buffer saline (PBS); #p<0.05, ##p<0.01 versus *Ldlr*^{-/-} mice fed an HFD treated with antagomir control (AC). Data are given as median (horizontal bar) and IQR (error bar). Samples were randomly selected in each group of mice.

(collagen staining with Sirius red, collagen quantification and gene expression of the fibrogenic-related genes *Tgfb* and *collagen-1 α 2*; figure 3) and of liver lipid accumulation (liver triglycerides and total cholesterol levels and histological evaluation; see online supplementary figure S2) were higher in *Ldlr*^{-/-} mice under HFD than in WT mice under chow diet.

Antagomir-21 strongly reduced markers of liver cell injury (figure 2A, B), of hepatic inflammation (figure 2C–J) and of liver fibrosis (figure 3), but had no effect on liver lipid accumulation or on fatty acid β -oxidation (liver gene expression of the β -oxidation-related genes carnitine palmitoyltransferase 1 (*Cpt1*) and acyl-coenzyme A oxidase 1 (*Acox1*) and liver total ketone bodies) (see online supplementary figure S2).

MiR-21^{-/-} mice under MCD diet have lower liver injury, liver inflammation and liver fibrogenic genes expression, but no change in liver lipid accumulation

We next analysed a second complementary model of NASH induced by MCD diet in *miR-21*^{-/-} versus WT mice (figure 4A). As shown in figure 4B, liver *miR-21* was increased by MCD diet in WT mice but remained undetectable in *miR-21*^{-/-} mice. Fasting serum glucose, total cholesterol, HDL and triglyceride levels were not different between *miR-21*^{-/-} and WT mice fed an MCD diet (see online supplementary table S3).

As expected in this mouse model of NASH, markers of liver injury (serum AST and ALT; figure 5A, B), of liver inflammation (liver MCP-1 and TNF- α expression, liver CD3 and CD11b cell number; figure 5C–H), of liver fibrosis (collagen deposition and *Tgfb* and *collagen-1 α 2* gene expression; figure 6) and of liver lipid accumulation (liver triglycerides and total cholesterol levels and histological evaluation; see online supplementary figure S3)

were increased in WT mice fed an MCD diet compared with WT mice fed a chow diet.

When compared with WT mice fed an MCD diet, *miR-21*^{-/-} mice fed an MCD diet had decreased liver injury and hepatic inflammation (figure 5). Although liver expression of the fibrogenic-related genes *Tgfb* and *collagen-1 α 2* was lower in *miR-21*^{-/-} than in WT mice fed an MCD diet, there was no significant difference in liver collagen deposition (figure 6). Liver lipid accumulation and liver expression of the fatty acid β -oxidation-related genes *Cpt1* and *Acox1* were not different between *miR-21*^{-/-} and WT mice fed an MCD diet (see online supplementary figure 3A–F). Liver total ketone bodies were slightly lower in *miR-21*^{-/-} than in WT mice fed an MCD diet (see online supplementary figure S3G).

PPAR α is responsible for miR-21 effect in NASH

To explore the molecular mechanisms through which miR-21 exerts its function in NASH, we used the miRWalk algorithm and identified 231 genes validated as having a miR-21 target site. We then searched MEDLINE to select among these genes those relevant to NASH, that is, those with a known downregulated liver expression in NASH. We used each one of the 231 gene names together with the terms “steatohepatitis OR NASH OR fatty liver”. We identified phosphatase and tensin homologue deleted on chromosome 10 (PTEN), caspase 9 and PPAR α . Out of these three potential targets, we screened those that would fit with our observation of increased miR-21 expression in biliary and inflammatory cells (figures 1 and 4). Only PPAR α fulfilled this criteria.^{22–25} In both mouse models, liver PPAR α gene and protein expressions were decreased in NASH and were restored by pharmacological inhibition or genetic suppression of miR-21 (figure 7).

