Alternative activation of macrophages by prostacyclin synthase ameliorates alcohol induced liver injury

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Abstract
Alcoholic liver disease (ALD) is a major cause of chronic liver disease worldwide. Macrophages exhibit different functional states and are classified as classically activated (M1) and alternatively activated (M2) macrophages. However, the mechanisms that govern M1/M2 polarization in chronic ALD remain to be elucidated. Prostacyclin (PGI2) synthase (PTGIS) is an enzyme of the prostaglandin pathway which catalyzes the conversion of Prostaglandin H2 (PGH2) to PGI2. PTGIS has anti-inflammatory properties. However, the function of PTGIS in ALD has not yet been determined. In this study, we demonstrated that PTGIS was downregulated in ALD and forced PTGIS expression in vivo using recombinant adeno-associated viral vector-packed PTGIS overexpression plasmid, which alleviated the inflammatory response and suppressed the macrophage M1 phenotype in mice. Loss- and gain-of function-experiments demonstrated that forced PTGIS expression inhibited the macrophage switch to the M1 phenotype and promoted M2 polarization. Furthermore, we identified the genes regulated by PTGIS through RNA-sequencing (RNA-seq) analysis. Gene ontology and KEGG pathway analyses showed that PTGIS regulates many genes involved in the immune response and is enriched in the Janus kinase/signal transducers and activators of transcription (JAK/STAT) signal transduction pathway, which plays an important role in regulating macrophage polarization. The proteins interacting with JAKs were predicted using the STRING database. The overlap between the RNA-seq and the STRING database was interleukin-6; this indicated that it was involved in macrophage polarization regulated by JAK/STAT signaling. We further explored the microRNAs that could regulate the expression of PTGIS through TargetScan. The results of luciferase assay illustrated that the expression of PTGIS was regulated by miR-140-3p.1. These results imply that PTGIS plays a pivotal role in ALD, partly by influencing macrophage polarization.

Introduction
Alcohol consumption is a leading risk factor for global disease burden and causes substantial health loss [1]. Alcoholic liver disease (ALD) is the leading cause of liver-related disorders and contributes to 25% of alcohol-related deaths worldwide [2]. ALD is characterized by progressive liver steatosis, fibrosis, and cirrhosis. Consumption of alcohol increases gut permeability, which leads to the translocation of both microbes and microbial products, including lipopolysaccharide (LPS), β-glucan, and other pathogen-associated molecular patterns, into the portal circulation, activating innate immune responses in the liver [3, 4]. Macrophages are highly heterogeneous, plastic populations that undergo pleiotropic coordinated responses to tissue damage through distinct programs of activation, known as classical activation (M1) or alternative activation (M2) [5, 6]. M1 activation is mainly induced by Th1 cytokines, such as interferon-gamma (IFNγ), or bacterial molecular patterns, including endotoxin/LPS.

The macrophage switch to an M1 phenotype plays a key role in releasing pro-inflammatory enzymes and cytokines, such as inducible nitric oxide synthase (iNOS), interleukin...
orectal cancer [28], and prostaglandin G/H synthase 2 (PGIS) plays an anti-inflammatory role in colonic inflammation and is involved in tissue repair and remodeling [5, 6]. Stimulation of the IFNγ receptor triggers Janus kinase (JAK)-mediated tyrosine phosphorylation and subsequent dimerization of (STAT)1[10]. IL-4Rα signals through a JAK-STAT6 pathway [11], and many of the genes associated with mouse M2 macrophages are regulated by STAT6, including arginase (Arg)1, macrophage mannose receptor 1 (also known as CD206), resistin-like-α (also known as Fizz1), and chitinase 3-like 3 (also known as Ym1) [12]. In addition, PI3K and PPARγ play an important role in M2 polarization [13, 14].

The polarization phenotype of macrophages in response to ethanol is complex, with increased expression of both M1 and M2 polarization markers as well as that of both pro- and anti-inflammatory mediators [15]. Lourvet et al. showed that activation of cannabinoid receptor 2 expressed in Kupffer cells inhibits M1 polarization and favors M2 response, thereby exerting anti-steatogenic roles by paracrine interactions with hepatocytes [15]. Limiting M1 macrophage polarization attenuates alcohol-induced liver injury, and IL-10 released from M2 polarized macrophages promotes M1 apoptosis [9]. Interestingly, ethanol-induced M1 polarization is modifiable; M1 macrophages can be shifted to an M2 phenotype [16]. However, little is currently known regarding the mechanisms of hyperpolarization and the impact of these complex phenotypes on the development of chronic ethanol-induced inflammation in the liver.

MicroRNAs (miRNAs), endogenous small non-coding RNAs, can inhibit target gene expression by directly binding to complementary sequences in the 3′ untranslated region (3′UTR) of mRNAs and thus are involved in many physiological processes, including cellular differentiation and immune cell activation [17]. miRNAs may play pivotal roles in modulating macrophage activation, polarization, tissue infiltration, and resolution of inflammation [18–20].

