Histone deacetylase inhibitor givinostat alleviates liver fibrosis by regulating hepatic stellate cell activation

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Abstract. Hepatic fibrosis, a common pathological manifestation of chronic liver injury, is generally considered to be the end result of an increase in extracellular matrix produced by activated hepatic stellate cells (HSCs). The aim of the present study was to target the mechanisms underlying HSC activation in order to provide a powerful therapeutic strategy for the prevention and treatment of liver fibrosis. In the present study, a high-throughput screening assay was established, and the histone deacetylase inhibitor givinostat was identified as a potent inhibitor of HSC activation in vitro. Givinostat significantly inhibited HSC activation in vivo, ameliorated carbon tetrachloride-induced mouse liver fibrosis and lowered plasma aminotransferases. Transcriptomic analysis revealed the most significantly regulated genes in the givinostat treatment group in comparison with those in the solvent group, among which, dermokine (Dmkn), mesothelin (Msln) and uroplakin-3b (Upk3b) were identified as potential regulators of HSC activation. Givinostat significantly reduced the mRNA expression of Dmkn, Msln and Upk3b in both a mouse liver fibrosis model and in HSC-LX2 cells. Knockdown of any of the aforementioned genes inhibited the TGF-β1-induced expression of α-smooth muscle actin and collagen type I, indicating that they are crucial for HSC activation. In summary, using a novel strategy targeting HSC activation, the present study identified a potential epigenetic drug for the treatment of hepatic fibrosis and revealed novel regulators of HSC activation.

Introduction

Cirrhosis is an increasing global health burden that accounts for >100 million deaths annually worldwide (1). Liver fibrosis is the result of wound-healing response to chronic liver impairment triggered by a variety of causes, including hepatitis virus, ethanol, drugs and poisons, parasites, metabolism and genetics, cholestasis and immune deregulation (2,3). Without diagnosis and treatment, hepatic fibrosis will ultimately progress to hepatic cirrhosis, and even to hepatocellular carcinoma (4). Thus, it is of great importance to actively intervene in liver fibrosis. Hepatic fibrosis is characterized by the deposition of extracellular matrix (ECM) proteins, which destroy the normal liver histological structure and functions (5). Hepatic stellate cells (HSCs) play a vital role in the development of liver fibrosis, and are the main producers of ECM (5). In the case of liver injury, certain cytokines and growth factors crucial for HSC activation are released, and promote HSC activation into myofibroblasts, which are responsible for the synthesis of ECM proteins, including α-smooth muscle actin (α-SMA, which is encoded by Acta2), collagen type I (Col1a1), matrix metalloproteinases and tissue inhibitor of metalloproteinases (6). Therefore, directly inactivating HSCs is of great importance for fibrosis resolution, representing a therapeutic strategy for the treatment of hepatic fibrosis.

Epigenetic modifications regulate patterns of gene expression by modulating DNA accessibility and chromatin structure. The epigenetic machinery, particularly certain epigenetic enzymes, has been demonstrated to be involved in myofibroblast activation and regulation of fibrotic gene expression (7,8). Blocking the expression of the DNA methyltransferase DNMT3B has been reported to significantly reduce α-SMA and Col1a1 expression in ischemic heart disease (9). In addition, the histone deacetylase (HDAC)
inhibitors (HDACis), MS-275 and trichostatin A have been found to reduce renal fibrosis by diminishing the accumulation of ECM proteins (10–12). By contrast, other studies have indicated that targeted inhibition of certain epigenetic enzymes might aggravate fibrosis. It has been reported that inhibition of type I protein arginine methyltransferases can aggravate renal fibrosis by reducing asymmetric dimethylarginine accumulation, increasing nitric oxide concentrations and enhancing the expression of profibrotic proteins (13). However, there is currently no effective high-throughput screening method to identify candidate compounds for the treatment and prevention of liver fibrotic diseases.

Aiming to identify a novel candidate compound for the treatment of hepatic fibrosis, the present study established a cell-based high-throughput assay based on HSC activation, and screened our in-house epigenetic compound library (14). The HDACi givinostat, which has been used in phase I/II clinical trials for the treatment of Duchenne muscular dystrophy (15), was identified as the most potent hit. Givinostat reduced the expression of α-SMA, fibronectin and collagen, which are markers of HSC activation in vitro. Carbon tetrachloride (CCl4) has been widely used to induce liver injury and fibrosis in mice for decades (16), and C57BL/6J inbred mice are frequently used for fibrosis studies in the CCl4 model of the ready availability of genetically-modified mice (17). In a chronic CCl4-challenged mouse model in the present study, mice developed mild liver fibrosis after 2 weeks of CCl4 treatment, and were then treated with givinostat for 6 weeks. Givinostat significantly ameliorated CCl4-induced mouse liver injury and fibrosis. RNA-sequencing (RNA-seq) analysis of the liver tissue from the givinostat treatment and solvent groups of CCl4-challenged mice revealed genes regulated by givinostat treatment, among which, dermokine (Dmkn), mesothelin (Msln) and uroplakin-3b (Upk3b) were further identified as crucial genes regulating HSC activation. Givinostat inhibited HSC activation and alleviated liver fibrosis in vivo and in vitro, making it a promising tool for developing a novel therapy for the treatment of hepatic fibrosis.