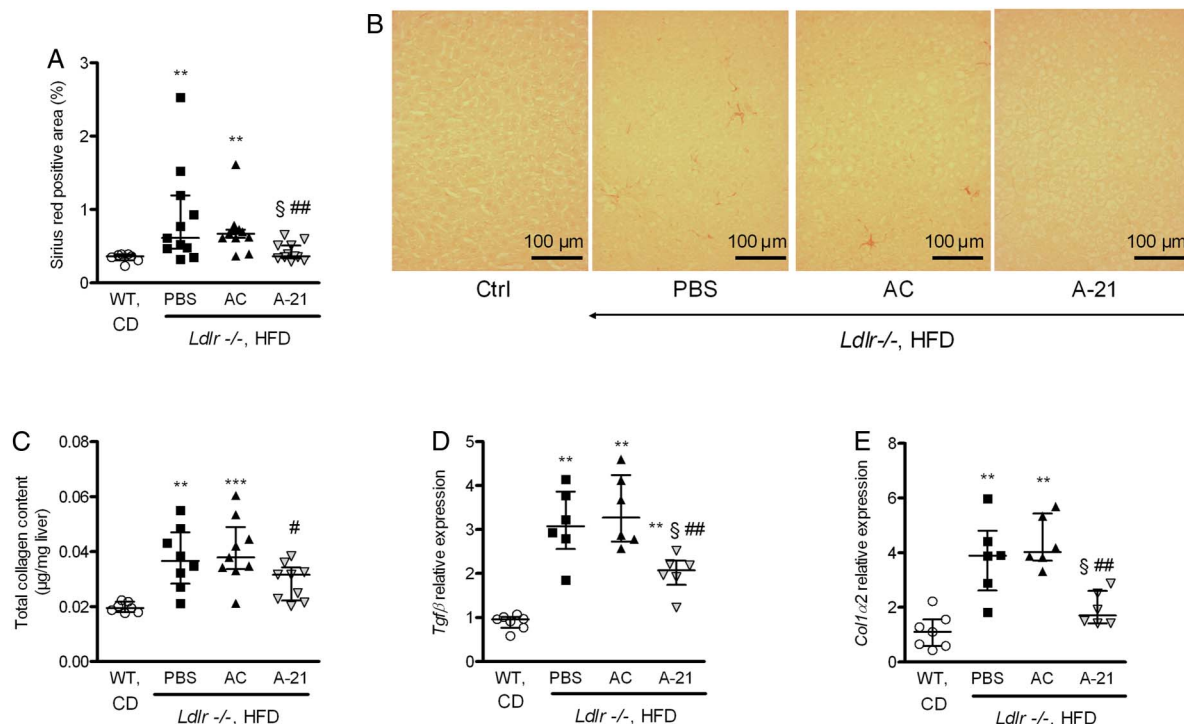
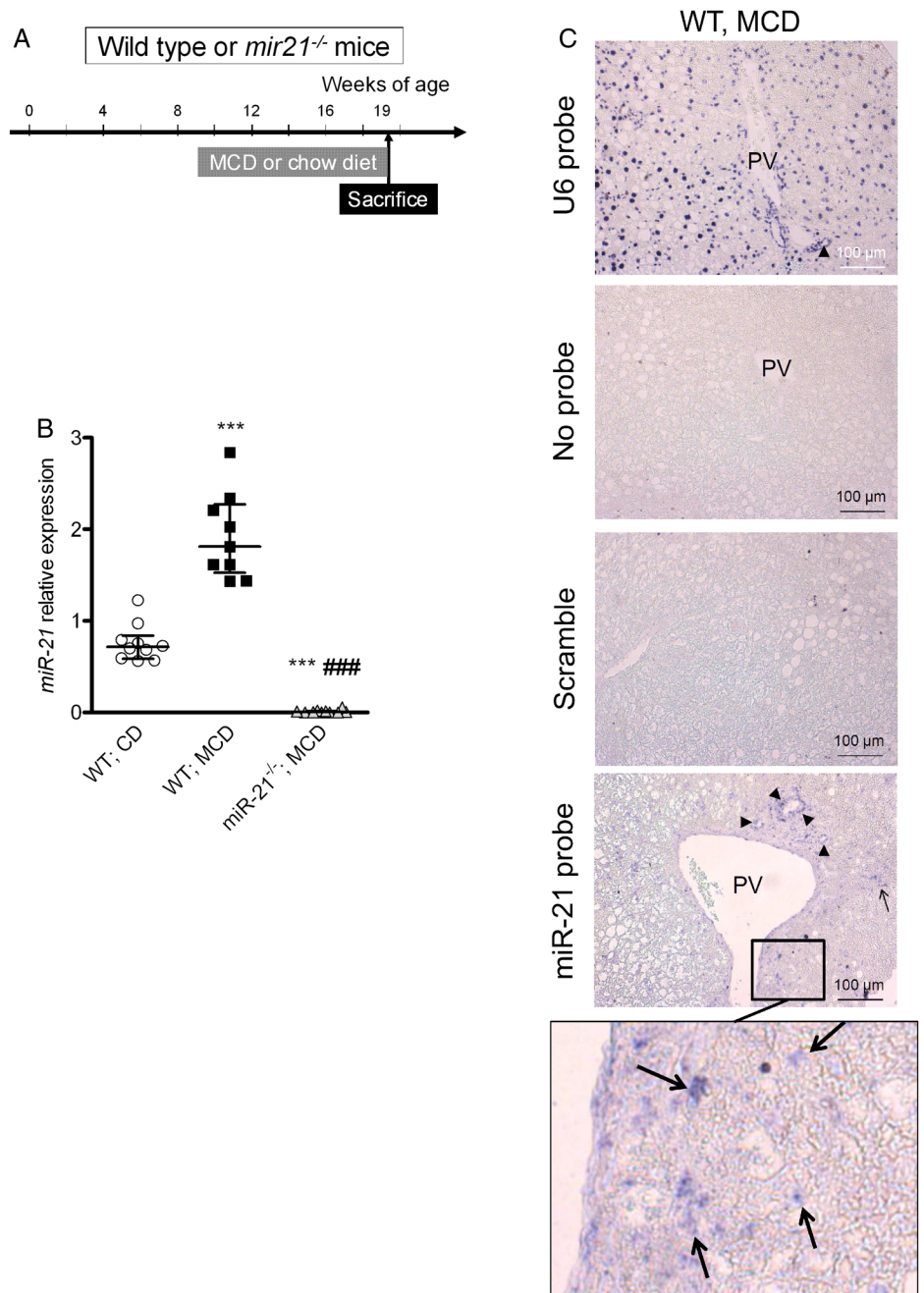


Figure 3 Antagomir-21 (A-21) reduced liver fibrosis and expression of fibrogenic-related genes in the liver of low-density lipoprotein receptor-deficient (*Ldlr*^{-/-}) mice fed a high-fat diet (HFD). A-21 reduced liver collagen staining (Sirius red; A, quantification; B representative images, original magnification $\times 200$), collagen deposition (C), as well as liver normalised *Tgfb* (D), and *collagen-1 α 2* (E) mRNA expression. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$ versus wild-type (WT) mice fed a chow diet (CD); $\$p < 0.05$ versus *Ldlr*^{-/-} mice fed an HFD treated with phosphate buffer saline (PBS); # $p < 0.05$, ## $p < 0.01$ versus *Ldlr*^{-/-} mice fed a HFD treated with antagomir control (AC). Data are given as median (horizontal bar) and IQR (error bar). Samples were randomly selected in each group of mice.

Figure 4 MicroRNA-21 (miR-21) was overexpressed in the liver of wild-type (WT) mice fed a methionine-choline-deficient (MCD) diet. (A) Mouse study design. (B) miR-21 expression in the liver of WT mice fed a chow diet (CD) or an MCD diet as well as in *miR-21*^{-/-} mice fed an MCD diet. ****p*<0.001 versus WT mice fed a chow diet (CD); ###*p*<0.001 versus WT mice fed an MCD diet. Data are given as median (horizontal bar) and IQR (error bar). (C) In situ hybridisation. Arrows point to liver inflammatory cells and arrow heads point biliary cells. Original magnification $\times 200$. PV, portal vein.



To firmly establish that the effect of miR-21 implicates PPAR α , WT and *Ppara*^{-/-} mice fed an MCD diet for 5 weeks were treated with antagomiR-21 or antagomir control (figure 8A). Liver *miR-21* expression was increased by MCD diet both in WT and in *Ppara*^{-/-} mice and was dramatically decreased upon antagomiR-21 treatment (figure 8B). In accordance with results obtained in *Ldlr*^{-/-} mice, *miR-21* was overexpressed in CD45⁺ cells, particularly in T cells, and was also detected in biliary cells (figure 8C, D).

When compared with WT mice fed an MCD diet, *Ppara*^{-/-} fed an MCD diet had lower serum HDL cholesterol and glucose levels, more steatosis and lower liver expression of the β -oxidation genes *Cpt1* and *Acox1*, but no significant change in liver cell injury, hepatic inflammation or liver fibrosis (see online supplementary table S4, figure S4 and figures 8 and 9). Similarly to what we observed in *Ldlr*^{-/-} mice, antagomiR-21 also improved liver cell injury, hepatic inflammation and liver fibrosis in WT mice fed an MCD diet. Importantly, this

effect was lost in *Ppara*^{-/-} mice fed an MCD diet (figures 8 and 9).

miR-21 is overexpressed in the liver of patients with NASH primarily in inflammatory cells

Online supplementary table S5 summarises the features of the 25 patients included in this study. As shown in figure 10A, miR-21 was overexpressed in the liver of patients with NASH compared with patients with no or mild abnormalities at histological examination or, interestingly, with bland steatosis. Liver miR-21 expression correlated better with liver activity (ballooning, $r=0.713$, $p<0.001$; lobular inflammation, $r=0.735$, $p<0.001$) than with steatosis ($r=0.539$, $p=0.005$) or fibrosis ($r=0.461$, $p=0.021$). Using fluorescent in situ hybridisation, we observed that miR-21 was primarily expressed in human biliary (CK19⁺) and inflammatory (CD3⁺) cells in the liver from patients with NASH, similarly to what was observed in *Ldlr*^{-/-} mice fed an HFD (figure 10B, C).