Prostaglandins are key lipid mediators that regulate physiological functions and inflammatory responses in health and disease [21]. They are synthesized from arachidonic acid via cyclooxygenases (1 and 2) to generate unstable prostaglandin H₂ (PGH₂), which is further processed by terminal synthases into the major prostaglandins (PGH₂, PGE₂, PGD₂, and PGF₂α) or thromboxane A₂ [21]. The PGHS pathway is frequently dysregulated in cancers [22]. PGI₂ synthase (PTGIS), also known as PGIS or CYP8A1, is a member of family 8 (CYP8) in the cytochrome P450 superfamily. PTGIS catalyzes the conversion of PGH₂ to PGI₂ [23]. Altered PTGIS plays an important role in preventing tumor growth and progression [24–27]. In addition, PTGIS plays an anti-inflammatory role in colorectal cancer [28], and prostaglandin G/H synthase 2 involved in the initial response to LPS stimulation [29]. However, whether PTGIS is involved in regulating macrophage polarization in ALD has not yet been determined. Therefore, in this study, we investigated whether PTGIS is involved in regulating macrophage polarization in ALD.

Materials and methods

Murine model of ALD

Six-to-eight-week-old male C57BL/6j mice (18–22 g weight) were purchased from the Laboratory Animal Center of Anhui Medical University. All animal procedures were reviewed and approved by the Institutional Animal Experimental Ethics Committee. Mice were housed in a temperature-controlled room (22 °C). After a week of adaptive breeding, mice were randomly divided into control diet (CD) and EtOH-fed groups (eight mice per group).

A mouse model of chronic plus single binge ethanol gavage was used as previously described [30]. For chronic alcohol administration, in the first 5 days, all mice were fed a liquid control diet. Thereafter, EtOH-fed mice were fed a liquid diet containing 5% v/v ethanol for 10 days, whereas the CD-fed mice were fed isocaloric maltose-dextrin. On day 16, EtOH-fed mice were administrated a single dose of ethanol gavage (5 g/kg body weight, 20% ethanol), while CD-fed mice were administrated an isocaloric dose of dextrin maltose gavage. Nine hours after gavage, the mice were euthanized, and blood and liver tissue samples were collected for further analysis.

Isolation of primary macrophages

Liver macrophages were isolated following a previously established method [31]. Briefly, a 20-G catheter was inserted through the mouse portal vein, and the inferior vena cava was cut. The liver was perfused with PB (40 mL of PBC diluted in 1 L ddH₂O). The PBC solution preparation method was as follows: NaCl (103.75 g), KCl (6.25 g), and Hepes (28.70 g) were dissolved in 350 mL H₂O; 1 M NaOH (75 mL) was added, and ddH₂O was subsequently added to make up a volume of 500 mL. Next, the liver was perfused with digestion buffer (type IV collagenase [35 mg], pronase [35 mg], 1 M CaCl₂ solution [1 mL], PB solution [99 mL]). After digestion, the liver was disrupted in 1% bovine serum albumin (BSA) solution. Single cells were passed through a 200-mesh sieve cell strainer and fractionated using 25 and 50% Percoll (Sigma-Aldrich, St. Louis, USA). The interface fraction was washed and adhered to plastic in medium. Macrophages adhered to the plastic, and the nonadherent fraction was washed off with Dulbecco’s Modified Eagle’s Medium (DMEM, Hyclone, China) with 15% fetal bovine...
serum (FBS; Biological Industries, Israel). The non-adherent cells were removed, and the remaining adhered cells were considered macrophages. Macrophages from three mice were pooled for protein and RNA isolation, because of the limited number of macrophages available from each animal.

**Cell culture and treatment**

RAW264.7 cells were cultured in DMEM containing 10% FBS at 37 °C in a humidified atmosphere containing 5% CO₂.

**Blood measurements**

The enzyme activities of alanine aminotransferase (ALT, C009-2-1), and aspartate aminotransferase (AST, C010-2-1), as well as triglyceride (TG, A-110-1-1) and total cholesterol (T-CHO, F002-1-1) levels were detected using commercial assay kits according to the manufacturers’ introductions (Nanjing Jiancheng Bioengineering Institute, Nanjing, China). The absorbance values at 510 nm were obtained with a micro-plate reader model 680 (Bio-Rad Laboratories, Hercules, CA, USA).

**Histopathology**

The specimen was sequentially fixed in 10% formalin for 24–48 h and embedded in paraffin for histopathological analysis. Thin sections (5 μM) were stained with hematoxylin and eosin staining (H&E) for histopathological analysis. The stained tissues were scanned using Panoramic MIDI (3D HISTECH, Hungary) and viewed using CaseViewer slice software.

**Oil-Red O staining**

Fresh liver tissues were immersed in freezing medium and stored at −80 °C. The liver tissues were sliced, washed with PBS three times (5 min), and stained with Oil Red O reagent according to the manufacturer’s protocol (Nanjing Jiancheng Bioengineering Institute, Nanjing, China). The stained tissues were scanned using Panoramic MIDI (3D HISTECH, Hungary) and viewed using CaseViewer slice software.