Materials and methods

Animals and treatment. Female C57BL/6J mice (8–9 weeks old, weighting 21–23 g, specific pathogen-free) were purchased from the Animal Center of the Chinese Academy of Medical Sciences. Animal care was carried out according to the guidelines of the Principles of Laboratory Animal Care (18), all experimental protocols were approved by the Institutional Animal Care and Use Committee at the Shanghai Institute of Materia Medica (approval no. 2018-12-002; Shanghai, China). The animals were fed a standard laboratory diet and tap water. All mice were kept in standard laboratory conditions (21±2°C, 12-h light/dark cycle), and were fed adaptively for 1 week before starting the experiments. A total of 24 mice were randomly divided into three groups with 8 mice per group: i) The normal control group; ii) the solvent group of CCl4-challenged mice; and iii) the givinostat treatment group of CCl4-challenged mice. In the givinostat treatment group, mice were i.p. injected with 10% CCl4 dissolved in olive oil at a dose of 1 ml/kg body weight twice a week for 8 weeks to trigger liver fibrosis (20). Givinostat or PBS solvent was i.p. injected after CCl4 treatment for 2 weeks, when mild fibrosis was shown. At the end of the experiment, the mice were sacrificed, and blood as well as liver samples were harvested. Although there was a total of 24 mice used overall, too little blood was collected during blood collection to be used for experiments so the number of experimental results displayed was n=8 in normal control group, n=6 in CCl4 group and n=7 in the CCl4 + givinostat group. All surgeries (blood and liver samples were harvested) were performed under sodium pentobarbital anesthesia (50 mg/kg), and then all mice were euthanized by 5% isoflurane (cat. no. HR135327; Haurui Chemical). Death of the mice was confirmed by checking whether their heartbeat had completely stopped and whether their pupils were dilated.

Liver histopathology and immunohistochemistry. Liver tissues were fixed in 4% paraformaldehyde for 24 h at 37°C, dehydrated and paraffin embedded. The liver tissue sections were deparaffinized using xylene (Wuhan Servicebio Technology Co., Ltd.), rehydrated with graded alcohol, treated with 0.3% endogenous peroxidase blocking solution (Sigma-Aldrich; Merck KGaA) for 10 min. Following high pressure freezing retrieval (125°C and 103 kPa) and blocking with 10% normal goat serum (Wuhan Servicebio Technology Co., Ltd.) at 37°C for 30 min, the sections were incubated overnight at 4°C with the following primary antibodies (Wuhan Servicebio Technology Co., Ltd.): anti-α-SMA (cat. no. GB13044; 1:100) and anti-Col1α1 (cat. no. GB11022-1; 1:100). Following washing with PBS, goat anti-rabbit non-biotinylated reagents (cat. no. G1213; 1:1,000; Wuhan Servicebio Technology Co., Ltd.) were used to react with the primary antibody for 2 h at 37°C. Images were captured by observers who were blinded to the experimental conditions at 6–8 non-consecutive random fields under a light microscope (magnification, x100), and were used to assess the histological changes using Image-Pro Plus 6.0 software (Media Cybernetics, Inc.). Representative views were displayed.

Cell culture. The human HSC LX-2 cell line and the rat HSC-T6 cell line were obtained from the FuHeng Cell Center, and were cultured in Dulbecco's modified Eagle's medium (DMEM; cat. no. L101KJ; Shanghai BasalMedia Technologies Co., Ltd.) supplemented with 10% fetal bovine serum (Gibco; Thermo Fisher Scientific, Inc.) and 1% penicillin and streptomycin (Gibco; Thermo Fisher Scientific, Inc.), at 37°C in a 95% air humidified atmosphere containing 5% CO2. For stimulation, the cells were starved in serum-free DMEM for 24 h before being treated with recombinant human TGF-β1 (10 ng/ml; cat. no. 100-21C; PeproTech, Inc.) and/or givinostat (900, 300 or 100 nM; cat. no. CSN16577; CSNpharm) for 24 h.