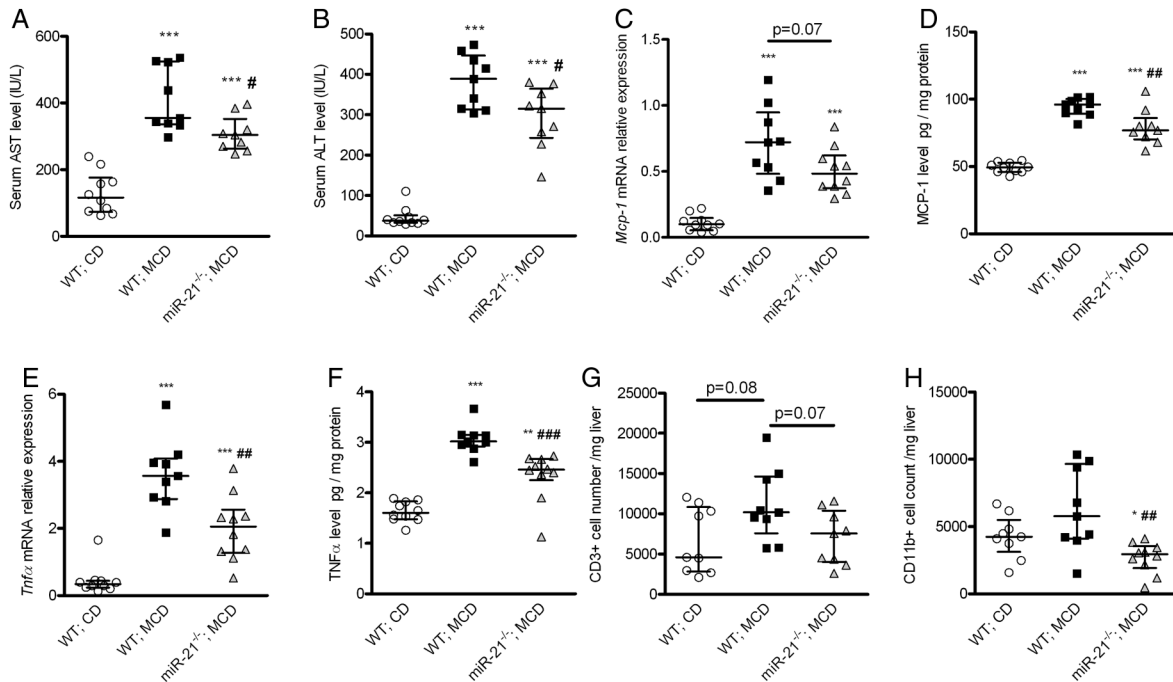


Figure 5 *MIR-21*^{-/-} mice fed a methionine-choline-deficient (MCD) diet had reduced liver injury and liver inflammation. When compared with wild-type (WT) mice, *miR-21*^{-/-} mice had reduced serum aspartate aminotransferase (AST) (A) and alanine aminotransferase (ALT) (B) levels, liver monocyte chemoattractant protein-1 (MCP-1) (C, D) and tumour necrosis factor (TNF)- α (E, F) gene and protein expressions, as well as lower CD3⁺ (G, not significant) and CD11b⁺ (H) cell infiltrates. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$ versus WT mice fed a chow diet (CD). # $p < 0.05$, ## $p < 0.01$, ### $p < 0.001$ versus WT mice fed an MCD diet. Data are given as median (horizontal bar) and IQR (error bar).

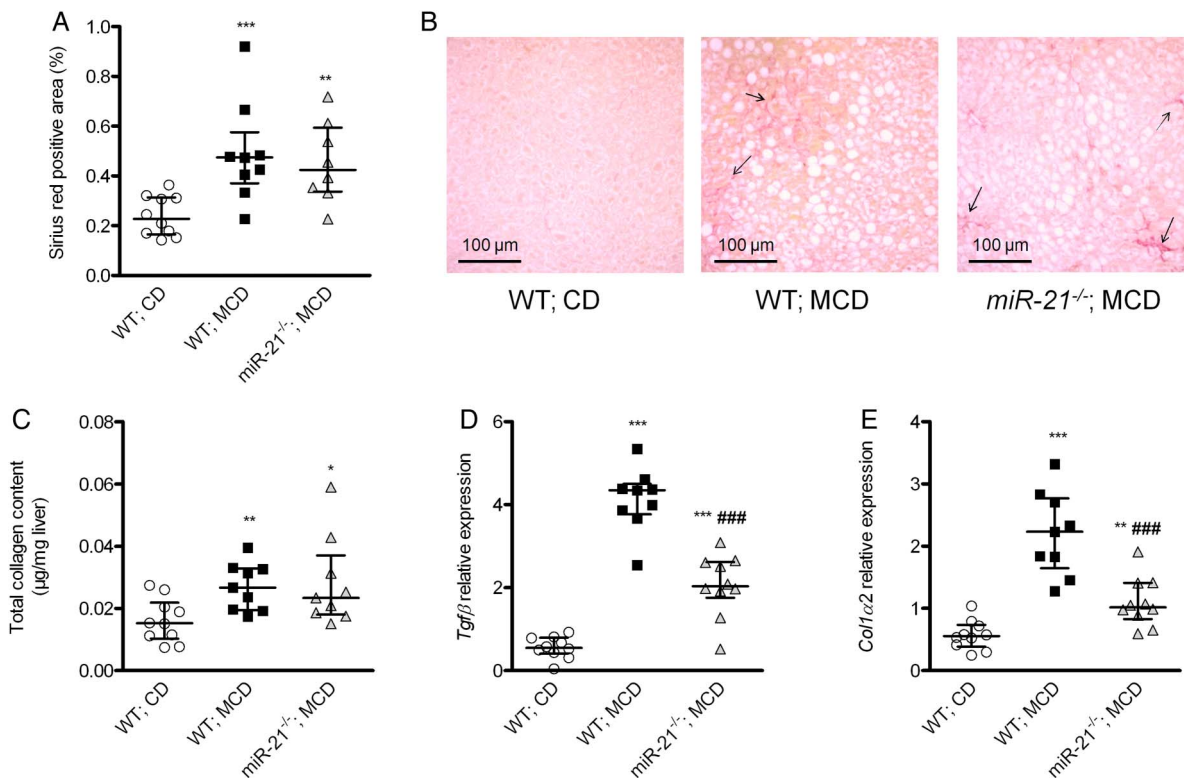


Figure 6 *MIR-21*^{-/-} mice fed a methionine-choline-deficient (MCD) diet had reduced expression of fibrogenic-related genes. When compared with wild-type (WT) mice, *miR-21*^{-/-} mice had similar liver collagen staining (Sirius red; A, quantification; B representative images, original magnification $\times 200$) and collagen deposition (C), but reduced liver normalised *Tgfb* (D) and *collagen-1 α 2* (E) mRNA expression. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$ versus WT mice fed a chow diet (CD). ## $p < 0.01$ versus WT mice fed an MCD diet. Data are given as median (horizontal bar) and IQR (error bar).