**Immunohistochemistry (IHC)**

IHC was performed on paraffin sections to examine the expression of F4/80 and PTGIS using a microwave-based antigen retrieval technique. Sections were incubated with rabbit anti-F4/80 (1:50 dilution) and anti-PTGIS (1:200 dilution) overnight at 4 °C. After incubation with the secondary antibody and 3, 30-diaminobenzidine tetrahydrochloride, the slides were counterstained with hematoxylin. Non-immune rabbit IgG instead of the primary antibody was used for the negative control (NC). The stained tissues were examined with an inverted fluorescence microscope (OLYMPUS IX83, Tokyo, Japan) or scanned using Panoramic MIDI (3D HISTECH, Hungary) and viewed using CaseViewer slice software.

**Transfection of PTGIS overexpression plasmid constructs and PTGIS siRNA into RAW264.7 cells**

Overexpression or knockdown of PTGIS was assessed through western blotting and real-time quantitative PCR (RT-qPCR) analysis after transfection of RAW264.7 cells with PTGIS overexpression plasmids or siRNA. All siRNAs and plasmid constructs were obtained from GenePharma Co., Ltd. Cells (3 × 10⁵/mL) were seeded in six-well plates and transfected with the PTGIS plasmid or siRNA and control constructs were mixed with lipofectamine 2000 transfection reagent (Invitrogen, Carlsbad, CA, USA) according to the manufacturer’s protocol. Cells were incubated with Opti-MEM at 37 °C and 5% CO₂ for 6 h. Subsequently, cells were cultured in DMEM containing 10% FBS. The PTGIS overexpressing or knockdown cells were treated with LPS (1 μg/mL) and IFNγ (10 ng/mL) for 24 h to polarize the M1 phenotype, or were cultured with IL-4 (15 ng/mL) for 24 h to induce M2 macrophages. The cells were subsequently harvested for western blotting, RT-qPCR, or other analyses.

**ELISA assay**

The concentrations of IL-1β, MCP-1, IL-10, and TGF-β in the serum and culture supernatant were determined using Joyee mouse ELISA kit (Joyee Biotechnics Co., Ltd, Anhui, China) according to the manufacturer’s introductions. Optical density values were measured at 450 nm. Three independent experiments were performed, and each sample was quantified with three replicates.

**Immunofluorescence**

To determine the expression of PTGIS in vivo, immunofluorescence staining was performed. Briefly, freshly dissected liver tissues were embedded in opti-mum cutting temperature compound, and the sections (8 μm thick) were cut with a cryotome Cryostat (Leica, CM 1950, Germany). Frozen tissue sections were incubated with 10% BSA for 1 h at room temperature (RT) to block nonspecific staining. After rinsing, the slides were incubated with primary antibodies for PTGIS (1:200 dilution) overnight at 4 °C. The slides were then washed twice with PBS (5 min) and
incubated with FITC (green)-conjugated secondary antibody (1:50 dilution) for 1.5 h at RT in the dark.

RAW264.7 cells were fixed and permeabilized in 4% paraformaldehyde and 0.2% TritonX-100 in PBS for 10 min. Nonspecific binding was blocked with 10% BSA for 1 h at RT. Subsequently, the cells were incubated with primary antibodies for PTGIS (1:200 dilution) overnight at 4 °C. Sections were washed twice with PBS and incubated with fluorescein-labeled secondary antibody at a dilution of 1:50 for 1.5 h at RT in the dark. Slides were mounted in mounting media with DAPI for 5 min at RT. After washing twice with PBS, the slides were covered with DABCO and the stained tissues were examined with an inverted fluorescence microscope (OLYMPUS IX83, Tokyo, Japan). Fluorescence density was measured using Image Pro Plus 6.

Western blotting analysis

Total proteins from cultured RAW264.7 cells and primary macrophages cells were extracted with radioimmunoprecipitation lysis buffer (containing 1% phenylmethylsulfonyl fluoride; Beyotime, China). The concentration of protein was determined using a bicinchoninic acid protein assay kit (Beyotime, Jiangsu, China). Proteins of each sample (30–50 μg) were separated via sodium dodecyl sulfate-polyacrylamide gel electrophoresis (10%) and transferred onto polyvinylidene fluoride (PVDF) membranes (Millipore Corp, Billerica, MA, USA). The membranes were blocked in 5% skim milk for 3 h to block nonspecific binding. Subsequently, membranes were incubated with the primary antibodies against PTGIS (Abcam, ab23668, 1:800), STAT1 (Bioworld, P42224, 1:800), p-STAT1 (Bioworld, AP0246P, 1:800), STAT6 (Bioss, bsm-52239R, 1:800), p-STAT6 (Bioworld, BS4181, 1:800) and β-actin overnight at 4 °C, followed by incubation with secondary antibody at RT for 1 h. Signals were captured with the ChemiDoc™ Image System (Bio-Rad, USA). The intensities of the western blotting bands were quantified and analyzed using the ImageJ software (NIH, Bethesda, MD, USA).