One-step reverse transcription-quantitative PCR (RT-qPCR). HSC LX-2 cells (5x10⁵ cells/well) were seeded in 96-well cell
culture plates (Cellvis) overnight and sequentially stimulated with TGF-β1 and compounds (2 µM) or with TGF-β1 solely. Cells were harvested 24 h after TGF-β1 stimulation with Total RNA extraction reagent (Vazyme Biotech Co., Ltd.). Cytolysis was then subjected to RT-qPCR using Transcript® Green One-Step qRT-PCR SuperMix (cat. no. AQ211; TransGen Biotech Co., Ltd.) (14). The compounds library containing 46 molecule probes targeting epigenetic proteins was screened to find small molecule compounds able to inhibit α-SMA expression.

**RNA-seq analysis.** Total RNA was isolated from flash-frozen mice liver tissues. Total RNA was isolated and purified using DNase I (Takara Bio, Inc.) and Dynabeads Oligo (dT) 25 (Thermo Fisher Scientific, Inc.). Subsequently, purified RNA (100 ng) was used for cDNA library construction, using the NEBNext Ultra™ RNA Library Prep kit for Illumina® (cat. no. E7530L; New England Biolabs, Inc.). Sequencing data was collected on an Illumina HiSeq 2500 instrument. The RNA integrity number (RIN) value was used to assess the quality of the isolated RNAs. Only RNAs with RIN ≥7.0 were used for sequencing. The sequencing reads were located to mm10 by STAR 2.5 (22), and gene counting was quantified using featureCounts (Subread package 2.0.0) (23). The edgeR R package (24) was used for differential gene expression analysis. The P-value was adjusted using the Benjamini and Hochberg method (25), and a 5% FDR cutoff value and fold-change >1.5 were set as the threshold of the significance analysis. The P-values were further analyzed by gene-annotation enrichment analysis using The Database for Annotation, Visualization and Integrated Discovery 6.8 bioinformatics platform (26). Cytoscape was used for network analysis (27). The original data generated using high-throughput sequencing methodologies has been submitted to the GEO database (https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE16981).

**Small interfering (si)RNA transfection.** Msln siRNA (sense, 5'-GCCUUGCUUGCAGAAACU-3' and antisense, 5'-AUG UUCGGAGAAAGGCG-3'); Msln antisense, 5'-GGACGUCCU AAAGCAUAA-3' and antisense, 5'-UUUAUGCUUGAG GACGUCC-3'), Dmkn siRNA (sense, 5'-GCAGAGACGAU CAGAACUA-3' and antisense, 5'-UGUUUCGCAGUCCGCUC UC-3' and sense, 5'-GCCUAUGGGGAAGAGACU-3' and antisense, 5'-AGACUCCUCCCACAUAGGC-3') and Upk3b siRNA (sense, 5'-GCCCUACACCAACACAA-3' and antisense, 5'-AUUCUGUGGGUGAGGC-3' and antisense, 5'-GGCUACUGAGCACCACCUAC-3') were synthesized by Shanghai GenePharma Co., Ltd. Transfection with siRNA against Msln, Upk3b or Dmkn, or with control siRNA (sense, 5'-UUUCUGCGAAGCGUCCUGGU-3' and antisense, 5'-ACUGAAGACGCUCCAGA-3') was performed according to the manufacturer's protocol. LX-2 cells were seeded in a 6-well plate at 60-80% confluence. Briefly, siRNA (20 µM, 1.5 µl) and 9 µl Lipofectamine® RNAiMAX transfection reagent (Invitrogen; Thermo Fisher Scientific, Inc.) were mixed with 150 µl Opti MEM (cat. no. 31985070; Gibco; Thermo Fisher Scientific, Inc.). Next, diluted siRNA was added to diluted Lipofectamine RNAiMAX reagent and cultured for 5 min at room temperature. siRNA-lipid complex was added to cells for 6-8 h at 37°C. Subsequent experiments were performed 24 h after transfection.

**RNA extraction and RT-qPCR.** Total RNA was extracted from HSC LX-2 cells, HSC-T6 cells or liver tissues using TRIzol® reagent (Invitrogen; Thermo Fisher Scientific, Inc.), according to the manufacturer's protocols. Total RNA (1,000 ng) was reverse transcribed into cDNA using a cDNA synthesis kit (Vazyme Biotech Co., Ltd.). The reverse transcription temperature protocol was as follows: 50°C for 15 min, followed by 80°C for 5 sec. RT-qPCR was performed using SYBR-Green (Vazyme Biotech Co., Ltd.). The thermocycling conditions were as follows: Pre-denaturation at 95°C for 10 min, 40 cycles of denaturation at 95°C for 15 sec, and annealing at 60°C for 30 sec, followed by extension at 72°C for 1 min. Subsequently, the expression values of mRNA were calculated using the 2^(-ΔΔCq) method (28). The expression of target genes was normalized to GAPDH expression. The primer sequences are shown in Table I.

