Inhibition of ferroptosis and iron accumulation alleviates pulmonary fibrosis in a bleomycin model

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ABSTRACT

Idiopathic pulmonary fibrosis (IPF) is a chronic progressive disease characterized by excessive proliferation of fibroblasts and excess accumulation of extracellular matrix (ECM). Ferroptosis is a novel form of cell death characterized by the lethal accumulation of iron and lipid peroxidation, which is associated with many diseases. Our study addressed the potential role played by ferroptosis and iron accumulation in the progression of pulmonary fibrosis. We found that the inducers of pulmonary fibrosis and injury, namely, bleomycin (BLM) and lipopolysaccharide (LPS), induced ferroptosis of lung epithelial cells. Both the ferroptosis inhibitor liproxstatin-1 (Lip-1) and the iron chelator deferoxamine (DFO) alleviated the symptoms of pulmonary fibrosis induced by bleomycin or LPS. TGF-β stimulation upregulated the expression of transferrin receptor protein 1 (TFRC) in the human lung fibroblast cell line (MRC-5) and mouse primary lung fibroblasts, resulting in increased intracellular Fe2+, which promoted the transformation of fibroblasts into myofibroblasts. Mechanistically, TGF-β enhanced the expression and nuclear localization of the transcriptional coactivator tafazzin (TAZ), which combined with the transcription factor TEA domain protein (TEAD)-4 to promote the transcription of TFRC. In addition, elevated Fe2+ failed to induce the ferroptosis of fibroblasts, which might be related to the regulation of iron export and lipid metabolism. Finally, we specifically knocked out TFRC expression in fibroblasts in mice, and compared with those in the control mice, the symptoms of pulmonary fibrosis were reduced in the knockout mice after bleomycin induction. Collectively, these findings suggest the therapeutic potential of ferroptosis inhibitors and iron chelators in treating pulmonary fibrosis.

1. Introduction

Idiopathic pulmonary fibrosis (IPF) is a fatal fibrotic lung disease characterized by a progressive decline in lung function, with a median survival time of 2–3 years after diagnosis. Moreover, in the past decades, the prevalence and incidence of this disease have been increasing year by year [1–3]. The pathogenesis of IPF is now thought to be initiated by injuries of the lung alveolar epithelium with dysregulated wound healing processes. Epithelial cell apoptosis and impaired epithelial cell progenitor function are believed to be critical in fibrogenesis [4–6]. The accumulation of scar tissue was associated with fibroblast-to-myofibroblast transformation, fibroproliferation and excessive accumulation of extracellular matrix (ECM) proteins in the distal lung parenchyma, eventually leading to a pathological remodeling of lung architecture and the obliteration of lung tissue [1,7,8]. The most important source of myofibroblasts was the activation and transformation of fibroblasts, including not only the fibroblasts originally residing in the lungs, but also the fibroblasts recruited by inflammatory...

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Factors and chemokines, which occupied the majority of myofibroblasts [9–11]. Although two anti-fibrosis drugs (Nintedanib and Pirfenidone) had been approved for the clinical treatment of IPF, they only delayed pulmonary function decline and failed to stop disease progression [12–14]. Therefore, it is essential to characterize the underlying molecular mechanisms of epithelium injury and fibroblast activation in order to explore new treatments to improve the prognosis and survival rate of patients.

Iron is an indispensable trace element for human beings, which regulates many physiological activities of the human body [15,16]. However, disorders of iron homeostasis could lead to a number of pathological conditions including cancer, hemochromatosis, sickle cell disease, fibrotic lung disease, etc [17–20]. Emerging evidences showed that iron metabolism disorder in the lung was closely related to the occurrence and progression of pulmonary fibrosis. Some studies observed extracellular iron and macrophage hemosiderin accumulation in histological samples from some IPF patients [20–22]. In addition, it had been confirmed that pulmonary fibrosis and functional decline were also related to abnormal iron overload in bleomycin-induced pulmonary fibrosis mouse model, and the similar phenomenon had also been verified in the iron overload model of homeostatic iron regulator (Hfe) gene-deficient mice [20].

Iron overload is also related to a new type of programmed cell death, ferroptosis. Membrane protein TFRC could recognize transferrin and deliver the Fe³⁺-transferrin complex composed of two molecules of Fe³⁺ and one molecule of transferrin into cells as an endosome. The reduction of Fe³⁺ to Fe²⁺ by lysosome-endosome fusion, which occurred through STEAP3 in the plasma membrane, was subsequently released into the cytoplasm and was incorporated into the labile iron pool (LIP). Furthermore, other molecules on the cell membrane could also transport Fe²⁺ into the cells as divalent metal transporter 1 (DMT1) and metal-ion transporter ZRT- and IRT-like protein 14 (ZIP14), while TFRC was relatively more important [23,24]. Besides, the amount of free iron in the cells could be regulated by other pathways. Ferritin is a complex of two basic subunits, the Ferritin heavy chain (FTH) and the Ferritin light chain (FTL1), which could bind and store intracellular free iron and thus reduce the risk of ferroptosis. In addition, the unique iron-removal channel protein called solute carrier family 40 member 1 (SLC40A1) or Ferroportin in the cell membrane could also reduce intracellular Fe²⁺ by transporting it out of cells. These iron regulatory elements could regulate intracellular free Fe²⁺ and affect the ferroptotic sensitivity of cells. Correspondingly, some iron-chelating agents such as DFO and deferoxipron (DFP) were known to significantly reverse ferroptosis by reducing the intracellular iron content [16,25–27].

In this study, we investigate the role of iron overload and ferroptosis in the development of pulmonary fibrosis. Both lipid peroxidation accumulation and iron overload are observed during the progress of bleomycin-induced pulmonary fibrosis. In an early inflammatory phase, bleomycin and LPS directly induce ferroptosis in lung epithelial cells, while the ferroptosis inhibitor displays significant resolution for bleomycin- and LPS-induced cell death of type II alveolar epithelial (ATII) cells and lung injury. In a late fibrotic phase, TGF-β enhances intracellular labile ferrous accumulation by upregulating TFRC via the TAZ-TEAD signaling, thus promoting the fibroblast-to-myofibroblast transition. Fibroblasts also survive in response to iron-induced ferroptotic stress by regulation of the ferroptosis-defense pathway. These findings pinpoint a novel mechanism by which ferroptosis and iron accumulation lead to epithelial injury and fibroblast activation in different phases of the bleomycin-induced pulmonary fibrosis model. This study also reveals the therapeutic potential of ferroptosis-prevention therapies and iron chelator in treating lung fibrosis.

2. Materials and methods

2.1. Cell culture

BEAS-2B and IMR90 cell lines were obtained from the American type culture collection (ATCC, United States), MRC-5 was obtained from Procell Life Science&Technology Co.,Ltd. These cells had been proven to be negative for mycoplasma contamination. All cells including BEAS-2B, MRC-5, IMR90 and PLFs were maintained in RPMI 1640 with 10% FBS (Gibco, United States), 2 mM L-glutamine, 100 U/ml of penicillin, and 100 μg/ml of streptomycin at 37 °C and 5% CO₂.

2.2. Isolation and culture of primary lung fibroblasts (PLFs)

Wild-type male C57 mice (less than 4 weeks) were selected. After the mice were euthanized, the lung tissues were rinsed with PBS and then infiltrated in serum-free medium. Lung tissue was cut into 1–3 mm pieces with sterile ophthalmic tweezers in a dish and added into serum-free medium with 200 U/ml penicillin, 200 μg/ml streptomycin and 1.5 mg/ml type IV collagenase. Lung tissues were incubated at 37 °C for 45 min and shook slightly every 15 min. 10% FBS was added to stop collagenase digestion and the digested lung fragments were passed through a 70 μm filter. Cells were centrifuged at 1500 rpm for 5 min. Red Cell Lysis Buffer (TIANGEN, Cat#RT122-02) was added to the cell pellets and the cells were suspended and incubated at room temperature for 5 min. Cells were then centrifuged at 1500 rpm for 5 min, and the supernatant was discarded. Cells were resuspended in complete medium for counting. 5 × 10⁵–1 × 10⁶ cells were added into each well of the 6-well plate, and the medium was supplemented to 2 ml per well. All procedures were performed on ice unless otherwise stated. Fibroblasts (a group of spindle cells) were obtained by repeated trypsinization and passage.