RNA extraction and RT-qPCR analysis

Total RNA was extracted from primary macrophages isolated from a mouse model, and cultured RAW264.7 cells were using TRIzol (Invitrogen, USA) according to the manufacturer’s instructions. The total RNA was uniformly quantified to 500 ng/μL using a Thermo Scientific NanoDrop 2000 Spectrophotometer (Thermo Scientific, USA). Single-stranded complementary DNA was synthesized from total RNA using the PrimeScript® RT reagent kit (TaKaRa, Japan) according to the manufacturer’s instructions. SYBR-Green RT-qPCR analysis was performed for Ptgis, Il-1β, Tnf-α, Il-6, iNo2, Il-10, Tgf-β, Arg-1, Cd163, and Gapdh mRNAs using SYBR Green Master Mix (TaKaRa, Japan) and the CFX Connect™ Real-Time System (Bio-Rad, USA). The primers used in this study were as follows: mus Ptgis: Forward: 5′-TCCTCAGAATCCGGAGCC-3′, Reverse: 5′-TCTTCTGTGGGAGTGTGGTC-3′; mus Il-1β: Forward: 5′-GAAGAGAGGCCCATCCTCTG-3′, Reverse: 5′-TCATCTCGAGGCCTGTAGTG-3′; mus Tnf-α: Forward: 5′-GACAGTGACCGGTACGTGG-3′, Reverse: 5′-TGAGACAGAGGCCAACCTGC-3′; mus Il-6: Forward: 5′-AGTCCGGAGAGGACAGTTCA-3′, Reverse: 5′-ATTTCCAGCTTCCAGAGGACGAG-3′; mus iNos: Forward: 5′-CCTTGTTTCAGCTACGCTTC-3′, Reverse: 5′-CTTCAGAGTCTGCCCATTCG-3′; mus Il-10: Forward: 5′-TGCAACTCCAAAGCCCAAG-3′, Reverse: 5′-TCAGTAAAGACAGGCAGCAT-3′; mus Tgf-β: Forward: 5′-CTGCTTCAGCTCCACAGAGA-3′, Reverse: 5′-CAGAAGTTGGCATGTAGCC-3′; mus Arg-1: Forward: 5′-GCAGTTGGGACATCTCCTG-3′, Reverse: 5′-GAGAAGGACACAGGTTGCC-3′; mus Cd163: Forward: 5′-ATGGTGGACAGAAATGAGT-3′, Reverse: 5′-AGCTCCAGCACCACAAG-3′; mus Gapdh: Forward: 5′-GGTTCCAGGCTTGAAGAGA-3′, Reverse: 5′-CCAATACGCGCAATCGGT-3′. The ratio for the mRNA of interest index was normalized to Gapdh and expressed as the mean ± s.e.m.

Recombinant adeno-associated virus-mediated PTGIS overexpression in mice

Mouse PTGIS over-expression plasmid labeled with green fluorescent protein (GFP) was obtained from Hanbio Biotechnology Co., Ltd. (Shanghai, China). The PTGIS plasmid was packaged with recombinant adeno-associated-virus for overexpression of PTGIS in vivo. Male C57BL/6j mice (age, six-to-eight weeks, weight, 18-22 g) were housed at the Animal Experimental Center for one week to adapt to the environment. The tails of mice were wiped with alcohol to expand the tail vein for injection. Mice were slowly injected with 100 μL of recombinant adeno-associated-virus-packaged PTGIS over-expression plasmid at a concentration of 1 × 10¹¹ v.g/mL/mouse through the tail vein using a 0.5 mL insulin syringe. One week later, mice were fed either control diet or EtOH diet.

RNA sequencing (RNA-seq) and computational analysis

Briefly, RAW264.7 cells transfected with pEX3 and pEX3-PTGIS plasmid were collected and lysed with TRIzol reagent (Invitrogen, USA) according to the manufacturer’s instructions. RNA was sequenced using the Illumina HiSeq 1000 at Novogene in Beijing (https://en.novogene.com). Raw data were analyzed using Illumina Casava 1.8, and...
three types of sequence reads—reads with adapter, no exact base information, or low-quality reads—were filtered and removed. RNA (3 μg) was used for library preparation with the NEBNext® UltraTM RNA Library Prep Kit for Illumina® (NEB, USA). The reference genome and gene model annotation files were downloaded directly from the genome website. The index of the reference genome was built using Hisat2 v2.0.5, and paired-end clean reads were aligned to the reference genome using Hisat2 v2.0.5. Differential expression analysis of two conditions/groups (two biological replicates per condition) was performed using the DESeq2 R package (1.16.1). The threshold we used to screen upregulated or downregulated mRNAs was log₂ fold-change | > 1 and p value < 0.05.

Bioinformatic analyses

The potential miRNA binding sites of PTGIS were obtained using the TargetScan 7.1 software (http://www.targetscan.org/). The protein interactions with PTGIS, JAK1, JAK2, and JAK3 were predicted using the online STRING database (https://string-db.org/).

Dual-luciferase reporter assay

The binding site of 3′-UTR-PTGIS and miRNA-140-3p.1 was detected via the dual-luciferase reporter assay. Briefly, the wt-PTGIS and mut-PTGIS sequences were inserted into the KpnI and HindIII sites of the pSI-Check2 promoter vector (Hanbio Biotechnology Co., Ltd., Shanghai, China) in a dual-luciferase reporter assay. Finally, the 293 T cells were grown in 96-well plates until they reached 50–70% confluence. Subsequently, 0.16 μg PTGIS plasmid was co-transfected with mmu-miR-140-3p.1 mimics/NC. The dual-luciferase assay was carried out 48 h after transfection. The experiment was independently performed three times.