**Biochemical analysis.** The levels of serum aspartate aminotransferase (AST; cat. no. C010-3-1) and alanine aminotransferase (ALT; cat. no. C009-3-1) were measured using the corresponding kits (Nanjing Jiancheng Bioengineering Institute) and were assessed using a Hitachi 7020 automatic analyzer (Hitachi, Ltd.).

**Western blotting.** Total protein was extracted from HSC LX-2 cells, HSC-T6 cells or liver tissues with 1X SDS sample loading buffer (250 mM Tris HCl pH 6.8, 10% SDS, 30% glycerol, 5% β-mercaptoethanol and 0.02% bromophenol blue). Protein concentration was determined using a BCA kit (Thermo Fisher Scientific, Inc.). The lysates (25 µg/lane) were separated via SDS-PAGE on 6, 10 or 12% gels, and subsequently transferred to a nitrocellulose membrane (EMD Millipore). Following blocking with 5% milk at room temperature for 1 h, membranes were incubated with primary antibodies at 4°C overnight. Subsequently, the membranes were incubated with a HRP-conjugated goat anti-Rabbit IgG secondary antibody (1:10,000; cat. no. D110058; Sangon Biotech Co., Ltd.) for 1 h at room temperature. The bands were visualized using an enhanced chemiluminescence kit (Thermo Fisher Scientific, Inc.) with a ChemiScope 3400 mini imaging system (Clinx Science Instruments Co., Ltd.). Densitometry was performed for each group using ImageJ software (v1.50b; National Institutes of Health). The following primary antibodies were used: Anti-α-SMA (1:1,000; cat. no. 19245; Cell Signaling Technology, Inc.), anti-Col1a1 (1:1,000; cat. no. 72026; Cell Signaling Technology, Inc.) and anti-GAPDH (1:5,000; cat. no. 8884; Cell Signaling Technology, Inc.), which was used as the loading control.

**Statistical analysis.** All numerical results are expressed as the mean ± standard deviation, and represent data from a minimum of three independent experiments. Two-tailed unpaired t-test was used to analyze differences between two groups. One-way ANOVA or two-way ANOVA were used to compare the means of multiple groups followed by LSD post hoc test. All analyses were performed using GraphPad.
Identification of givinostat as an inhibitor of HSC activation using cell-based high-throughput screening. Aiming to find potential candidate compounds that can inhibit HSC activation for the treatment of liver fibrosis, the present study established a high-throughput screening assay based on one-step RT-qPCR for detecting α-SMA mRNA expression as a readout for HSC activation, since activated HSCs are characterized by the expression of α-SMA (21). A library of small molecule inhibitors targeting epigenetic enzymes was screened, and givinostat was identified as the most potent compound that inhibited TGF-β1-induced α-SMA expression (Fig. 1A). Givinostat, a pan-HDACi that belongs to the hydroxamic acids family with HDAC type I and type II inhibitory activity, has been used in phase I/II clinical trials for the treatment of Duchenne muscular dystrophy. To validate the inhibitory effect of givinostat on HSC activation, the mRNA and protein expression levels of Col1α1 and α-SMA after givinostat treatment were examined. Givinostat dose-dependently inhibited Col1α1 and α-SMA mRNA expression in the presence of TGF-β1 (Fig. 1B). Consistently, western blotting also confirmed that givinostat inhibited Col1α1 and α-SMA protein expression levels in a dose-dependent manner (Fig. 1C). Based on these data, givinostat was identified as a potent inhibitor of HSC activation in vitro.

Table I. RT-qPCR primer sequences used in the present study.