2.3. Reagents and drugs

Reagents and Drugs used were bleomycin (Selleck, S1214), LPS (Sigma-Aldrich, L4005), recombinant Human TGF-β1 Protein (Peprotech, AF-100-21C-100), Hematoxylin (BASO, BA-4097), Eosin (BASO, BA-4098), Masson’s trichrome Stain Kit (Solarbio, GI340), Hoechst 33342 (Beyotime, C1022), SYTOX™ Green (Thermo Fisher, S34860), Liproxstatin-1 (Selleck, S7699), Ferrostatin-1 (MCE, HY-100579), Deferoxamine (Sigma-Aldrich, Y0001937), Ferric ammonium citrate (FAC) (Sigma-Aldrich, F5879), SB431542 (MCE, HY-10431), Necrostatin-1 (MCE, HY-15760), Cell Counting Kit-8 (CCK-8) (Engreen, EC008).

2.4. Real-time quantitative PCR

Total RNA in cells was collected using the RNA extraction Kit (E.Z.N.A. Total RNA Kit, OMEGA, R6934) according to manufacturer’s instructions. RNA concentration was measured by Nanodrop 2000C (Thermo Scientific). cDNA synthesis was performed using a reverse transcription kit (Prime-Script RT Master Mix, TaKara, RR036A). RT-PCR was performed using SYBR Premix Ex Taq™ (TaKara, DRR081A). The relative expression of each target gene was normalized to GAPDH expression. The following primers were used: TFRC (human), Forward: 5′-ACCATTTGCTATATACCCGGTTC-3′; Reverse: 5′-CAATAGCCAAATGTCAGCACT-3′. TRFC (mouse), Forward: 5′-GGTGAATGATCCTCTGCTGCG-3′; Reverse: 5′-CCCGAGAAGAATTTGCTGAAAGG-3′. α-SMA (human), Forward: 5′-CTTAGAGGCT ATGCCCTGCC-3′; Reverse: 5′-GCTACAGTATAGAAAGGAA-3′. α-S MA (mouse), Forward: 5′-GTTCAGTGTGCCCTGTCTG-3′; Reverse: 5′-AAGGAGATGTTGAAAAGG-3′. COL1A1 (human), Forward: 5′-GAG GCACAGAGGAAGACATC-3′; Reverse: 5′-CAGATACGTCATGCA CAAC-3′. COL1A1 (mouse), Forward: 5′-GTCTTCGTTGCTCTGTTG-3′;
Reverse: 5'-GACGTTGACTCCTCTT-3'. GAPDH (human), Forward: 5'-CAG GTG TTCCT GACGTT-3'; Reverse: 5'-CCA AAT TCG TGT TCA TAC CA-3'. GAPDH (mouse), Forward: 5'-TGACCTCAACTGATGGCTCTACA-3'; Reverse: 5'-CTTCCCATTTGCGGCTTG-3'.

2.5. Western blot and antibodies

The cultured cells were washed with PBS and then Radio-Immunoprecipitation Assay (RIPA) lysis buffer was added. Cell samples were collected with a cell scraper. Lung tissue from mice was weighed and ten times the mass of RIPA lysis buffer was added. Then the lung tissue was homogenized in a pre-cooled tissue homogenizer for 2 × 30 s at 50 Hz. Subsequently, cell or tissue samples were placed in ice, lysed for 10 min and centrifuged at 16 000 g for 15 min, and the supernatant was collected. These protein samples were separated by 10% SDS-PAGE and then were analyzed by Western blot. Primary antibodies used were anti-α-tubulin (Proteintech, 66031-1-lg), anti-GAPDH (HUAbio, R1210-1), anti-α-SMA (abcam, ab7817), anti-COL1A1 (abcam, ab34710), anti-TFRC (abcam, ab214039), anti-Ferritin (abcam, ab75973), anti-SLC40A1 (abcam, ab78066) and anti-4-HNE (abcam, ab48506).

2.6. Migration and invasion assay

Transwell chamber (Millipore, P18PO1250) with or without Matrigel gel (Corning, 354 234) were used for transwell migration or invasion assay. MRC-5 cells were seeded in the upper portion of the transwell chamber in a 24-well plate suspended in 500 µl serum-free medium. Then 500 µl of RPMI 1640 with 10% FBS was added to the lower portion of the transwell chamber and the drug was simultaneously added into medium. After 48 h of incubation, the transwell chamber was washed with PBS. Cells on the upper portion of the chamber were discarded with a cotton swab. The fluorescence intensity was proportional to the concentration of Fe²⁺ in the cells. For mouse lung tissue, it was first cut into 2.5 mm pieces, then ground and extruded, and made into a single cell suspension, and the cells were incubated at 37 °C for 1 h away from light. The FeRhoNox-1 promoter-Luciferase Reporter plasmid (Promega, Glomax-20/20) using a Dual Luciferase Reporter Assay System (Promega, E1910 and E1960) according to manufacturer’s instructions. Renilla luciferase was be used as a reference for normalization of transfection efficiency.

2.7. Measurement of Fe²⁺ in cells

The treated cells were collected by trypsin digestion. After the cells were washed with PBS, 5 µM FeRhoNox-1 (MKBio, MX4558) was added to cell suspension, and the cells were incubated at 37 °C for 1 h away from light. The FeRhoNox-1 probe entering the cell would react with Fe²⁺ to produce an irreversible orange-red fluorescent substance (Ab = 540 nm, FL = 575 nm). After the MRC-5 cells were washed with PBS again, flow cytometry (Fortessa, BD Biosciences) was able to analyze fluorescence in single cell, and the fluorescence intensity was proportional to the concentration of Fe²⁺ in the cells. For mouse lung tissue, it was first cut into 1–3 mm pieces, then ground and extruded, and made into a single cell suspension through a 70 µm filter. To detect Fe²⁺ in the α-SMA⁺ cells from mouse lung tissue, cells were fixed with fixation buffer (Biolegend, 420 801) at 4 °C for 30 min and the special cells were incubated with a α-SMA primary antibody (abcam, ab7817) at 4 °C for 1 h. The fluorescent secondary antibody used is Alexa Fluor 488 Donkey Anti-mouse IgG (Invitrogen, A21202). After washing with PBS, FeRhoNox-1 was incubated in 20 µl of 1× binding buffer. In addition, depending on the probe, intracellular Fe²⁺ can also be reflected by fluorescence microscopy under same condition.

2.8. Measurement of ROS and lipid ROS

After the cells were digested by trypsin and collected, the cells were washed with PBS. Then the cells were resuspended in serum-free medium containing 10 µM 2′,7′-dichlorodihydrofluorescein diacetate (CM-H2DCFDA, Invitrogen, C6827) and incubated at 37 °C for 1 h. Intracellular reactive oxygen species (ROS) could oxidize H2DCFDA to highly fluorescent 2′,7′-dichlorofluorescein (DCF), thus flow cytometry could analyze the ROS content in a single cell. Similarly, C11 BODIPY 581/591 (Invitrogen, D3861) at a concentration of 10 µM could specifically bind to and react with the lipid reactive oxygen species (lipid ROS) in the plasma membrane. As a result, the fluorescence peak of C11 BODIPY 581/591 shifted from ~590 nm to ~510 nm, which could be detected by flow cytometry.