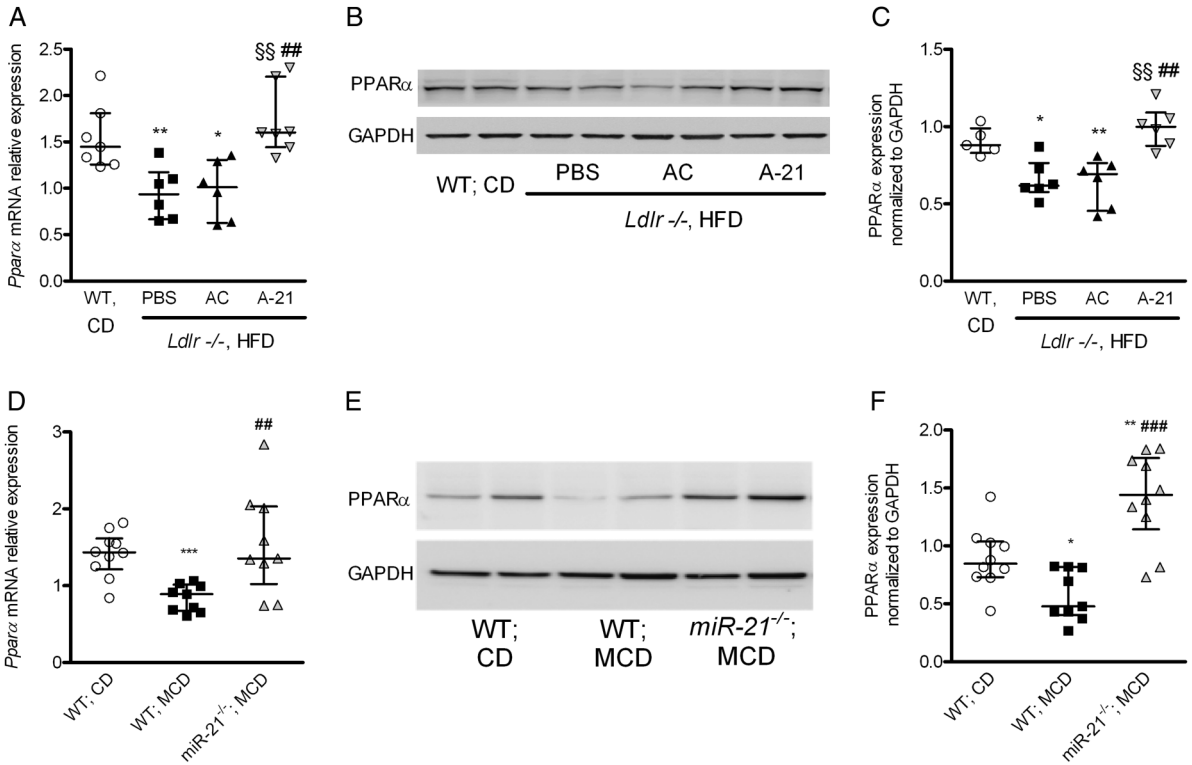


Figure 7 MicroRNA-21 (miR-21) inhibition or suppression restored liver peroxisome proliferation-activator receptor α (PPAR α) expression. (A–C) Antagomir-21 (A-21) restored liver PPAR α gene (A) and protein (B and C) expression. * $p < 0.05$, ** $p < 0.01$ versus wild-type (WT) mice fed a chow diet (CD); $\$p < 0.05$; $\$\$p < 0.01$ versus low-density lipoprotein receptor-deficient (*Ldlr*^{-/-}) mice fed a high-fat diet (HFD) treated with phosphate buffer saline (PBS); $\#\#p < 0.01$ versus *Ldlr*^{-/-} mice fed an HFD treated with antagomir control (AC). (D–F) *Mir-21*^{-/-} mice fed a methionine-choline-deficient (MCD) diet had restored liver PPAR α gene (D) and protein (E and F) expression. * $p < 0.05$, ** $p < 0.01$, versus WT mice fed a chow diet (CD). $\#\#\#p < 0.001$ versus WT mice fed an MCD diet. Data are given as median (horizontal bar) and IQR (error bar). Liver samples randomly selected in each group of mice.

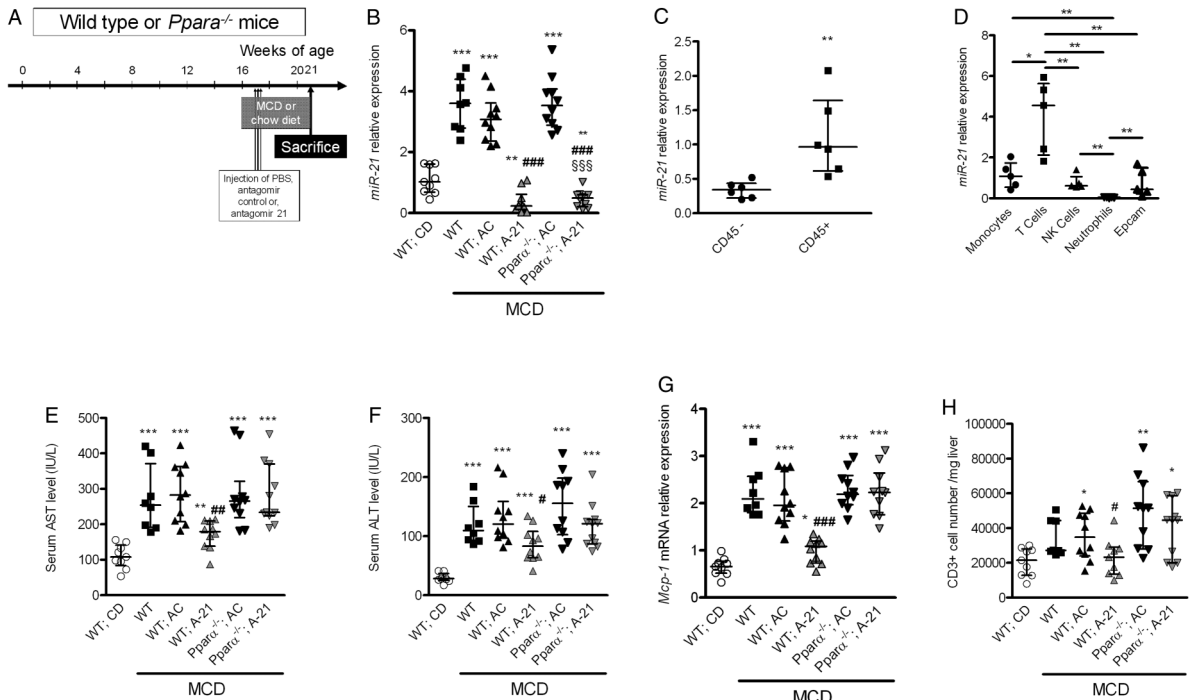


Figure 8 Antagomir-21 (A-21) did not improve liver cell injury or liver inflammation in *Ppara*^{-/-} mice fed a methionine-choline-deficient (MCD) diet. (A) Mouse study design. (B) Liver microRNA-21 (miR-21) expression. (C) miR-21 expression in sorted liver inflammatory (CD45⁺) and non-inflammatory (CD45⁻) cells from wild-type (WT) mice fed an MCD diet for 5 weeks. (D) miR-21 expression in subpopulations of liver inflammatory (CD45⁺) cells and in biliary cells (Epcam⁺) from WT mice fed an MCD diet for 5 weeks. A-21 did neither reduce serum aspartate aminotransferase (AST) (E) or alanine aminotransferase (ALT) (F) levels, nor liver *Mcp1* gene expression (G) or CD3⁺ (H) cell infiltrates in *Ppara*^{-/-} mice. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$ versus WT mice fed a chow diet (CD). $\#p < 0.05$, $\#\#p < 0.01$, $\#\#\#p < 0.001$ versus WT mice fed an MCD diet and treated with antagomir control (AC). Data are given as median (horizontal bar) and IQR (error bar).