| Gene          | Primer sequences (5’→3’)                                                                 |
|---------------|-------------------------------------------------------------------------------------------|
| Acta2 (α-SMA) | **F**: GCTGAAGTATCCCGTATAGAACAGC <br> **R**: GCTCTAAACATAATCTGGGTC                        |
| Collα1        | **F**: TCACAGGCGCGAAGATAAAAGCT <br> **R**: CCCCAAGTTCGGTGGAAGCA                           |
| GAPDH         | **F**: GTGTAAGGCTGTTGTTGGAACGGA <br> **R**: CCAAAGTTGTCTAGGATGACTG                         |
| Sult3a1       | **F**: TATTGGAGCTATGGGAGAAG <br> **R**: GTGATGCTATTTGATGGAGAT                             |
| Upk3b         | **F**: CACTTGCTAGTGGGCTTT <br> **R**: GGTAGTGGGATGACTG                                      |
| Adgrd1        | **F**: GTCTGCTCTGTGGGA <br> **R**: TAAGGACCCGAGGTTGA                                       |
| Snai3         | **F**: GTTCCTCAACTACGGGGAAC <br> **R**: CGTAAAGGGCTACCTGGGATT                             |
| Msln          | **F**: CACCGCGAGGAACTTGAAG <br> **R**: CTCGGTGAGGATGACTCAAC                               |
| Dmkn          | **F**: AGCTGACCACTTCT <br> **R**: GCCAGTTGGAAGTGGTACC                                     |
| Upk1b         | **F**: AACAGGAAAATTCCCTTGGG <br> **R**: AAAAGTGCAGGTTGTGCT                                |
| Cebpe         | **F**: ATTCGGCTATACCTTCCACAC <br> **R**: GTAGCCTCGACTG                                     |
| Eif4ebp3      | **F**: GTCCACGAGTGGCCCAATTC <br> **R**: GGGGTAGTGGGCTATAGGTTG                             |
| Slc2a5        | **F**: TCTCAATTGAGTCAACG <br> **R**: GCCGTAAGGAACTGCAAC                                   |
| Ntrk1         | **F**: GCCAACCATCGTGAGAGT <br> **R**: CCAACGCATGGGACGAG                                   |

Table I. Continued.

| Gene          | Primer sequences (5’→3’)                                                                 |
|---------------|-------------------------------------------------------------------------------------------|
| MSLN          | **F**: CAGAGGAGGCTCAAGAGAGCTA <br> **R**: GCCCCACAGGACCCAC                                 |
| DMKN          | **F**: CCAAGGAGGACGAGACGGAG <br> **R**: CCAAGTGTTTTCCAGAGCAT                                |
| UPK1B         | **F**: GAACCTCTCAACCTGGGAGG <br> **R**: TGGATCAGGAAACCCAAA                                    |
| CEBPE         | **F**: CTTGGAGTGGAGGAGG <br> **R**: GTGCTGGGAGGAGGAGG                                      |
| EIF4EBP3      | **F**: CCACTAGCTGGCCGAGTTCC <br> **R**: GGTAGTGGGAGTACG                                      |
| SLCA25        | **F**: CAAGAAAGTTGATGATGAGGTG <br> **R**: CAAGAAGAGTTGATGATGAGGTG                          |
| NTRK1         | **F**: CCATCTCTGCAGACTGACC <br> **R**: GCCAAGGAGGAGGAGG                                     |

F, forward; R, reverse; HSC, hepatic stellate cell; RT-qPCR, reverse transcription-quantitative PCR; α-SMA, α-smooth muscle actin; Collα1, collagen type I; Dmkn, dermokine; Msln, mesothelin; Upk3b, uroplakin-3b.

Prism 7.0 statistical software (GraphPad Software, Inc.). P<0.05 was considered to indicate a statistically significant difference.

**Results**

Identification of givinostat as an inhibitor of HSC activation using cell-based high-throughput screening. Aiming to find potential candidate compounds that can inhibit HSC activation for the treatment of liver fibrosis, the present study established a high-throughput screening assay based on one-step RT-qPCR for detecting α-SMA mRNA expression as a readout for HSC activation, since activated HSCs are characterized by the expression of α-SMA (21). A library of small molecule inhibitors targeting epigenetic enzymes was screened, and givinostat was identified as the most potent compound that inhibited TGF-β1-induced α-SMA expression (Fig. 1A). Givinostat, a pan-HDACi that belongs to the hydroxamic acids family with HDAC type I and type II inhibitory activity, has been used in phase I/II clinical trials for the treatment of Duchenne muscular dystrophy. To validate the inhibitory effect of givinostat on HSC activation, the mRNA and protein expression levels of Collα1 and α-SMA after givinostat treatment were examined. Givinostat dose-dependently inhibited Collα1 and α-SMA mRNA expression in the presence of TGF-β1 (Fig. 1B). Consistently, western blotting also confirmed that givinostat inhibited Collα1 and α-SMA protein expression levels in a dose-dependent manner (Fig. 1C). Based on these data, givinostat was identified as a potent inhibitor of HSC activation in vitro.
Givinostat treatment alleviates chronic liver injury and fibrosis in mice treated with CCl4. Whether givinostat could inhibit HSC activation in vivo and alleviate liver fibrosis, which has not been extensively studied, was further examined...
in the present study, which investigated its potential role in the treatment of chronic liver fibrotic diseases. A widely used mouse liver fibrosis model with i.p. repeated injection of CCl4 was used to evaluate the efficacy of givinostat for the treatment of hepatic injury and liver fibrosis (29). C57BL/6J mice were i.p. injected with CCl4 for 2 weeks; at that time there was already mild liver fibrosis (30). Mice were then treated with givinostat for the following 6 weeks after CCl4 treatment (Fig. 2A). As shown in Fig. 2B, persistent i.p. injection of CCl4 for 8 weeks caused hepatocyte steatosis (Fig. 2B middle panel), varying degrees of central venous wall thickening and collagen deposition (Fig. 2C middle panel), which are consistent with previous reports (31,32). The results indicated that givinostat treatment markedly ameliorated the extent of hepatocyte steatosis (Fig. 2B right panel). Since the deposition of ECM (primarily collagens) is the major characteristic
of liver fibrosis, the collagen contents were evaluated based on Sirius Red staining. Images (n=6–8 images per group) showed that hepatic lobules maintained a normal physiological structure in the liver of control mice, while higher collagen accumulation was observed in the liver of CCl4-treated mice (P<0.0001; Fig. 2C). In the givinostat treatment group, the
hepatic structure was improved and collagen was decreased (P<0.0001; Fig. 2C). There was a significant increase in the levels of serum ALT (n=6; 469±41.65; P<0.0001) and AST (n=6; 536.8±95.53; P<0.01) in CCl4-treated mice, whereas treatment with givinostat significantly decreased these serum markers of liver injury, ALT (n=6; 137.7±71.72; P<0.01) and AST (n=6; 156.5±71.44; P<0.01) (Fig. 2D). Furthermore, no systemic toxicity was observed at the dose of givinostat used in this experiment (data not shown). Thus, givinostat treatment significantly alleviated liver fibrosis and injury in vivo.