2.9. Measurement of cell death and cell viability

BEAS-2B cells were seeded in 12-well plates (5 × 10⁴ cells/well) and were treated with bleomycin (10 µg/ml) or LPS (5 µg/ml) with or without Fer-1 (5 µM). MRC-5 cells were seeded in 12-well plates (5 × 10⁴ cells/well) and were treated with FAC with or without DFO (50 µM). After that, cell death could be analyzed by dual staining with Hoechst33342 and SYTOX Green followed by microscopy at special node and the percentage of dead cells was counted by ImageJ software. MRC-5 cells were seeded in 96-well plates (1 × 10⁵ cells/well) and were treated with FAC with or without DFO (50 µM) for 72 h. Then cell viability was determined by CCK8 according to manufacturer’s instructions.

2.10. RNA interference

MRC-5 cells were transfectected with siRNA (20 µM) through jetPRIME transfection reagent (Polyplus Transfection, PT-114-15) following the manufacturer’s instructions. The human-specific siRNAs targeting TFRC were designed and synthesized by GenePharma (Shanghai). Sequences of siRNAs were as follows: siRNA1 (Sense: 5′-GGCGGACAAAGUUAGAAGAATT-3'); siRNA2 (Sense: 5′-GCGGGACAGUUUGAGAATT-3'); siRNA3 (Sense: 5′-GGCGGACAAAGUUUGAGAATT-3'); Negative Control (Sense: 5′-UUUCGGAACAGUUCAGUUGUUGAATT-3').

2.11. Lentivirus transfection

MRC-5 cells were seeded in a 6-well plate at a density of 1 × 10⁵ cells/well. Subsequently, 5 µl of 1 × 10⁸ TU lentivirus (ShangHai Gen-eChem, China) and 40 µl of 25 × HitransG A lentivirus infection regent were added to each well. Transfection efficiency was assessed by green fluorescent protein (GFP) detection using fluorescence microscopy, and cells were selected with fluorescence-activated cell sorting. The expression levels of relevant proteins (TFRC) were examined by Western blot.

2.12. Dual luciferase reporter assay

293T cells were seeded in 96-well plates at a density of 1 × 10⁴ cells/well. When the cell density reached 60–80%, the mixture of TFRC Promoter-Luciferase Reporter plasmid (50 ng), the pCMV6-Entry-TEAD4 plasmid (0/50/100/200 ng) and Renilla plasmid (2.5 ng) was co-transfected according to jetPRIME Transfection reagent operation instructions. After 24 h, luciferase activity was measured by luminometer (Promega, Glomax-20/20) using a Dual Luciferase Reporter Assay System (Promega, E1910 and E1960) according to manufacturer’s instructions. Renilla luciferase was be used as a reference for normalization of transfection efficiency.

2.13. Tissue iron content detection

Fresh mouse lung tissue of similar quality was collected. After adding 250 µl extracting solution, lung tissues were homogenized in a pre-cooled tissue homogenizer for 2 × 30 s at 50 Hz. A tissue iron assay kit was used for subsequent extraction following the manufacturer’s instruction (Solarbio, BC4355). Fe²⁺ in tissue could be reduced to Fe²⁺ that could be detected colorimetrically with 2,2’-bipyridine at 520 nm which indicated the iron content in tissue.
2.14. Detection of hydroxyproline in lung

Fresh or frozen lung tissue including the middle and lower lobes of the right lung was selected for examination. According to the manufacturer’s instructions (Nanjing Jiancheng, A030-2-1), the sample was added to the hydrolysate and heated at 100 °C for 20 min. The pH value of the lysate was adjusted to 6.0–6.8 and the lysate was centrifuged at 5000 rpm for 10 min. After subsequent reaction, the supernatant could be used to detect the absorbance value at 550 nm wavelength.

2.15. Immunofluorescence staining in cells

MRC-5 cells or PLFs were seeded in a 35 mm confocal dish (5 × 10^4 cells/dish). After treating with drug for 48 h or 72 h, the cells were washed with PBS and fixed with 4% paraformaldehyde for 10 min, and permeabilized with 0.2% Triton X-100 for 8 min. Then the cells were blocked with 2% goat serum at room temperature for 20 min. Subsequently, the cells were incubated with primary antibody at 4 °C overnight and were incubated with fluorescent secondary antibody at room temperature for 1 h after washing. Confocal microscope could analyze the expression intensity and location of molecules in cells. The concentrations of antibodies used in immunofluorescence staining of cells were anti-α-SMA (abcam, ab7817, 1:300), anti-TAZ (CST, 83669, 1:100), Donkey anti-Mouse IgG Alex Fluor 555 (Invitrogen, A-31570, 1:200) and Donkey anti-Rabbit IgG Alexa Fluor 488 (Invitrogen, A-21206, 1:200). DAPI (Beyotime, C1002, 1:500) was used to label the nucleus. The cellular fluorescent images were acquired by a Nikon Eclipse Ti confocal microscopy (Nikon, Tokyo, Japan).

2.16. H&E staining and Masson’s staining

After the mice were euthanized, the whole lung was taken and fixed overnight in 4% paraformaldehyde. Then gradient dehydration was carried out according to standard procedures and the tissue was paraffin embedded. Next, the tissue was sliced into 3 μm slices. After routine dewaxing and hydration, H&E staining was performed with hematoxylin (BASO, BA-4097) for 7 min and eosin (BASO, BA-4098) for 7 min. Masson’s staining was performed according to the Masson’s trichrome staining Kit (Solarbio, G1340). The blue collagen tissue was visible under a light microscope. H&E staining scanning was performed with PANNORAMIC 250 Flash III (3DHISTECH Ltd.), Masson’s staining image was collected with an OLYMPUS IX73 microscope (Olympus, Tokyo, Japan).

2.17. Immunohistochemistry (IHC) and immunofluorescence staining for lung tissue

The drug-induced fibrotic lung tissues of mice were embedded in paraffin and sectioned in 3 μm sections, which can be used for immunohistochemistry and immunofluorescence staining. After routine dewaxing and hydration, the antigen retrieval step was performed in citrate buffer (pH6.0) (ZSGB-BIO, ZL19065) at ~115 °C for 2 min which could enhance the binding of primary antibody to a specific epitope. Then the section was blocked with goat serum at room temperature for 1 h. Subsequently, the section was incubated with primary antibody at 4 °C overnight. After washing, the section was incubated with HRP-labeled Goat Anti-Mouse/Rabbit IgG for 1 h at room temperature and Streptavidin/HRP for 20 min at room temperature. Finally, the DAB chromogenic kit (ZSGB-BIO, ZL9019) was used for chromogenic reaction. The Immunohistochemistry kit (ZSGB-BIO, SP-9000) could perform specific histochemical staining. The concentrations of primary antibodies were anti-TGFβ1 (abcam, ab215715, 1:500), anti-α-SMA (abcam, ab7817, 1:300), anti-FAPs (abcam, ab207178, 1:200). Different from IHC, the section would be incubated with fluorescent secondary antibody at room temperature for 1 h after washing the primary antibody in immunofluorescence staining. The concentrations of antibodies were anti-α-SMA (abcam, ab7817, 1:3000), anti-TFR (abcam, ab214039, 1:200), Donkey anti-Mouse IgG Alex Fluor 488 (Invitrogen, A-21202, 1:200) and Donkey anti-Rabbit IgG Alex Fluor 555(Invitrogen, A-31572, 1:200). DAPI (Beyotime, C1002, 1:500) was used to label the nucleus. The images were collected with a Nikon Eclipse Ni-U microscope (Nikon, Tokyo, Japan).