Statistical analysis

Data are expressed as the mean ± s.e.m. One-way analysis of variance and the Newman–Keuls post hoc test (Prism 8.3 GraphPad Software, Inc, San Diego, CA, USA) were used to analyze the results. p < 0.05 was considered statistically significant.

Results

Alcohol consumption causes liver injury, steatosis, and inflammation

Mice with chronic ethanol feeding were characterized by liver injury. To investigate the degree of liver injury induced by ethanol consumption, we performed H&E staining and Oil Red O staining, measured serum ALT, AST, TG, and T-CHO levels, and determined the body weights and liver-to-body weight ratios. Histopathological analysis showed extensive steatosis, liver cell cord derangement, and intercellular space dilatation in EtOH-fed mice (Fig. 1A). Serum ALT, AST, TG, and T-CHO levels were substantially elevated in the EtOH-fed mouse group (Fig. 1B). As shown in Fig. 1C, body weight decreased in the early stage and subsequently increased in the EtOH-fed group. However, the liver-to-body weight ratio was elevated in EtOH-fed group (Fig. 1D). We further investigated macrophages infiltration in liver tissues using IHC analysis. Intriguingly, chronic alcohol feeding caused a marked elevation in the total number of macrophages in C57BL/6j mice, as assessed via F4/80 immunostaining (Fig. 1A), associated with an obvious increase in M1-polarized macrophages and a moderate rise of M2-polarized macrophages (Fig. 1E and F). The in vitro results were consistent with the in vivo results (Fig. 1G). The results indicate that chronic alcohol consumption leads to liver injury, steatosis, and significant macrophage M1 polarization in vivo and in vitro.

Chronic alcohol consumption decreases PTGIS expression both in vivo and in vitro

To identify the expression changes of PTGIS induced by chronic alcohol consumption, western blotting, RT-qPCR, and IHC analyses were performed. As shown in Fig. 1H–J, PTGIS expression significantly decreased in the EtOH-fed group. As shown in Fig. 1G, PTGIS mRNA expression was significantly decreased in RAW264.7 cells cultured with EtOH and LPS. The immunosignals of PTGIS in EtOH-fed mice substantially decreased (Fig. 1J). The immunofluorescence signal of PTGIS significantly decreased in M1-polarized RAW264.7 cells and increased in M2-polarized RAW264.7 cells (Fig. 1K). These results indicate that chronic alcohol consumption decreased PTGIS expression both in vivo and in vitro.

Recombinant adeno-associated virus-mediated overexpression of PTGIS protects against chronic alcohol consumption-induced liver injury, steatosis, and inflammation in vivo

The expression of PTGIS, both at protein and mRNA levels, was significantly downregulated by EtOH consumption both in vivo and in vitro. We investigated whether PTGIS-targeted therapy could prevent chronic alcohol-induced liver injury in vivo. The eGFP signal in the liver transfected with the PTGIS overexpression plasmid indicated a successful infection of the virus (Fig. S1A). PTGIS expression was substantially increased.
Fig. 1 PTGIS expression and macrophage polarization in vivo and in vitro. A Representative pictures of H&E, Oil Red O and F4/80 staining of liver tissue sections were shown. B Serum ALT, AST, TG, and T-CHO levels were measured. C Body weight were shown. D The ratio of liver-to-body weight. E Serum cytokines were detected by ELISA. F The mRNA levels of M1 markers and M2 markers in primary macrophages isolated from liver tissues were detected by RT-qPCR analysis. G PTGIS mRNA expression and M1, M2 markers in vitro were measured by RT-qPCR analysis. The mRNA (H) and protein levels (I) of PTGIS in primary macrophages were measured by RT-qPCR and western blotting analysis. J Representative pictures of PTGIS immune staining were shown. K Protein level of PTGIS in vitro was detected by immunofluorescence. Values represent the mean ± s.e.m for 3–4 independent experiments. *p < 0.05, **p < 0.01, ***p < 0.001 as indicated.
in primary macrophages isolated from recombinant adenoviral-associated viral vector (rAAV8)-PTGIS-transfected liver compared to rAAV8-empty-vector-treated liver tissue at both the protein and mRNA levels (Fig. 2A, B). As shown in Fig. 2C, overexpression of PTGIS in liver tissues alleviated alcohol-induced liver injury (H&E staining) and inhibited liver steatosis (Oil Red O staining) and macrophage infiltration (F4/80 staining) compared to empty-virus-treated mice in the EtOH-fed group. Forced expression of PTGIS dramatically elevated the body-weight and suppressed the liver-to-body weight ratio of mice compared to the empty-virus-infected group in the EtOH-fed group (Fig. S1B, S1C). As shown in Fig. 2D, E, forced PTGIS expression in vivo could inhibit M1 polarization and promote macrophage switch to the M2 phenotype. The ELISA results were consistent with those of the RT-qPCR analysis (Fig. 2F). These results indicated that forced expression of PTGIS in vivo inhibited M1
polarization and promoted M2 polarization and may be a potential therapeutic target for chronic alcohol-induced liver injury.