**Givinostat inhibits HSC activation in mice with CCl4-induced liver fibrosis.** Quiescent HSCs are activated on account of liver injury and considered to be the primary source of ECM yielding during hepatic fibrosis. Since givinostat significantly inhibited HSC activation in vitro and alleviated liver fibrosis in vivo, the present study assessed whether it inhibited HSC activation in vivo. α-SMA and Col1α1 are the most abundant ECM proteins in liver tissue, and are markers of HSC activation (33,34). Thus, α-SMA or Col1α1-positive cells were detected by morphometric quantification to evaluate the accumulation of activated HSCs in mouse liver tissues. Immunohistochemical staining for α-SMA (n=6; 65.800±4.861; P<0.01) or Col1α1 (n=6; 8.215±1.069; P<0.0001) showed that positively stained brown-colored cells were notably increased in the liver tissues of mice treated with CCl4 (Fig. 3A middle panel). By contrast, immunohistochemical staining for α-SMA (n=6; 12.886±5.603; P<0.01) or Col1α1 (n=6; 3.140±859.9; P<0.01) in the liver tissue of givinostat-treated mice was much weaker compared with that of solvent-treated mice, almost at a level similar to that of the normal control group (Fig. 3A right panel). RT-qPCR analysis confirmed that the increase in mRNA expression of α-SMA and Coll1α1 in the liver tissues of CCl4-challenged mice was significantly reduced by givinostat treatment (P<0.01; Fig. 3B). Consistently, western blot analysis further confirmed that the protein expression levels of α-SMA and Col1α1 in mouse liver tissues were increased in the CC14-challenged group, and were reduced by givinostat treatment (P<0.05; Fig. 3C). Taken together, these results demonstrated that givinostat alleviated liver fibrosis and inhibited HSC activation in vivo.

**Identification of crucial genes for HSC activation that are regulated by givinostat via transcriptomic analysis.** To explore the mechanism underlying the improvement of hepatic fibrosis by givinostat treatment in CC14-challenged mice, RNA-seq analysis was performed to compare the gene expression profile of liver tissues from CC14-challenged mice with or without givinostat treatment. Differential gene expression analysis identified genes upregulated or downregulated in givinostat-treated group compared with their expression in the solvent group in CC14-challenged mice. The most significantly regulated genes by givinostat treatment are shown in Fig. 4A. RT-qPCR analysis confirmed that givinostat treatment inhibited or upregulated the mRNA expression of these genes in liver tissues (Fig. 4B), which was consistent with the RNA-seq results.

The present study next examined whether givinostat regulated the transcription of these genes in vitro in HSCs. The mRNA expression of these genes was analyzed via RT-qPCR in HSC LX-2 cells stimulated by TGF-β1 in the absence or presence of givinostat treatment. The mRNA expression levels of Msln, Dmkn and Upk3b were also reduced in HSC LX-2 cells (Fig. 4C), which was consistent with the findings in liver tissues of mice treated with givinostat. As for the genes that were most notably upregulated by givinostat treatment in vitro, their mRNA expression was increased in vitro to a much lower extent (Fig. 4C). As shown in Fig. 4D, givinostat dose-dependently inhibited Msln, Dmkn and Upk3b mRNA expression in the presence of TGF-β1. These results confirmed that givinostat inhibited Msln, Dmkn and Upk3b gene expression in vivo and in vitro.