2.18. Multicolor immunofluorescence staining for lung tissue

Similarly, the lung tissues of mice were embedded in paraffin and sectioned in 3 μm sections, following gradient dehydration and hydration. Then heat-induced antigen retrieval was performed using citrate (pH = 6) or Tris-EDTA (pH = 9) buffer. After that, the section was blocked with 5% goat serum and was incubated with primary antibody at 37 °C for 1 h, Opal Polymer HRP Mouse + Rabbit at room temperature for 10 min, and TSA Amplification at room temperature for 10 min. The Opal 7-color Manual IHC Kit (Akyoa Biosciences, NEL811001KT) was purchased for the assay. Nuclei were stained with DAPI. The sample was washed by TBST for 2 min × 3 per interval. The primary antibodies used were anti-TGFβ1 (abcam, ab215715, 1:500), anti-α-SMA (abcam, ab7817, 1:3000), anti-FAPs (abcam, ab207178, 1:200), anti-Sftpc (abcam, ab90716, 1:200) and anti-FSP1 (abcam, ab197896, 1:2000). Images were collected by AKOYA BIOSCIENCES Vectra Polaris and analyzed by InForm 2.4.9 software or HALO™ software (Akyoa Biosciences).

2.19. Animal experiments

For earlier experiment, wild-type male C57 mice aged 10 weeks and weighing 25 g were selected as experimental subjects, which were purchased from SPF (Beijing) Biotechnology Co.,Ltd. In the first experiment, the mice were randomly divided into two groups including control (Ctrl) and bleomycin. In the second experiment, the mice were randomly divided into four groups: control, bleomycin, bleomycin + Lip-1 and bleomycin + DFO. Finally, the mice were randomly divided into three groups: control, LPS, LPS + Lip-1. The mice were intratracheally injected with normal saline (total volume 50 μl) or bleomycin (total volume 50 μl, 5 mg/kg) or LPS (total volume 50 μl, 5 mg/kg). For the inhibitor group, mice were intraperitoneally injected with Lip-1 (total volume 150 μl, 15 mg/kg) three times a week or DFO (total volume 200 μl, 30 mg/kg) daily. When Lip-1 was injected from day 0, lung tissue samples from the mice were collected at 4, 7, 14 and 21 days. When Lip-1 was injected from day 10, samples were collected at 10, 14, 18 and 21 days.

C57BL/6J-FSP1-CreERT transgenic mice were purchased from Shanghai Model Organisms Center. C57BL/6J-TFRC<sup>flox/flox</sup> transgenic mice were purchased from Cyagen Biosciences. Crossing the TFRC<sup>flox/flox</sup> mouse with the FSP1-CreERT transgenic mice could generate FSP1-CreERT; TFRC<sup>flox/flox</sup> mice. And Tamoxifen (MCE, HY-13757A) could induce the knockout of TFRC in FSP1<sup>−/−</sup> cells in FSP1-CreERT; TFRC<sup>flox/flox</sup> mice, described as TFRC<sup>−/−</sup> mice, according to the manufacturer’s instructions. Bleomycin induced pulmonary fibrosis in TFRC<sup>−/−</sup> mice and TFRC<sup>flox/flox</sup> mice under similar condition.

2.20. Statistical analyses

All statistics were presented with mean ± SD, which were performed through Prism 7.0 GraphPad Software. Unpaired Student’s t-test was used for comparison between two groups and one-way ANOVA was used for comparison between multiple groups, followed by Tukey test. P < 0.05 was considered as statistically significant.
Fig. 1. The accumulation of lipid peroxides and iron overload in lungs from bleomycin-induced pulmonary fibrosis mice. (a) Upper, lungs of mice from control (Ctrl) group and bleomycin (BLM) group on Day 21 after bleomycin administration. Bottom, representative images of H&E staining, Masson staining, and IHC staining for α-SMA, FAPα and TGF-β of lung tissue from control group and bleomycin-treated group at different days. Scale bars, 100 μm. (b) Scatter plot showing the percentage of the positive area of Masson staining and IHC staining in lung tissue. n = 3–4 mice per group. Statistical analysis was performed using one-way ANOVA followed by multiple comparisons. *P < 0.05, **P < 0.01, ***P < 0.001, ****P < 0.0001. (c) Left, representative images of multicolor immunofluorescence for DAPI (blue), α-SMA (green) and TGF-β (orange) in lung tissue of mice on Day 14 after bleomycin administration. Scale bars, 50 μm. Right, scatter plot showing the percentage of α-SMA+ and TGF-β+ cells, mean fluorescence intensity (MFI) of α-SMA and the positive area of TGF-β in lung tissue of mice on Day 14 after bleomycin administration. Each dot represents a microscopic image, n = 15 images per group from 3 mice. Statistical analysis was performed using the Student’s t-test. *P < 0.05, ****P < 0.0001. (d) Left, immunoblotting analysis of 4-HNE in lung tissues of mice on Day 7 and Day 14 after bleomycin administration. Right, quantitative analysis of 4-HNE from three mice. Statistical analysis was performed using the Student’s t-test. *P < 0.05, **P < 0.01. (e) Intracellular Fe2+ levels in lung tissue of mice were assessed by flow cytometry using FeRhoNox-1 on Day 7 and Day 14 after bleomycin administration. Upper, representative flow cytometry result for Fe2+. Bottom, scatter plot showing the results from three mice. Statistical analysis was performed using one-way ANOVA followed by multiple comparisons. **P < 0.01, ****P < 0.0001. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)
3. Results

3.1. Lipid peroxidation accumulation and iron overload during bleomycin-induced pulmonary fibrosis

To explore the role and mechanism of iron and ferroptosis in IPF, we established a pulmonary fibrosis model by treating mice with bleomycin, which is commonly applied to induce pulmonary fibrosis in rodents [8, 28, 29]. H&E staining showed that the alveolar structure was destroyed, the lung interstitium was thickened, and the lung texture was abnormal (Fig. 1a). Bleomycin also gradually increased the deposition of collagen, which was measured by Masson’s staining (Fig. 1a and b). After bleomycin administration, a significant increase in the expression of two widely accepted markers of fibroblasts, α-smooth muscle actin (caption on next page)
(α-SMA) and fibroblast activation protein-α (FAPα), was observed within 7 days, with peak levels identified at 14 days (Fig. 1a and b).

TGF-β expression was also increased in bleomycin-induced lung fibrosis, occurring 4 days after bleomycin treatment, which was earlier than the upregulation of fibroblast marker expression (Fig. 1a and b). Fourteen days after the administration of bleomycin, multicolor immunofluorescence analyses showed that the expression of TGF-β and the populations of α-SMA fibroblasts were increased, consistent with the immunohistochemistry results (Fig. 1c). These data confirmed the establishment of pulmonary fibrosis induced with bleomycin, and a suitable time point for assessing lung fibrosis was demonstrated.

We then measured a putative biomarker of lipid peroxidation, 4-hydroxynonenal (4-HNE) [30, 31], in the lung tissues of bleomycin-induced pulmonary fibrosis mice at different time points. Western blot analysis showed that the content of 4-HNE in the bleomycin-treated lung tissue increased compared with that in the control group tissue at 7, 14 and 21 days, suggesting the accumulation of lipid peroxides in lung tissues during bleomycin-induced pulmonary fibrosis (Fig. 1d, Supplementary Fig. 1a). In addition to elevated lipid peroxide levels, flow cytometry demonstrated the accumulation of ferrous iron (Fe^{2+}) in cells in bleomycin-treated lung tissue at both 7 and 14 days, as indicated by Fe^{2+}-specific fluorescent probes (Fig. 1e, Supplementary Figs. 1b and 1c). These data confirmed the accumulation of lipid peroxides and iron overload in bleomycin-induced pulmonary fibrosis. Notably, the production of lipid ROS and accumulation of ferrous iron were much lower in the lungs 14 days after bleomycin administration than 7 days after bleomycin administration. These data suggest that lipid peroxide production and iron overload tend to occur within a relatively short period after bleomycin administration, which may be subsequently partially relieved.