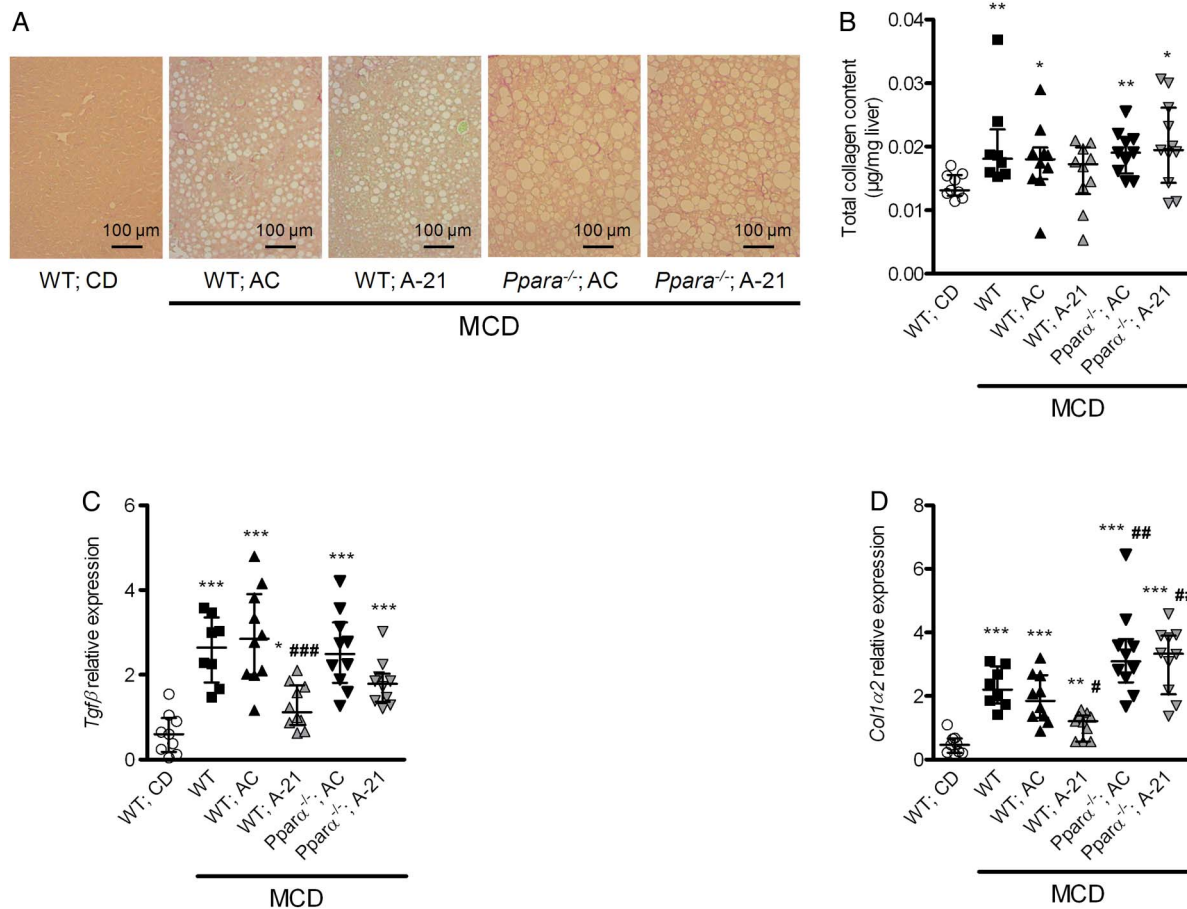


Figure 9 Antagomir-21 (A-21) did not improve liver fibrosis in *Ppara*^{-/-} mice fed a methionine-choline-deficient (MCD) diet. A-21 did not reduce liver collagen staining (A; Sirius red, representative images, original magnification $\times 200$), collagen deposition (B) or liver normalised *Tgfb* (C) and *collagen-1 α 2* (D) mRNA expression in *Ppara*^{-/-} mice fed an MCD diet. * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$ versus wild type (WT) mice fed a chow diet (CD). # $p < 0.05$, ### $p < 0.001$ versus WT mice fed an MCD diet and treated with antagomir-21. Data are given as median (horizontal bar) and IQR (error bar).

DISCUSSION

This study demonstrates that miR-21 inhibition or suppression decreases liver injury, inflammation and fibrosis, in different murine models of NASH, by restoring PPAR α expression. These findings support the hypothesis that miR-21 is involved in the pathogenesis of NASH. We also show that miR-21 is primarily expressed in liver inflammatory and biliary cells in murine models and in patients with NASH, thereby enhancing the clinical relevance of our findings. Taken all together, these findings point to the potential of antagomir-21 as a possible treatment for NASH.

A first major finding in this study was that miR-21 inhibition or suppression reduced liver cell injury, liver inflammation and liver fibrogenesis in complementary models of NASH. The effect of the pharmacological inhibition of miR-21 appeared stronger than the genetic inhibition. Indeed, *Ldlr*^{-/-} mice fed an HFD and treated with antagomir-21 had normal markers of liver injury, liver inflammation and liver fibrosis, while these parameters were reduced but not normalised in *miR-21*^{-/-} mice fed an MCD diet. There was even no change in collagen deposition in *miR-21*^{-/-} mice despite a reduction in liver expression of fibrogenic-related genes. We can speculate that genetic deletion of miR-21 occurring in early life leads to the development of compensatory mechanisms that could account for that observation.