To further validate whether these givinostat-regulated genes play crucial roles during HSC activation, the effect of Msln, Dmkn and Upk3b depletion on TGF-β1-induced HSC activation was examined. RT-qPCR analysis confirmed the knockdown efficiency of Msln, Dmkn and Upk3b by siRNA, which resulted in >70% reduction in mRNA levels in HSC LX-2 cells compared with that of the siControl group (P<0.0001; Fig. 5A). RT-qPCR analysis showed that the knockdown of Msln, Dmkn and Upk3b inhibited the mRNA levels of α-SMA and Coll1α1 in HSC-LX2 cells stimulated with TGF-β1 compared with the siControl + TGF-β1 group (P<0.0001; Fig. 5B and C). These results demonstrated that givinostat inhibited Msln, Dmkn and Upk3b expression, and these genes were shown to be crucial for HSC activation.

**Discussion**

As a chronic hepatic disease with limited treatment options, liver fibrosis affects millions of individuals worldwide. Fibroblasts and myofibroblasts, which are primarily responsible for the synthesis of ECM proteins, have been identified as pivotal fibrotic effectors in multiple organs (35). HSCs, following activation and transformation to a myofibroblast phenotype, as marked by expression of α-SMA, are the dominating cell type that produce ECM proteins during liver fibrogenesis (36,37). Thus, the inactivation of HSCs is currently regarded as a potential treatment strategy for the treatment of hepatic fibrosis (21).

The present study established high-throughput screening assays to identify compounds that inhibited HSC activation. In view of the potential regulatory role of epigenetic mechanisms in modulating HSC function and reprogramming, an epigenetic inhibitor library was screened. The HDACi givinostat, which has been used in phase I/II clinical trials for Duchenne muscular dystrophy (15), was identified and validated as a potent inhibitor of HSC activation in vitro and in vivo. In the screening assay, givinostat was the most potent inhibitor of HSC activation, and it was more potent than other HDACi drugs. It was also observed that givinostat treatment alleviated liver fibrosis and liver injury in CC14-injected mice, which had shown mild liver fibrosis. The liver organization architecture and function in mice with hepatic fibrosis were markedly improved after givinostat treatment. A previous study reported that givinostat can inhibit the proliferation and induce the apoptosis of HSC cells, thereby inhibiting liver fibrosis in mice subjected to a high-fat diet combined with intraperitoneal injection of CC14 (38). However, ingestion of an obesogenic diet in that model could lead to the development
**Figure 4. Identification of genes for HSC activation that are regulated by givinostat in vivo and in vitro via transcriptomic analysis.** (A) RNA-sequencing analysis of liver tissue extracted from CCl4 (n=4) and CCl4 + givinostat-treated mice (n=4). Diagram of the top genes with the most significant changes with adjusted P<0.05 and absolute value of log2 fold-change >1.5. (B) RT-qPCR analysis was performed to validate the expression of the genes exhibiting major changes following givinostat treatment in vivo in liver tissues (n=4). (C) RT-qPCR analysis was performed to validate the expression of the genes displayed in panel (B) in HSC LX-2 cells treated with givinostat in the presence of TGF-β1. (D) RT-qPCR analysis of the expression of three genes in HSC LX-2 cells treated with different concentrations of givinostat. The results are normalized to β-actin and error bars indicate the SD of three independent experiments. Data are expressed as the mean ± SD.* P<0.05, ** P<0.01, *** P<0.001, **** P<0.0001 vs. the TGF-β1 group. HSC, hepatic stellate cell; RT-qPCR, reverse transcription-quantitative PCR; CCl4, carbon tetrachloride; Dmkn, dermokine; Msln, mesothelin; Upk3b, uroplakin-3b.

### A) Gene Expression Analysis

| Gene    | Log2Fold-change | Inhibited by givinostat | Upregulated by givinostat |
|---------|-----------------|-------------------------|---------------------------|
| **Msf** | -6.701855899   |                         |                           |
| **Dmkn**| -6.10974243    |                         |                           |
| **Upk3b**| -5.201986187  |                         |                           |
| **Upk1b**| -4.35607744    |                         |                           |
| **Adgr1**| -2.966815175   |                         |                           |
| **Cebpe**| -2.110823927   |                         |                           |
| **Eil4ebp3**| 2.140357193   |                         |                           |
| **Slc2a5**| 2.150984153    |                         |                           |
| **Sult3a1**| 2.458027153    |                         |                           |
| **E030018b13rik**| 2.835231844 |                         |                           |
| **Sna3** | 2.874938114   |                         |                           |
| **Ntrk1**| 3.183048653    |                         |                           |