**The expression of PTGIS is regulated by miR-140-3p.1**

Bioinformatic tools (TargetScan) were used to screen for differentially expressed miRNAs targeting PTGIS. As shown in Fig. 3A, miR-140-3p.1 was the only one conserved and was found to be related to PTGIS expression. Next, we determined the expression of miR-140-3p.1 in the LPS- and EtOH-treated RAW264.7 cells via RT-qPCR. As shown in Fig. 3B, miR-140-3p.1 expression was substantially increased in the LPS- and EtOH-treated group. The transfection efficiency of miR-140-3p.1 mimics and inhibitors were shown in Fig. 3C. We further explored the effect of miR-140-3p.1 on alcohol-induced inflammation in RAW264.7 cells cultured with EtOH and LPS. As shown in Fig. S2A, miR-140-3p.1 mimics promoted M1 polarization and inhibited M2 phenotype, whereas miR-140-3p.1 inhibitors promoted M2 marker expression and inhibited M1 polarization (Fig. S2B). TargetScan showed that the 3′UTR of PTGIS included a putative conserved target site for miR-140-3p.1, and the binding site was conserved (Fig. 3D). Furthermore, miR-140-3p.1 decreased wild-type PTGIS 3′UTR luciferase activity, and the reduction was not observed in a PTGIS 3′UTR reporter, which had mutations in the miR-140-3p.1-binding site (Fig. 3D). In addition, we observed that miR-140-3p.1 mimics significantly decreased PTGIS expression, while miR-140-3p.1 inhibitors markedly elevated PTGIS expression at protein and mRNA levels in RAW264.7 cells cultured with LPS and EtOH (Fig. 3E–H). These results suggest that miR-140-3p.1 is upregulated in LPS- and EtOH-treated RAW264.7 cells and could regulate PTGIS expression.

**The influence of altered PTGIS expression on M1-polarized macrophages**

To investigate the influence of PTGIS on M1 polarization, we performed loss- and gain-of-function-experiments. The PTGIS plasmid significantly increased PTGIS expression at the mRNA and protein levels in M2-polarized macrophages (Fig. 5A, B), and forced PTGIS expression significantly elevated anti-inflammatory gene expression (Fig. 5C, D). Furthermore, PTGIS-siRNA significantly inhibited PTGIS expression in M2-polarized RAW264.7 cells (Fig. 5E, F), and PTGIS silencing induced decreased expression of anti-inflammatory genes (Fig. 5G, H). These results indicated that PTGIS could promote M2 macrophage polarization.

**PTGIS may regulate macrophage polarization via the JAK/STAT pathway**

To further evaluate the mechanism of PTGIS in regulating macrophage polarization, we forced PTGIS expression in RAW264.7 cells cultured with LPS and EtOH and subsequently isolated total mRNA for RNA-seq (Fig. 6A). As shown in Fig. 6B, we detected 11,244 genes in the PTGIS overexpression group and 11,123 genes in the pEX3-transfected group. As shown in the volcano plot, 290 mRNAs were upregulated and 99 mRNAs were downregulated in the RAW264.7 cells transfected with the PTGIS plasmid compared to those of the pEX3-transfected group (log2FC > 1, p < 0.05) (Fig. 6C). Gene ontology analysis revealed that PTGIS overexpression affected many genes associated with inflammatory responses, chemokine activity, chemokine receptor activity, and cytokine activity and so on (Fig. 6D). KEGG pathway analysis revealed that forced PTGIS expression could alter JAK-STAT signaling, NOD-like receptor signaling, PI3K-AKT signaling, cytokine-cytokine receptor interaction,
IL-17 signaling, etc. (Fig. 6E). As the JAK/STAT pathway plays an important role in macrophage polarization, we focused on the influence of forced PTGIS expression on the JAK/STAT pathway. The proteins interacting with JAK were predicted by the STRING database (Fig. 6F). The overlap between RNA-seq and the STRING database was IL-6 (Fig. 6G). Furthermore, we assessed the effects of altered PTGIS expression on JAK/STAT signaling via western blotting analysis (Fig. S4A–S4D). The results revealed that forced PTGIS expression inhibited JAK/STAT1 activation in M1-polarized macrophages and promoted JAK/STAT6 activation in M2-polarized macrophages (Fig. S4A and S4C). PTGIS silencing promoted JAK/STAT1 activation in M1-polarized macrophages and inhibited JAK/STAT6 activation in M2-polarized macrophages (Fig. S4B and S4D). These results suggest that PTGIS might regulate macrophage polarization through JAK/STAT signaling.
Discussion

Alcohol consumption is a leading risk factor for death and disability worldwide. A systematic analysis of the global burden of alcohol revealed that the level of consumption that minimizes health loss is zero [1]. ALD appears as a wide range of disorders, ranging from simple liver steatosis to more severe forms of liver injury, including alcoholic hepatitis, liver fibrosis, cirrhosis, and hepatocellular carcinoma [33]. Ethanol metabolism-associated oxidative stress, glutathione depletion, abnormal methionine metabolism, malnutrition, ethanol-mediated induction of gut endotoxin leakage, and subsequent activation of macrophages play important roles in the pathogenesis of ALD [34–39]. In this study, we illustrated for the first time, to our knowledge, that PTGIS decreased in a chronic alcohol-consuming mouse model, and overexpression of PTGIS in vivo alleviated liver steatosis, inflammatory cell infiltration, and macrophage switch to the M1 phenotype (Fig. 7).