A second major finding in this study was that miR-21 expression was elevated in the liver in patients with NASH compared

with those with bland steatosis or with no or mild abnormalities at liver histological examination. The difference between patients with NASH and normal liver had been previously suggested by microRNA microarray assay, but not demonstrated by real-time quantitative PCR, and patients with bland steatosis had not been analysed.⁷ This result contrasts with the normal circulating miR-21 levels reported in patients with NASH.²⁶ Importantly, we showed here in mice and in patients that this augmented liver miR-21 expression resides predominantly in inflammatory and biliary cells, and not in hepatocytes. This result is reinforced by the very significant correlation between liver miR-21 expression and activity score in the liver of patients. Liver infiltration with inflammatory cells is a key feature of NASH since this is one of the minimal criteria for the histological diagnosis of adult NASH.^{17, 27} MiR-21 has been reported to be upregulated in many inflamed states, including lungs in lipopolysaccharide-treated mice,²⁸ allergic airway inflammation,²⁹ osteoarthritis,³⁰ psoriasis and atopic eczema,³¹ disease-active UC tissue,³² *Helicobacter pylori*-associated gastric cancer,³³ cardiac muscle injury³⁴ and cardiac hypertrophy.³⁵ Interestingly, miR-21 expression is upregulated by lipopolysaccharide in macrophages and human peripheral blood mononuclear cells.³⁶ This feature might explain the upregulation of miR-21 in NASH since plasma lipopolysaccharide levels are higher in patients and mice with NAFLD than controls.^{37, 38}

Biliary cells also seem to play a key role in NASH. In patients with NASH, there is an increased ductular reaction compared

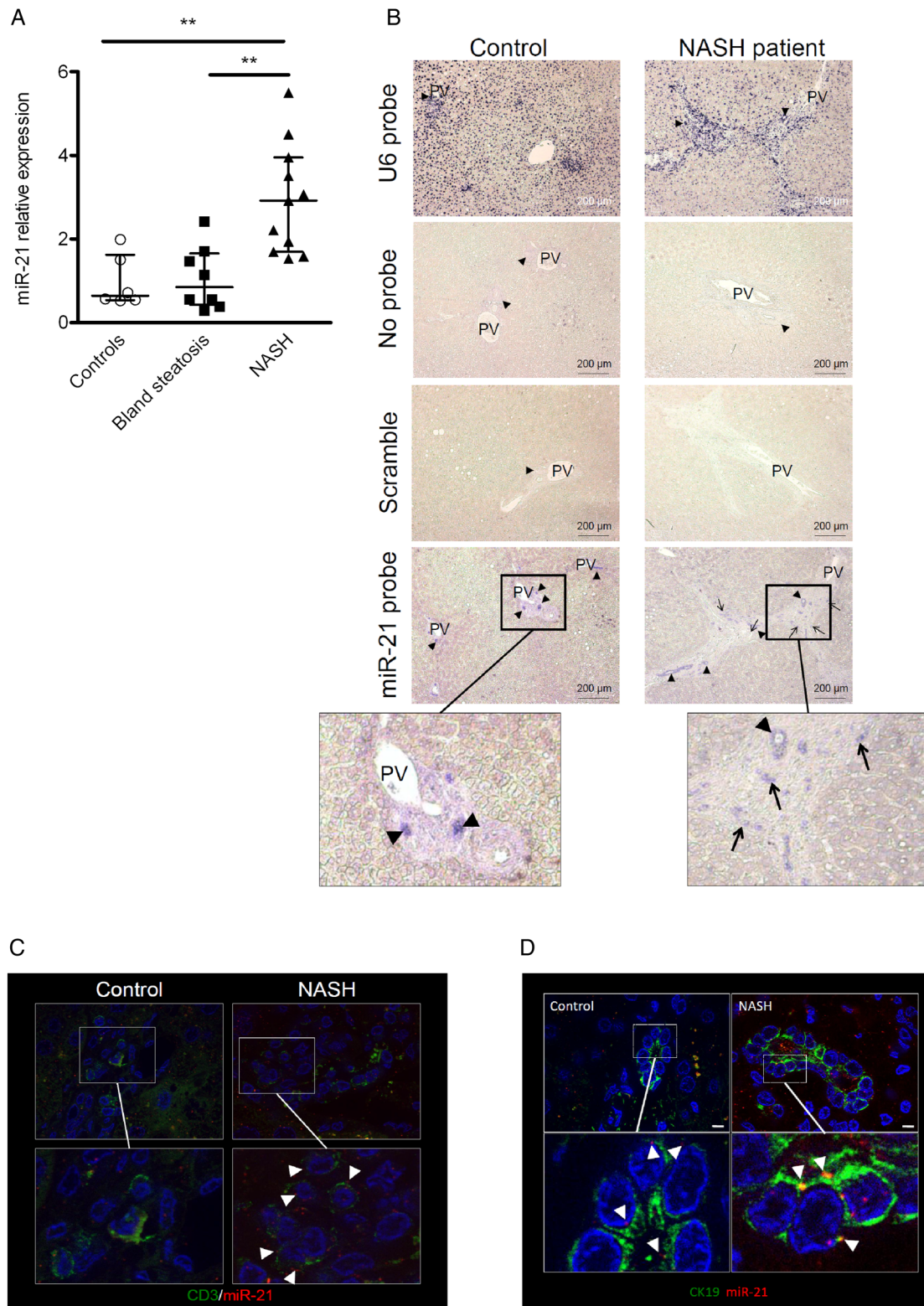


Figure 10 MicroRNA-21 (miR-21) was overexpressed in the liver of patients with non-alcoholic steatohepatitis (NASH), primarily in inflammatory and biliary cells. (A) miR-21 expression normalised to U6 snRNA in the liver of patients with no or mild abnormalities at histological examination (control) or with bland steatosis or with NASH. * $p < 0.05$, ** $p < 0.01$. (B) In situ hybridisation. miR-21 was detected in inflammatory (arrow) and biliary cells (arrow head). U6 probe is the positive control and no probe the negative control. Original magnification $\times 200$. (C and D) Fluorescence in situ hybridisation (FISH). Combined FISH was performed for miR-21 and CD3 (lymphocyte marker) or CK19 (biliary cells marker) in liver sections from controls or patients with NASH. miR-21 expression (red) was detected in CD3-positive cells (C, green) from patients with NASH and in CK19-positive cells from patients with NASH and from controls (D, green). Original magnification $\times 630$. Representative of three experiments. PV, portal vein.

with simple steatosis, which correlates with the stage of fibrosis, with the grade of NASH activity, as well as with the grade of portal inflammation.³⁹ MiR-21 is expressed in cholangiocytes in

a basal manner and overexpressed following exposure to lipopolysaccharide.⁴⁰ We can speculate that endotoxemia associated with NASH could increase miR-21 expression in biliary cells.