3.2. Bleomycin and LPS induce lipid ROS accumulation and ferroptosis in alveolar epithelial cells

Chronic injury of epithelial cells, especially ATII cells, is thought to be a key early driver of the aberrant wound-healing response and pathogenesis of IPF [32, 33]. Emerging data indicate that oxidative damage might trigger the apoptosis of ATII cells, contributing to TGF-β-induced pulmonary fibrosis [34, 35]. Since we demonstrated that lipid peroxides and iron accumulated in bleomycin-induced pulmonary fibrosis, we investigated whether ferroptosis was involved in bleomycin-induced lung epithelial cell damage. Human lung epithelial BEAS-2B cells were exposed to bleomycin and then evaluated for expression of ferroptosis markers. As shown in Fig. 2a and b, bleomycin significantly enhanced the production of ROS and lipid ROS in BEAS-2B cells, and the effects on lipid ROS were abolished by the lipid peroxide-trapping agent ferrostatin-1 (Fer-1). Moreover, intracellular ferrous iron was detected by FerroNox-1 probe staining. The accumulation of ferrous iron in bleomycin-stimulated BEAS-2B cells was inhibited by both Fer-1 and DFO (Fig. 2c). Notably, treatment with Fer-1 partially prevented the cell death triggered by bleomycin (Fig. 2d), suggesting that ferroptosis was involved in bleomycin-induced cell death. We then used LPS, another commonly used lung injury and pulmonary fibrosis inducer, to repeat the aforementioned experiment, and the results were basically consistent (Fig. 2e–h). Necroptosis inhibitor Necrostatin-1 (Necro-1) could also diminish cell death triggered by bleomycin or LPS, demonstrating that bleomycin- or LPS-induced cell death may include both ferroptosis and necroptosis (Supplementary Fig. 1d). Previous articles have proved that bleomycin could induce apoptosis of ATII cells, indicating that multiple cell death modes may contribute to injury of alveolar epithelium in bleomycin-induced pulmonary fibrosis [32, 36]. Collectively, these results indicate that ferroptotic cell death of lung epithelial cells is involved in the lung injury and destruction of alveolar epithelium induced by both bleomycin and LPS.

3.3. Ferroptosis inhibitor and iron chelator induce significant resolution of bleomycin- and LPS-induced lung injury and fibrosis in mouse models

To examine whether blocking ferroptosis protects lung tissue from injury and fibrosis, we treated mice with the potent ferroptosis inhibitor Lip-1 three times per week following the administration of bleomycin. Consistent with our hypothesis, a decrease in the number of surfactant protein C (SPC, encoded by Sftpc)-positive type II alveolar epithelial cells was observed from 4 days through 14 days (Fig. 3a). Lip-1 treatment rescued SPC-positive type II alveolar epithelial cells after bleomycin-induced acute lung injury, indicating recovery of the alveolar epithelium (Fig. 3a). H&E staining of lung sections showed that the fibrotic symptoms characterized by alveolar septal thickening and pulmonary consolidation were relieved by Lip-1 treatment (Fig. 3b). In addition, we found that the levels of both hydroxyproline and lipid ROS in bleomycin-induced pulmonary fibrosis were significantly reduced in the Lip-1-treated group (Fig. 3c and d). Masson’s trichrome staining revealed a substantial resolution in collagen deposition in the lung sections of the Lip-1-treated mice compared with that in the bleomycin-treated mice (Fig. 3e and f). Notably, the time-dependent increases in the levels of α-SMA, FAPα and TGF-β were reduced in Lip-1-treated mice compared with bleomycin-treated mice, reflecting the decreased severity of the pulmonary fibrosis (Fig. 3e and f).

To determine whether Lip-1 modulates pulmonary fibrosis in the fibrotic stage, we intraperitoneally injected Lip-1 on the 10th day after bleomycin treatment. By H&E staining, we found that Lip-1 mitigated bleomycin-induced pulmonary fibrosis in the fibrotic phase (Fig. 4a). The elevated level of hydroxyproline in the lungs of the bleomycin-
Lip-1 inhibited collagen deposition and α-SMA+ fibroblasts in mice lung tissue from control group, bleomycin group and bleomycin + Lip-1 group at different days. Left, representative images of multicolor immunofluorescence for DAPI (blue), α-SMA (green) and SPG (yellow) in lung tissue of mice at different days after bleomycin administration. Scale bars, 50 μm. Right, scatter plot showing the percentage of SPG+ mouse fibroblast cells in lung tissue from control group, bleomycin group and bleomycin + Lip-1 group at different days. Each dot represents a microscopic image, n = 16–34 images per group from 4 mice. Statistical analysis was performed using one-way ANOVA followed by multiple comparisons. *P < 0.01, **P < 0.001. (b) Histopathological study of lung tissue in control group, bleomycin group and bleomycin + Lip-1 group at different days was performed by H&E staining. Representative photographs of lungs on Day 14 were showed. Scale bars, 1 mm. (c) Hydroxyproline (HYP) content from control group, bleomycin group and bleomycin + Lip-1 group at different days. n = 4 mice per group. Statistical analysis was performed using one-way ANOVA followed by multiple comparisons. *P < 0.05, **P < 0.001, ****P < 0.0001. (d) Immunoblotting analysis of 4-HNE in lung tissue of mice on Day 7 and Day 14 after bleomycin administration. Left, representative Western blot result for 4-HNE. Right, scatter plot showing the results for all samples examined. Each lane represents one mouse, n = 3 mice per group. Statistical analysis was performed using one-way ANOVA followed by multiple comparisons. *P < 0.05, **P < 0.001. (e) Representative images of Masson’s staining and IHC staining for α-SMA, FAP and TGF-β in lung tissue from control group, bleomycin group and bleomycin + Lip-1 group at different days. Scale bars, 100 μm. (f) Scatter plot showing the percentage of the positive area of Masson’s staining and IHC staining in lung tissue from control group, bleomycin group and bleomycin + Lip-1 group at different days. n = 4 mice per group. Statistical analysis was performed using one-way ANOVA followed by multiple comparisons. **P < 0.01, ****P < 0.0001. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)

We then exposed MRC-5 cells to TGF-β and detected intracellular Fe2+ with an FeRhoNox-1 probe. As shown in Fig. 5c, the fluorescence intensity of the FeRhoNox-1 probe was increased with TGF-β stimulation, while both SB431542, a blocker targeting the TGF-β receptor, and DFO attenuated TGF-β-induced intracellular labile Fe2+ accumulation. Similarly, intracellular Fe2+ was increased by FAC, and this effect was reduced by DFO (Fig. 5d). In other words, both TGF-β and exogenous iron supplementation increased intracellular labile Fe2+ accumulation. To investigate the effect of elevated Fe2+ on fibroblasts, we stimulated MRC-5 cells with different concentrations of TGF-β and observed positive effects of TGF-β on the fibroblast-to-myofibroblast transition in all three types of cells. TGF-β stimulation increased both the mRNA and protein expression levels of α-SMA and COL1A1, which are classical markers of fibroblasts, in a dose-dependent manner (Supplementary Figs. 4a–d). Moreover, TGF-β significantly enhanced the migratory and invasive potential of the fibroblasts (Supplementary Fig. 4e).