Ethanol increases fatty acid synthesis in hepatocytes via the upregulation of sterol regulatory element-binding protein 1c [40], a transcription factor that promotes fatty acid
Fig. 5 The influence of alternative PTGIS expression on M2 polarization. A The mRNA levels of PTGIS in M2 polarized RAW264.7 cells transfected with PTGIS overexpression plasmid. B The protein levels of PTGIS in M2 polarized macrophages. C The mRNA levels of M2 markers in PTGIS overexpression group. D Serum cytokine of M2 markers were detected by ELISA. E PTGIS mRNA expression in M2 polarized RAW264.7 cells transfected with PTGIS-RNAi were measured by RT-qPCR. F The protein levels of PTGIS were detected. G The mRNA levels of M2 markers in PTGIS silencing group. H Serum cytokines of M2 markers were detected by ELISA. Values represent the mean ± s.e.m. *p < 0.05, **p < 0.01, ***p < 0.001, ###p < 0.001 as indicated.
Fig. 6 Expression profiles of mRNAs on pEX3 and pEX3-PTGIS overexpressed plasmid transfected RAW264.7 cells. A RNA-seq was performed with RNA extracts from RAW264.7 cells transfected with pEX3 or pEX3-PTGIS overexpressed plasmid. Hierarchical cluster analysis of significantly differentially expressed mRNAs: bright blue, under-expression; white, no change; bright red, over-expression. B Venn diagram showed mRNAs detected in pEX3 and PTGIS overexpression plasmid transfected group. C Volcano plot was presented. D, E GO and KEGG analysis of altered mRNAs in pEX3 and PTGIS overexpressed transfected group. F The proteins interacted with JAK were predicted by STRING database. G The overlap between RNA-seq and STRING was IL-6.
M2-polarized macrophages to increase simultaneously that chronic alcohol consumption induces M1- and in vitro (Fig. 1H
in vitro (Figs. 2D and 5E). Forced PTGIS expression inhibited JAK/STAT1 activation in M1-polarized macrophages and promoted JAK/STAT6 activation in M2-polarized macrophages (Fig. 4A S4A and S4C). PTGIS silencing promoted JAK/STAT1 activation in M1-polarized macrophages and inhibited JAK/STAT6 activation in M2-polarized macrophages (Fig. S4B S4D). The overlap between the altered genes associated with PTGIS and the genes interacted with JAKs was IL-6 (Fig. 6G). The activation of STAT1 (pSTAT1) and STAT3 (pSTAT3) plays a pivotal role in controlling liver steatosis and liver inflammation [51]. STAT3, which is activated by IL-6, plays an important role in hepatoprotection, liver regeneration, and glucose homeostasis by inducing a variety of anti-apoptotic and mitogenic proteins [51]. Miller et al. demonstrated that deletion of IL-6 or hepatic STAT3 increased STAT1 activation in HFD-fed IL-10−/− mice [33]. IL-10 is a well-documented anti-inflammatory cytokine and IL-6 is a pro-inflammatory cytokine [52]. However, IL-6 can act as an anti-inflammatory cytokine by inhibiting the activation of STAT1 [33]. Therefore, we hypothesized that chronic alcohol consumption leads to liver injury and inflammation and forced PTGIS expression can inhibit M1 macrophage polarization by inhibiting the activation of STAT1 through IL-6, and IL-6 acts as an anti-inflammatory cytokine on this occasion.

A range of evidence over the last decade indicates a pivotal role of miRNAs as key mediators of macrophage differentiation, infiltration, and activation. Initial studies have indicated the property of miRNAs to regulate the magnitude of the innate response, participating as integral components of feedback loop regulatory mechanisms, which significantly shape the inflammatory response [53]. In addition, recent studies have demonstrated that miRNAs regulate macrophage differentiation and polarization [54]. In short, a complex and highly regulated network of miRNAs regulating inflammatory pathways by targeting components of TLR signaling and influencing downstream inflammatory cytokine production has been defined. In this study, we demonstrated that the regulation of PTGIS in macrophage polarization could be influenced by miRNA-140-3p.1 (Fig. 3).

Macrophages are hindered by their versatility, and can adapt their phenotype (to become pro-inflammatory or anti-inflammatory) in response to molecular signals in healthy or injured livers. Historically, the plasticity of macrophages has been described using the terms classically activated (M1 type) and alternatively activated (M2 type). However, this nomenclature is too simple to describe the polarization

**Fig. 7 Schematic presentation of the effects of PTGIS on macrophage polarization on alcohol induces liver injury.** Ethanol and LPS induced miR-140-3p.1 expression. Subsequently, miR-140-3p.1 inhibited PTGIS expression. Overexpression of PTGIS in vivo and in vitro inhibits macrophage M1 polarization and promotes M2 polarization through JAK-STAT signaling.

synthesis via the upregulation of lipogenic genes and the inhibition of PPARα [41, 42], a nuclear hormone receptor that controls the transcription of a range of genes involved in free fatty acid transport and oxidation. In addition, ethanol can affect the activities of enzymes involved in fat metabolism by inhibiting AMP-activated protein kinase, which reduces fat metabolism and fatty liver [43, 44]. In this study, we demonstrated that PTGIS overexpression in vivo alleviated fatty acid accumulation in the liver, as determined by Oil Red O staining (Figs. 1A and 2C).