To explore the molecular mechanism involving miR-21 in NASH, we first identified 231 genes validated as having an miR-21 target site. When selecting those relevant to NASH, that is, those for which liver expression was downregulated in NASH, we identified phosphatase and tensin homologue deleted on chromosome 10 (PTEN), caspase 9 and PPAR α . However, in fatty liver, PTEN expression is decreased specifically in fatty hepatocytes.⁴¹ This did not fit with our findings, showing that miR-21 is overexpressed specifically in biliary and inflammatory cells. Similarly, caspase 9 has a reduced liver expression only in ballooned hepatocytes in NASH.⁴² Therefore, it was unlikely that PTEN and caspase 9 were responsible for miR-21 effect in NASH. PPAR α is a well-known target of miR-21.^{43–44} Previous studies reported that PPAR α expression is decreased in the liver in patients and rats with NASH, and inversely correlates with NASH severity.^{22–23} As PPAR α is expressed not only by hepatocytes, but also by biliary cells and inflammatory cells, this prompted us to test the hypothesis that miR-21 contributes NASH by targeting PPAR α .^{24–25–45} We first observed in the murine models of NASH that changes in liver PPAR α mRNA and protein expression mirrored those of miR-21. Pharmacological inhibition and genetic suppression of miR-21 restored liver PPAR α expression in the murine NASH models. Second, although not specific, changes in liver PPAR α levels were accompanied by opposite changes in the downstream molecules of the PPAR α pathway, including TNF- α and MCP-1.^{44–46} Third, PPAR α is known to be expressed in inflammatory and biliary cells, that is, the cells where we observed an miR-21 upregulation.^{24–25} Interestingly, a recent report has shown that PPAR α inhibits hepatic inflammatory responses leading to liver injury and progression to fibrosis by a mechanism independent of its effect on lipid accumulation in hepatocytes.⁴⁵ This latter study suggests that steatosis is related to decreased expression of PPAR α in hepatocytes and that inflammation and fibrosis are related to decreased expression of PPAR α in inflammatory cells. Fourth and most importantly, we observed that the beneficial effect of antagomir-21 on liver cell injury, hepatic inflammation and liver fibrosis was lost in *Ppara*^{-/-} mice fed an MCD diet. Altogether, this suggests that miR-21, expressed mainly in the liver in inflammatory cells, regulates liver inflammation and fibrosis via PPAR α inhibition in these cells, but has no effect on steatosis because miR-21 is weakly expressed in hepatocytes.

MiR-21 is considered a druggable target microRNA.⁵ The present results suggest that inhibition of the miR-21 pathway might be an interesting new avenue for treating patients with NASH in the future, similarly to miR-122 inhibition in patients with chronic hepatitis C virus infection.⁴⁷ Such a strategy may overcome the limitations of PPAR α pharmacological agonists in NASH. Indeed, even though the novel dual PPAR α/δ agonist GFT505 seems promising, trials with fibrates, that is, PPAR α activators, have yielded mixed results in humans so that they are currently not recommended in patients.^{1–48–49} Yet, while hepatitis C virus can be eradicated, one would speculate that NASH would recur after treatment interruption as long as the risk factors for NASH persist. Therefore, long-term miR-21 antagonism treatment would be necessary with inherent risks of long-term side effects. Careful study design and safety monitoring would be critical.

In conclusion, we demonstrated here using complementary mouse models of NASH that inhibition and suppression of miR-21 strongly decreased liver injury, liver inflammation and fibrogenesis, through liver PPAR α normalisation. We also showed that miR-21 was overexpressed in the liver of patients with NASH

primarily in inflammatory and in biliary cells. Antagomir-21 might be a possible therapeutic strategy for NASH.

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Correction notice This article has been corrected since it published Online First. The Open Access licence has been added.

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Contributors XL, P-ER, and AT obtained funding, conceived and designed the study, and interpreted the data. XL and P-ER drafted the manuscript. P-ER collected the patients clinical and laboratory data. CMB and AT contributed to the writing of the manuscript. XL, A-CV, CH, NC, CLG, CD, SO, JS, MR and J-LP carried out experiments and analysed the data. MER provided *miR-21*^{-/-} mice. CPB carried out hepatic ketonic bodies measurement and discussed data and manuscript. EB and BS provided *Ppara*^{-/-} mice and discussed the data and the manuscript. PM, FD, PB and VP provided liver samples and contributed to the analysis of the data. All authors critically revised the manuscript and approved its final version.

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Competing interests None declared.

Patient consent Obtained.

Ethics approval The study protocol conformed to the ethical guidelines of the 1975 Declaration of Helsinki.

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Data sharing statement We agree to share all primary data and protocols employed in this study.

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REFERENCES

- 1 Chalasan N, Younossi Z, Lavine JE, *et al*. The diagnosis and management of non-alcoholic fatty liver disease: Practice Guideline by the American Association for the Study of Liver Diseases, American College of Gastroenterology, and the American Gastroenterological Association. *Hepatology* 2012;55:2005–23.
- 2 Ratziu V, Bellentani S, Cortez-Pinto H, *et al*. A position statement on NAFLD/NASH based on the EASL 2009 special conference. *J Hepatol* 2010;53:372–84.
- 3 Weston SR, Leyden W, Murphy R, *et al*. Racial and ethnic distribution of nonalcoholic fatty liver in persons with newly diagnosed chronic liver disease. *Hepatology* 2005;41:372–9.
- 4 Bartel DP. MicroRNAs: target recognition and regulatory functions. *Cell* 2009;136:215–33.