### B) Liver Tissue RT-qPCR Analysis

- **Msf**, **Dmkn**, **Upk3b**, **Upk1b**, **Adgr1**, **Cebpe**

### C) HSC-LX2 Cell RT-qPCR Analysis

- **Msf**, **Dmkn**, **Upk3b**, **Upk1b**, **Adgr1**, **Cebpe**

### D) HSC-LX2 Cell RT-qPCR Analysis (Different Concentrations of Givinostat)

- **Msf**
- **Dmkn**
- **Upk3b**

**Legend:**
- +: Treatment with givinostat
- -: Control treatment
- CCl4: Carbon tetrachloride
- RT-qPCR: Reverse transcription-quantitative PCR
- TGF-β1: Transforming growth factor-beta 1
of steatosis, which might result in steatohepatitis and progressive fibrosis (16); thus, the alleviation of liver fibrosis might also be derived from the reduction of steatosis by givinostat.

To evaluate the effects of givinostat on a CCl4-induced liver fibrosis model, the present study showed a reduction in HSC activation and alleviation of liver fibrosis by givinostat in vivo, and also ruled out the possibility that the alleviation of fibrosis by givinostat was due to reduced steatosis. It has also been reported that HDAC9 knockdown can inhibit HSC activation and decrease fibrogenic gene expression in HSC LX-2 cells (39). Moreover, in agreement with the anti-fibrotic effects of givinostat shown in the present study, givinostat was reported to decrease endothelial-to-mesenchymal transition and reduce cardiac fibrosis, leading to improved heart performance and protection of blood vessels from apoptosis in a mouse model of acute myocardial injury (19). Moreover, single or repeated oral administration of givinostat in humans has been found to be safe, as shown in a previous clinical trial (29). In the present mouse model, no systemic toxicity was observed at the dose of givinostat used in this experiment. Thus, the present study suggested that givinostat inhibited hepatic fibrosis and HSC activation in vivo, and raised the possibility that an existing drug can be repurposed as a new treatment of hepatic fibrosis.

Transcriptomic analysis revealed the most significantly regulated genes in the givinostat treatment group in comparison with the solvent group, among which, Dmkn, Msln and Upk3b were validated in vitro in HSC LX-2 cells as crucial genes regulating HSC activation. When Msln, Dmkn or Upk3b expression was knocked down, the increased mRNA expression of α-SMA and Col1α1 in response to TGF-β1 stimulation was significantly reduced in HSC LX-2 cells, suggesting that these three genes may play crucial roles in the activation of HSCs. To the best of our knowledge, the role of Msln, Dmkn and Upk3b in HSC activation was reported for the first time in the present study. Moreover, givinostat treatment significantly reduced the mRNA expression of Dmkn, Msln
Upk3b in both a mouse model and HSC-LX2 cells. Certain genes that were significantly affected by givinostat treatment in vivo were not affected in vitro in HSC LX-2 cells, which may be unrelated to HSC activation or could be the result of other cell types in the liver, such as endothelial, Kupffer and bile-duct cells (40,41). Thus, the identification of givinostat as an inhibitor of HSC activation and its use as a chemical probe led to the identification of novel regulators of HSC activation.

In summary, the present study established a high-throughput cell-based assay for the identification of a compound targeting HSC activation, and identified givinostat as a potent inhibitor of HSC activation in vitro and in vivo. Novel regulators of HSC activation were identified using givinostat as a probe, and these findings illustrated the efficacy of an epigenetic strategy that targets HSC activation for the treatment of hepatic fibrosis.

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Availability of data and materials

The datasets generated and/or analyzed during the current study are available in the GEO repository, https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE161981. The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors' contributions

HMH, YJL, LPL, LY and JJP performed the immunofluorescence staining, western blotting, siRNA transfection and mouse liver fibrosis experiments, analyzed the corresponding data and wrote the manuscript. XRZ, SJF and JH contributed to manuscript writing and modification and analyzed the RNA-seq data. GML, CL, CCS and YYZ conceived and supervised the project, and revised the manuscript. The present article was conducted in accordance with the ARRIVE guideline checklist. The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. HMH, XRZ and LPL confirm the authenticity of all the raw data. All authors read and approved the final manuscript.

Ethics approval and consent to participate

Animal care was carried out according to the guidelines of the Principles of Laboratory Animal Care, and the protocol was approved by the Institute Animal Care and Use Committee at the Shanghai Institute of Materia Medica (approval no. 2018-12-LC-11; Shanghai, China).

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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