Alcohol consumption not only causes enteric dysbiosis and bacterial over-growth [45] but also increases gut permeability and the translocation of bacteria-derived LPS from the gut to the liver [46]. LPS interacts with Toll-like receptor 4 (TLR4) to activate the MyD88-independent signaling pathway, leading to the production of pro-inflammatory cytokines, including IL-1β and TNF-α [47–49]. Interestingly, activation of TLR4 and complement factors also causes macrophages to produce hepatoprotective cytokines, such as IL-6, and anti-inflammatory cytokines, such as IL-10 [33, 50]. In this study, the results indicated that chronic alcohol consumption induces M1- and M2-polarized macrophages to increase simultaneously (Fig. 1E–G) and inhibit PTGIS expression in vivo and in vitro (Fig. 1H–K). Overexpression of PTGIS could alleviate liver injury in vivo (Figs. 2C, S1B, and S1C). Forced PTGIS expression inhibited macrophage M1 polarization and promoted M2 polarization both in vivo and in vitro (Figs. 2D–F, 4, and 5). A range of evidence demonstrated that the JAK/STATs signaling pathway plays a pivotal role in regulating macrophage polarization. RNA-seq analysis revealed that the genes regulated by PTGIS were enriched in the JAK/STAT pathway (Fig. 6D and E). Forced PTGIS expression inhibited JAK/STAT1 activation in M1-polarized macrophages and promoted JAK/STAT6 activation in M2-polarized macrophages (Fig. S4A and S4C). PTGIS silencing promoted JAK/STAT1 activation in M1-polarized macrophages and inhibited JAK/STAT6 activation in M2-polarized macrophages (Fig. S4B and S4D). The overlap between the altered genes associated with PTGIS and the genes interacted with JAKs was IL-6 (Fig. 6G). The activation of STAT1 (pSTAT1) and STAT3 (pSTAT3) plays a pivotal role in controlling liver steatosis and liver inflammation [51]. STAT3, which is activated by IL-6, plays an important role in hepatoprotection, liver regeneration, and glucose homeostasis by inducing a variety of anti-apoptotic and mitogenic proteins [51]. Miller et al. demonstrated that deletion of IL-6 or hepatic STAT3 increased STAT1 activation in HFD-fed IL-10−/− mice [33]. IL-10 is a well-documented anti-inflammatory cytokine and IL-6 is a pro-inflammatory cytokine [52]. However, IL-6 can act as an anti-inflammatory cytokine by inhibiting the activation of STAT1 [33]. Therefore, we hypothesized that chronic alcohol consumption leads to liver injury and inflammation and forced PTGIS expression can inhibit M1 macrophage polarization by inhibiting the activation of STAT1 through IL-6, and IL-6 acts as an anti-inflammatory cytokine on this occasion.

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Macrophages are hindered by their versatility, and can adapt their phenotype (to become pro-inflammatory or anti-inflammatory) in response to molecular signals in healthy or injured livers. Historically, the plasticity of macrophages has been described using the terms classically activated (M1 type) and alternatively activated (M2 type). However, this nomenclature is too simple to describe the polarization
phenotypes of macrophages driven by many different stimuli in the microenvironment. Ly-6C is a cell surface glycoprotein widely used to identify functionally discrete murine circulating monocyte populations: Ly-6C<sup>hi</sup> monocytes and Ly-6C<sup>lo</sup>. Ly-6C<sup>hi</sup> monocytes are recruited early to inflammatory environments and are thought to be proinflammatory, whereas Ly-6C<sup>lo</sup> monocytes are a patrolling cell type and can supplement resident tissue macrophages [55, 56]. Further studies are needed to investigate the influence of altered PTGIS expression on differential Ly-6C expression in chronic ALD.

In conclusion, as shown in Fig. 7, we demonstrated that PTGIS expression was downregulated in chronic alcohol-consuming mice via elevated miR-140-3p.1 expression. Forced PTGIS expression in vivo and in vitro can inhibit macrophage switch to the M1 phenotype and promote M2 polarization. RNA-seq results showed that PTGIS might regulate macrophage polarization through the JAK/STAT pathway, and IL-6 might be involved in inhibiting the activation of STAT1. This study revealed that PTGIS might be a therapeutic target for ALD.

Data availability

The datasets used and analyzed in this study are available from the corresponding author on reasonable request.

Author contributions X.Y.P. conceived and carried out experiments, analyzed the data, and wrote the manuscript. L.W., H.M.Y. and M.C. helped to model establishment and isolated primary macrophages. Y.Y. and C.H. participated in the design of the study. J.L. conceived the study and revised the manuscript.

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Compliance with ethical standards

Conflict of interest The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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