3.4. TGF-β is involved in fibroblast activation by promoting intracellular Fe2+ accumulation

It is widely accepted that the bleomycin-induced lung injury model comprises two phases: an early inflammatory phase characterized by infiltration of inflammatory cells into the lungs within 7 days, followed by a late fibrotic phase characterized by increased fibroblast proliferation and differentiation to myofibroblasts, as well as synthesis of ECM during the second week [8,39,40]. Increasing evidence also suggests that alveolar epithelial injury might be the primary defect in IPF, which occurs relatively early in the inflammatory phase [41]. However, the number of α-SMA-positive fibroblasts increased from 7 days through 14 days but not within 4 days, in contrast to the control group (Fig. 3f). Flow cytometry demonstrated the accumulation of Fe2+ in α-SMA–positive fibroblasts in bleomycin-treated lung tissue on both Days 7 and 14, and the accumulation was decreased on Day 14 compared with the level on Day 7 (Fig. 5a and b). These results encouraged us to explore the reason that fibroblasts were abnormally activated under excessive iron, which is believed to be a trigger for lipid peroxidation and ferroptosis. During the initiation of the fibrotic phase, epithelial cell injury and activation of inflammation promote the production of the main profibrogenic cytokine TGF-β, which plays a critical role in the process of fibroblast differentiation into myofibroblasts and induces collagen production, causing a loss of lung elasticity and impairment of respiratory function [42,43]. To confirm the activation of fibroblasts induced by TGF-β, we stimulated two different human fibroblast cell lines (MRC-5 and IMR90 cell lines) and mouse primary lung fibroblasts (PLFs) with different concentrations of TGF-β and observed positive effects of TGF-β on the fibroblast-to-myofibroblast transition in all three types of cells. TGF-β stimulation increased both the mRNA and protein expression levels of α-SMA and COL1A1, which are classical markers of fibroblasts, in a dose-dependent manner (Supplementary Figs. 4a–d). Moreover, TGF-β significantly enhanced the migratory and invasive potential of the fibroblasts (Supplementary Fig. 4e).
Fig. 4. Liproxstatin-1 alleviates bleomycin-induced pulmonary fibrosis in the fibrosis stage. (a) H&E staining scanning of lung tissue from control group, bleomycin group and bleomycin + Lip-1 group at 10, 14, 18, and 21 days after bleomycin administration. Scale bars, 1 mm. (b) Hydroxyproline content from control group, bleomycin group and bleomycin + Lip-1 group at 10, 14, 18, and 21 days after bleomycin administration. n = 3–4 mice per group. Statistical analysis was performed using one-way ANOVA followed by multiple comparisons. *P < 0.05, **P < 0.01. (c) Representative images of Masson’s staining of mice lung tissue from control group, bleomycin group and bleomycin + Lip-1 group at 10, 14, 18, and 21 days after bleomycin administration. Scale bars, 100 µm. (d) Representative images of IHC staining for α-SMA of mice lung tissue from control group, bleomycin group and bleomycin + Lip-1 group at 10, 14, 18, and 21 days after bleomycin administration. Scale bars, 100 µm. (e) Scatter plot showing the percentage of the positive area of Masson’s staining in lung tissue from control group, bleomycin group and bleomycin + Lip-1 group at 10, 14, 18, and 21 days after bleomycin administration. n = 4 mice per group. Statistical analysis was performed using one-way ANOVA followed by multiple comparisons. *P < 0.05, ****P < 0.0001. (f) Scatter plot showing the percentage of the positive area of IHC staining in lung tissue from control group, bleomycin group and bleomycin + Lip-1 group at 10, 14, 18, and 21 days after bleomycin administration. n = 4 mice per group. Statistical analysis was performed using one-way ANOVA followed by multiple comparisons. **P < 0.01, ****P < 0.0001.

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(a) Fe²⁺ (α-SMA⁺ cells)

7 D

Normalized

Percentage of RhoNox-1 (F₂⁻⁺)

Ctrl vs BLM

14 D

Normalized

Percentage of RhoNox-1 (F₂⁻⁺)

Ctrl vs BLM

(c) Ctrl TGF-β TGF-β +SB TGF-β +DFO

(d) Ctrl FAC (50 μM) FAC+DFO

(e) FAC (μM)

Cell viability (CCK8)

(f) Ctrl FAC (50 μM) FAC+DFO

(g) MRC-5 Ctrl FAC (12.5 μM) + DFO FAC (25 μM) + DFO FAC (50 μM) + DFO

α-SMA COL1A1 α-tubulin

(kDa)

42 -100

55

IMR90

α-SMA COL1A1 α-tubulin

(kDa)

42 -100

55

(h) Ctrl TGF-β TGF-β +SB TGF-β +DFO

α-SMA COL1A1 α-tubulin

(kDa)

42 -100

55

(M) Ctrl TGF-β TGF-β +SB TGF-β +DFO

Migration

Invasion

(i) DAPI α-SMA Merge

Ctrl TGF-β TGF-β +SB TGF-β +DFO

(j) MRC-5 Ctrl TGF-β TGF-β +SB TGF-β +DFO

Migration

Invasion

(caption on next page)
accumulation in fibroblasts, we detected the expression of several iron-fibroblast activation markers. TFRC, a cell surface membrane protein, is involved in iron storage protein ferritin is composed of 24 subunits, including heavy-chain ferritin (FTH) and light-chain ferritin (FTL1), which can be assessed by Western blot after 72 h of treatment with FAC and DFO. FAC: 50 μM. DFO: 50 μM. Scale bars, 200 μm = 3 biological replicates. Statistical analysis was performed using one-way ANOVA followed by multiple comparisons. **P < 0.01. (f) Cell viability was determined by CCK8 after 72 h of treatment with FAC and DFO. FAC: 50 μM. DFO: 50 μM n = 4 biological replicates. Statistical analysis was performed using one-way ANOVA followed by multiple comparisons. *P < 0.05, **P < 0.01, ***P < 0.001. (g) Expression of α-SMA and COL1A1 in MRC-5 and IMR90 was assessed by Western blot after 72 h of treatment with the indicated doses of FAC and DFO. DFO: 50 μM. (h) Expression of α-SMA and COL1A1 in MRC-5 was assessed by Western blot after 72 h of treatment. TGF-β1: 10 ng/ml. SB: 1 μM. DFO: 50 μM. (i) Confocal microscopy showing α-SMA in MRC-5 after 72 h of treatment with TGF-β1, SB and DFO. DAPI (blue). α-SMA (red). TGF-β: 10 ng/ml. SB: 1 μM. DFO: 50 μM. Scale bars, 20 μm. (j) Representative microscopic images and the quantitation of MRC-5 cells that transferred through the transwell in the migration assay and invasion assays. Scale bars, 200 μm. TGF-β1: 10 ng/ml. SB: 1 μM. DFO: 50 μM n = 6 visual fields. Statistical analysis was performed using one-way ANOVA followed by multiple comparisons. ***P < 0.001. (For interpretation of the references to color in this figure legend, the reader is referred to the Web version of this article.)

3.5. The upregulation of TFRC expression via the TAZ-TEAD complex contributes to the function of TGF-β to promote iron accumulation and fibroblast activation

To clarify the mechanism involved in TGF-β-regulated ferrous iron accumulation in fibroblasts, we detected the expression of several iron-related proteins in MRC-5 cells stimulated with TGF-β1. As shown in Fig. 6a, the expression of TFRC and ferritin was upregulated upon TGF-β1 stimulation. TFRC, a cell surface membrane protein, is involved in cellular iron uptake by transporting transferrin-bound iron into the cell. Iron storage protein ferritin is composed of 24 subunits, including heavy-chain ferritin (FTH) and light-chain ferritin (FTL1), which can bind and store intracellular iron [26,27,44]. Thus, the elevation of FTH implied an increase in intracellular Fe

3.6. Regulation of the ferroptosis-defense pathway during fibroblast activation induced by iron overload

As shown in Fig. 7a and b, even low doses of FAC that induced fibroblast activation resulted in an increase in lipid peroxidation in fibroblasts, while DFO reduced the production of lipid peroxides. However, the accumulation of lipid ROS failed to trigger ferroptosis, indicating that fibroblasts might be protected from iron-induced oxidative stress by a certain ferroptosis-defense pathway (Fig. 7c). To investigate the molecular basis underlying the defense against ferroptosis, we first tested the expression of iron-related proteins in MRC-5 cells under long-term stimulation with TGF-β1. Prolonged treatment of MRC-5 cells with TGF-β1-stimulated high levels of α-SMA, SLC40A1 and FTH, and the lipid peroxidation inhibitor Fer-1 attenuated the effect of TGF-β1 (Fig. 7d). The upregulation of SLC40A1 and FTH suggested increased iron export and cytosolic iron storage, thus reducing the risk of ferroptosis caused by iron accumulation.