- 5 Wang XW, Heegaard NH, Orum H. MicroRNAs in liver disease. *Gastroenterology* 2012;142:1431–43.
- 6 Szabo G, Sarnow P, Bala S. MicroRNA silencing and the development of novel therapies for liver disease. *J Hepatol* 2012;57:462–6.
- 7 Cheung O, Puri P, Eicken C, et al. Nonalcoholic steatohepatitis is associated with altered hepatic MicroRNA expression. *Hepatology* 2008;48:1810–20.
- 8 Dattaray D, Pourhoseini S, Das S, et al. Micro-RNA 21 inhibition of SMAD7 enhances fibrogenesis via leptin-mediated NADPH oxidase in experimental and human nonalcoholic steatohepatitis. *Am J Physiol Gastrointest Liver Physiol* 2015;308:G298–312.
- 9 Wang B, Majumder S, Nuovo G, et al. Role of microRNA-155 at early stages of hepatocarcinogenesis induced by choline-deficient and amino acid-defined diet in C57BL/6 mice. *Hepatology* 2009;50:1152–61.
- 10 da Costa Martins PA, Salic K, Gladka MM, et al. MicroRNA-199b targets the nuclear kinase Dyrk1a in an auto-amplification loop promoting calcineurin/NFAT signalling. *Nat Cell Biol* 2010;12:1220–7.
- 11 Loyer X, Potteaux S, Vion AC, et al. Inhibition of microRNA-92a prevents endothelial dysfunction and atherosclerosis in mice. *Circ Res* 2014;114:434–43.
- 12 Krutzfeldt J, Rajewsky N, Braich R, et al. Silencing of microRNAs in vivo with 'antagomirs'. *Nature* 2005;438:685–9.
- 13 Lu TX, Hartner J, Lim EJ, et al. MicroRNA-21 limits in vivo immune response-mediated activation of the IL-12/IFN-gamma pathway, Th1 polarization, and the severity of delayed-type hypersensitivity. *J Immunol* 2011;187:3362–73.
- 14 Schaefer A, Jung M, Miller K, et al. Suitable reference genes for relative quantification of miRNA expression in prostate cancer. *Exp Mol Med* 2010;42:749–58.
- 15 Bonauer A, Carmona G, Iwasaki M, et al. MicroRNA-92a controls angiogenesis and functional recovery of ischemic tissues in mice. *Science* 2009;324:1710–13.
- 16 Thum T, Gross C, Fiedler J, et al. MicroRNA-21 contributes to myocardial disease by stimulating MAP kinase signalling in fibroblasts. *Nature* 2008;456:980–4.
- 17 Bedossa P, Poitou C, Veyrie N, et al. Histopathological algorithm and scoring system for evaluation of liver lesions in morbidly obese patients. *Hepatology* 2012;56:1751–9.
- 18 Kleiner DE, Brunt EM, Van Natta M, et al. Design and validation of a histological scoring system for nonalcoholic fatty liver disease. *Hepatology* 2005;41:1313–21.
- 19 Bustin SA, Benes V, Garson JA, et al. The MIQE guidelines: minimum information for publication of quantitative real-time PCR experiments. *Clin Chem* 2009;55:611–22.
- 20 Vandesompele J, De Preter K, Pattyn F, et al. Accurate normalization of real-time quantitative RT-PCR data by geometric averaging of multiple internal control genes. *Genome Biol* 2002;3:Research0034.
- 21 Williamson DH, Bates MW, Krebs HA. Activity and intracellular distribution of enzymes of ketone-body metabolism in rat liver. *Biochem J* 1968;108:353–61.
- 22 Francque S, Verrijken A, Caron S, et al. PPARalpha gene expression correlates with severity and histological treatment response in patients with non-alcoholic steatohepatitis. *J Hepatol* 2015;63:164–73.
- 23 Yeon JE, Choi KM, Baik SH, et al. Reduced expression of peroxisome proliferator-activated receptor-alpha may have an important role in the development of non-alcoholic fatty liver disease. *J Gastroenterol Hepatol* 2004;19:799–804.
- 24 Lefebvre P, Chinetti G, Fruchart JC, et al. Sorting out the roles of PPAR alpha in energy metabolism and vascular homeostasis. *J Clin Invest* 2006;116:571–80.
- 25 Firrincieli D, Zuniga S, Poupon R, et al. Role of nuclear receptors in the biliary epithelium. *Dig Dis* 2011;29:52–7.
- 26 Cermelli S, Ruggieri A, Marrero JA, et al. Circulating microRNAs in patients with chronic hepatitis C and non-alcoholic fatty liver disease. *PLoS ONE* 2011;6:e23937.
- 27 Brunt EM, Tiniakos DG. Histopathology of nonalcoholic fatty liver disease. *World J Gastroenterol* 2010;16:5286–96.
- 28 Moschos SA, Williams AE, Perry MM, et al. Expression profiling in vivo demonstrates rapid changes in lung microRNA levels following lipopolysaccharide-induced inflammation but not in the anti-inflammatory action of glucocorticoids. *BMC Genomics* 2007;8:240.
- 29 Lu TX, Munitz A, Rothenberg ME. MicroRNA-21 is up-regulated in allergic airway inflammation and regulates IL-12p35 expression. *J Immunol* 2009;182:4994–5002.
- 30 Iliopoulos D, Malizos KN, Oikonomou P, et al. Integrative microRNA and proteomic approaches identify novel osteoarthritis genes and their collaborative metabolic and inflammatory networks. *PLoS ONE* 2008;3:e3740.
- 31 Sonkoly E, Wei T, Janson PC, et al. MicroRNAs: novel regulators involved in the pathogenesis of psoriasis? *PLoS ONE* 2007;2:e610.
- 32 Wu F, Zikusoka M, Trindade A, et al. MicroRNAs are differentially expressed in ulcerative colitis and alter expression of macrophage inflammatory peptide-2 alpha. *Gastroenterology* 2008;135:1624–35 e24.
- 33 Zhang Z, Li Z, Gao C, et al. miR-21 plays a pivotal role in gastric cancer pathogenesis and progression. *Lab Invest* 2008;88:1358–66.
- 34 Ji R, Cheng Y, Yue J, et al. MicroRNA expression signature and antisense-mediated depletion reveal an essential role of MicroRNA in vascular neointimal lesion formation. *Circ Res* 2007;100:1579–88.
- 35 Cheng Y, Ji R, Yue J, et al. MicroRNAs are aberrantly expressed in hypertrophic heart: do they play a role in cardiac hypertrophy? *Am J Pathol* 2007;170:1831–40.
- 36 Sheedy FJ, Palsson-McDermott E, Hennessy EJ, et al. Negative regulation of TLR4 via targeting of the proinflammatory tumor suppressor PDCD4 by the microRNA miR-21. *Nat Immunol* 2010;11:141–7.
- 37 Thuy S, Ladurner R, Volynets V, et al. Nonalcoholic fatty liver disease in humans is associated with increased plasma endotoxin and plasminogen activator inhibitor 1 concentrations and with fructose intake. *J Nutr* 2008;138:1452–5.
- 38 Mouzaki M, Comelli EM, Arendt BM, et al. Intestinal microbiota in patients with nonalcoholic fatty liver disease. *Hepatology* 2013;58:120–7.
- 39 Richardson MM, Jonsson JR, Powell EE, et al. Progressive fibrosis in nonalcoholic steatohepatitis: association with altered regeneration and a ductular reaction. *Gastroenterology* 2007;133:80–90.
- 40 Zhou R, Hu G, Liu J, et al. NF-kappaB p65-dependent transactivation of miRNA genes following *Cryptosporidium parvum* infection stimulates epithelial cell immune responses. *PLoS Pathog* 2009;5:e1000681.
- 41 Vinciguerra M, Sgroi A, Veyrat-Durebex C, et al. Unsaturated fatty acids inhibit the expression of tumor suppressor phosphatase and tensin homolog (PTEN) via microRNA-21 up-regulation in hepatocytes. *Hepatology* 2009;49:1176–84.
- 42 Kakisaka K, Cazanave SC, Werneburg NW, et al. A hedgehog survival pathway in 'undead' lipotoxic hepatocytes. *J Hepatol* 2012;57:844–51.
- 43 Sarkar J, Gou D, Turaka P, et al. MicroRNA-21 plays a role in hypoxia-mediated pulmonary artery smooth muscle cell proliferation and migration. *Am J Physiol Lung Cell Mol Physiol* 2010;299:L861–71.
- 44 Zhou J, Wang KC, Wu W, et al. MicroRNA-21 targets peroxisome proliferator-activated receptor-alpha in an autoregulatory loop to modulate flow-induced endothelial inflammation. *Proc Natl Acad Sci USA* 2011;108:10355–60.
- 45 Pawlak M, Bauge E, Bourguet W, et al. The transrepressive activity of peroxisome proliferator-activated receptor alpha is necessary and sufficient to prevent liver fibrosis in mice. *Hepatology* 2014;60:1593–606.
- 46 Marx N, Kehrl B, Kohlhammer K, et al. PPAR activators as antiinflammatory mediators in human T lymphocytes: implications for atherosclerosis and transplantation-associated arteriosclerosis. *Circ Res* 2002;90:703–10.
- 47 Janssen HL, Reesink HW, Lawitz EJ, et al. Treatment of HCV infection by targeting microRNA. *N Engl J Med* 2013;368:1685–94.
- 48 Kallwitz ER, McLachlan A, Cotler SJ. Role of peroxisome proliferator-activated receptors in the pathogenesis and treatment of nonalcoholic fatty liver disease. *World J Gastroenterol* 2008;14:22–8.
- 49 Staels B, Rubenstunck A, Noel B, et al. Hepato-protective effects of the dual PPARalpha/delta agonist GFT505 in rodent models of NAFLD/NASH. *Hepatology* 2013;58:1941–52.