To investigate the mechanism of ferroptosis defense in iron-accumulating fibroblasts, we performed RNA sequencing in parental and FAC-stimulated MRC-5 cells with or without DFO treatment (Supplementary Fig. 5a). Heatmaps comparing the overall differences in gene expression among the three groups indicated that FAC stimulation caused many changes in the transcriptome of MRC-5 cells, which were reversed by DFO (Supplementary Fig. 5a). Data analysis revealed that the expression of a total of 78 genes was upregulated in the FAC group and that their expression was recovered in the FAC + DFO group; these differences were significantly different among the three groups, as determined through one-way ANOVAs (Fig. 7e). Through a Kyoto Encyclopedia of Genes and Genomes (KEGG) enrichment analysis of these significantly different genes, we unexpectedly found that FAC stimulation significantly altered certain pathways related to lipid metabolism and cholesterol metabolism in fibroblasts, which ranked in the top 20 enriched pathways (Fig. 7e). Differentially upregulated gene expression in the FAC group and differentially downregulated gene expression in the FAC + DFO group were associated with lipid metabolism, fatty acid metabolism and unsaturated fatty acid synthesis (Fig. 7f). These differences offered us some ideas. The main characteristics of ferroptosis are iron overload and the accumulation of lipid peroxides, and lipid metabolism greatly affects the synthesis of lipid peroxides and the sensitivity of ferroptosis. Therefore, we inferred that, in general, excessive accumulation of lipid peroxides in MRC-5 cells might be inhibited through the regulation of certain pathways related to the transcriptional regulation of TFRC was investigated with a dual-luciferase reporter assay system. pGL3-TFRC (−3000/−250) was cotransfected with pCMV6-Entry-TEAD4 into 293T cells. As shown in Fig. 6d, transfection of the TEAD4 plasmid significantly increased the luciferase activity of the TFR4 promoter region.
lipid metabolism when intracellular Fe\(^{2+}\) is moderately elevated; that is, a ferroptosis-defense pathway may be activated in these cells.

Subsequently, pairwise comparisons were performed within the control group and the FAC group and the FAC + DFO group, and the identified genes that might affect the accumulation of lipid peroxides and ferroptotic sensitivity are listed. For example, high expression of stearoyl-CoA desaturase (SCD) in the FAC group catalyzed the synthesis of monounsaturated fatty acids (MUFAs) in fibroblasts, which may protect cells from ferroptosis [48]. Fatty acid desaturase 2 (FADS2) is a key enzyme that regulates the biosynthesis of polyunsaturated Fatty acids (PUFAs). Some studies related to lung cancer have proven that knockdown of FADS2 can significantly reduce the expression of certain protective genes and increase the levels of lipid ROS in lung cancer cells, thus enhancing the sensitivity of cells to ferroptosis [49,50]. This outcome suggested that the upregulated expression of FADS2 in the FAC group might reduce the ferroptotic sensitivity of lung fibroblasts. In addition, acetyl-CoA acyltransferase 2 (ACAT2), a member of the acyl-coenzyme A-cholesterol acyltransferase (ACAT) family, can catalyze the generation of cholesterol esters, and it has been confirmed that high expression of ACAT2 can promote cell proliferation [51] (Supplementary Figs. 5b and c). Furthermore, we detected the mRNA expression of several key genes in MRC-5 cells in vitro after FAC and DFO stimulation, which was consistent with the RNA sequencing results (Supplementary Fig. 5d). Moreover, a Gene Set Enrichment Analysis (GSEA) provided some relevant evidence. Compared with the control group, genes involved in glutathione metabolism and the PPAR signaling pathway were enriched in the FAC group. Acyl-CoA synthetase long-chain family member 3 (ACSL3), with upregulated expression in the FAC group, converted MUFAs into phospholipid monounsaturated fatty acids (PL-MUFAs), thereby protecting cells from ferroptosis (Fig. 7g and h).

Evidence shows that overexpression of ACSL3 in melanoma cells may inhibit lipid oxidative stress through absorption of oleic acid, and high expression of ACSL3 has been associated with poor prognosis of patients with one of various tumors [52-54]. Compared with that in the FAC group, the expression of cell cycle and PPAR signaling pathway genes was decreased in the FAC + DFO group (Fig. 7i and j). On the one hand, this proves the effect of Fe\(^{2+}\) changes on cell proliferation; on the other hand, it demonstrates the effect of FAC on cell lipid metabolism.

3.7. Loss of TFRC in fibroblasts inhibits bleomycin-induced pulmonary fibrosis

Fibroblast-specific protein 1 (FSP1, also identified as S100A4) is considered as a marker of fibroblasts in different organs undergoing tissue remodeling and may serve as a sensitive and specific marker for lung fibroblasts [55]. As shown in Fig. 8a, multicolor immunofluorescence staining confirmed the colocalization of FSP1 and \(\alpha\)-SMA in bleomycin-induced fibrotic lung tissue. To determine the role played by TFRC and TFRC-associated iron uptake in the activation of fibroblasts in pulmonary fibrosis progression, we generated mice that lack TFRC only in fibroblasts by crossing mice expressing a conditional TFRC knockout allele (TFRC\(^{flox/flox}\)) with mice that express Cre recombinase driven by the FSP1 promoter (FSP1-CreER\(^T\)). The offspring (TFRC\(^{flox/flox}\), FSP1-CreER\(^T\)) are referred to hereafter as TFRC\(^{-/-}\), and TFRC in FSP1 \(^{-/-}\) cells were specifically knocked out after consecutive intraperitoneal injection of tamoxifen for 5 days. The bleomycin-induced pulmonary fibrosis model was then established with TFRC\(^{-/-}\) mice and TFRC\(^{flox/flox}\) mice (Fig. 8b). Western blot analysis of primary fibroblasts from TFRC\(^{flox/flox}\) and TFRC\(^{-/-}\) mice confirmed efficient conditional knockout (Fig. 8c). Comparing lung tissue 4, 7, 14, and 21 days after bleomycin administration, we found that the symptoms in TFRC\(^{-/-}\) mice were alleviated, compared to those in the TFRC\(^{flox/flox}\) mice, on Day 21 (Fig. 8d). H&E staining demonstrated that specific knockout of TFRC in fibroblasts alleviated the symptoms of bleomycin-induced pulmonary fibrosis, which was particularly significant in the later stage (Fig. 8e).

In addition, hydroxyproline content in lung tissues decreased in TFRC\(^{-/-}\) mice compared to TFRC\(^{flox/flox}\) mice 21 days after bleomycin administration (Fig. 8f). Fourteen days after bleomycin administration, \(\alpha\)-SMA, FSP1 and TFRC in the lung tissues of both groups were labeled by multicolor immunofluorescence. The expression of TFRC was significantly decreased in \(\alpha\)-SMA\(^{-/-}\) and FSP1\(^{+}\) cells (Fig. 8g). Masson’s staining showed that collagen deposition was relatively reduced in the lung tissues of the TFRC\(^{-/-}\) mice (Fig. 8h and i). The expression of \(\alpha\)-SMA and FSP1 was also decreased in the bleomycin-induced fibrotic lung tissues of the TFRC\(^{-/-}\) mice, as indicated by IHC staining (Fig. 8h and i). These results demonstrate that TFRC expression in fibroblasts effectively promotes the activation of fibroblasts and the occurrence of pulmonary fibrosis.

4. Discussion

The mechanisms involved in pulmonary fibrosis are not completely understood, and the effects of drugs on pulmonary fibrosis are unsatisfactory. As a novel type of programmed cell death driven by iron-dependent lipid peroxidation, ferroptosis is implicated in various diseases related to organ damage, including ischemia/reperfusion damage, drug-induced kidney injury, rhabdomyolysis-associated renal damage, and liver damage in chronic liver diseases [56-58]. Iron overload induces significant increases in lipid peroxide accumulation and ferroptosis in mouse primary hepatocytes, contributing to ferroptosis-related liver damage and fibrosis in mouse liver fibrosis models established either by high dietary iron or carbon tetrachloride (CCL4) injections [59-61]. In this study, we found that both bleomycin and LPS-induced iron overload and ferroptosis in lung epithelial cells, contributing to
epithelial damage in the early inflammatory phase of pulmonary fibrosis. Interestingly, the population of fibroblasts was increased even under iron-induced oxidative stress. This phenomenon was probably due to the following reasons. The half-life of bleomycin ranges from a few hours to 21 h in vivo [62,63]. It triggers epithelial injury, which is a key event in the onset of pulmonary fibrosis. AECs occupy more than 95% of the alveolar surface area in normal lung tissues and are most likely to come into direct contact with lethal external substances [64,65]. Notably, in the early stage of bleomycin-induced pulmonary fibrosis, AEC death was not limited to ferroptosis, as demonstrated in this study, and apoptosis and necroptosis of AECs have been confirmed by other studies [32,36,66]. In previous studies, LPS is often used to induce acute lung injury models, but it has been reported that LPS could also induce symptoms of pulmonary fibrosis, which may last shorter than bleomycin-induced pulmonary fibrosis. In our study, LPS was just selected as a supplement to the bleomycin-induced pulmonary fibrosis model in order to compare the protective effects of Liproxstatin-1 in different models of pulmonary fibrosis. Our results also confirmed that LPS-induced pulmonary fibrosis models supported our conclusions. During the following fibrotic phase in bleomycin-induced pulmonary fibrosis, the level of bleomycin remaining in lungs may not be high enough to trigger the death of activated fibroblasts. We also found that both the production of 4-HNE and the accumulation of ferrous iron decreased in the lungs of mice on Day 14 compared with that in the lungs on Day 7 after bleomycin administration. In other words, although fibroblasts might show accumulation of ferrous iron in this fibrotic phase, the iron concentration was more moderate than that in the early inflammatory phase. Notably, the increase in ferrous iron induced by low-dose FAC caused the production of lipid ROS without triggering ferroptosis. Mechanistically, our study revealed that activation of ferroptosis-defensive and iron exporting pathways protected fibroblasts from ferroptotic stress induced by moderate iron accumulation. Intracellular PUFAs are transformed into phospholipid polyunsaturated fatty acids (PL-PUFAs) by a series of catalytic processes and are eventually oxidized to lipid ROS or PL-PUFA-OH, which ultimately leads to ferroptosis. In contrast, PL-MUFAs derived from MUFAs contribute to cell resistance to ferroptosis. Undoubtedly, ferroptosis is greatly affected by cellular lipid metabolism [54]. From RNA-sequencing data, we observed that the expression of certain genes related to fatty acid biosynthesis and lipid metabolism pathways, such as SCD, FADS2, SQLE, APOE, ACAA, and FABP4, was altered after iron level elevation. A series of studies proved that the upregulated expression of these genes promoted the survival, migration and invasion of cancer cells and was associated with poor prognosis in people of tumors [48-50,67-71]. These upregulated genes inhibited the ferroptotic sensitivity of cells by influencing lipid metabolism and lipid peroxidation. In previous bioinformatics studies, researchers compared the gene signature in bronchoalveolar lavage fluid (BALF) or lung tissue between IPF patients and normal controls and found that certain differentially expressed ferroptosis-related genes were highly expressed in the IPF patients, which was associated with poor prognosis. Among these genes in our study, some were upregulated in FAC-stimulated fibroblasts, including MYC, AIFM2, PRKCA, RG54 and ZFP36. This result suggests that the expression of these genes may be upregulated when fibroblasts are activated by Fe<sup>2+</sup> in vivo, regulating ferroptotic risk and promoting pulmonary fibrosis [72-74]. MRC-5 cells exhibited upregulated expression of SLC40A1, which transports iron out of the cell, and FTH, which stores intracellular free iron, under long-term stimulation with TGF-β. The increase of these genes regulating iron homeostasis may protect fibroblasts against lipid oxidative stress and ferroptotic cell death by decreasing Fe<sup>2+</sup> levels. Chen et al. [72] analyzed and compared the ferroptosis-related gene signature in BALF between IPF patients and normal controls and thus identified 19 differentially expressed genes associated with poor prognosis. Among these differentially expressed genes, we found that SLC40A1 was highly expressed in the IPF patient group, which is consistent with the results of the in vitro experiments. This finding suggests that SLC40A1 may relieve the pressure of lipid peroxidation and the risk of ferroptosis by transporting iron out of the fibroblasts during the fibrotic stage in IPF patients.

Numerous studies have confirmed that the transition of fibroblasts to myofibroblasts is a key step for the excessive accumulation of ECM in fibrotic pathological diseases [75-77]. As mentioned above, Fer-1 inhibited the activating effect during the long-term stimulation of MRC-5 cells by TGF-β, suggesting that lipid peroxidation might positively regulate the activation of fibroblasts. Correspondingly, Lip-1 also alleviated pulmonary fibrosis when intraperitoneal injection was performed from the 10th day after bleomycin administration, which may have been due to the inhibition of fibroblast activation by Lip-1 not the ferroptosis of the epithelial cells in vivo. In fact, lipid peroxidation has been confirmed to promote the activation and transformation of in other fibroblast cell lines into myofibroblasts, which is consistent with our results [78]. Recently, COVID-19-induced acute respiratory distress syndrome was associated with clinical, radiographic, histopathological, and ultrastructural hallmarks of pulmonary fibrosis. The accumulation of monocyte-derived macrophages with enrichment of fibrosis-associated gene signatures, such as TGF-β, in IPF was significantly similar to the macrophage populations found in severe COVID-19. The lungs of patients with severe COVID-19 also had a pronounced expansion and activation of myofibroblasts and fibroblasts, similar to that of the pulmonary fibrosis models [79,80]. Therefore, characterizing the molecular mechanisms involved in fibroblast activation may further fuel the development of potential therapies to reverse phenotype transition in fibrotic tissue and control the progression of fibrotic diseases. A series of in vitro and in vivo studies have shown that TGF-β is a critical cytokine for the accumulation of ECM and the occurrence of fibrosis. External stimulation triggered damage to AECs and subsequent...
inflammatory responses, resulting in the secretion of profibrotic factors represented by TGF-β, which not only recruits and activates fibroblasts to transform into myofibroblasts but also promotes the epithelial-mesenchymal transition (EMT) [81,82]. Our findings revealed a novel mechanism that was involved in TGF-β-regulated fibroblast activation. TGF-β enhanced intracellular labile ferrous accumulation by upregulating TFRC via TAZ-TEAD signaling and promoted the fibroblast-to-myofibroblast transition in the fibrotic phase. Several previous studies have reported fibrosis-enhancing effects of iron. For example, Gao et al. [83] revealed that the iron overload of hepatic stellate cells (HSCs) was able to activate HSCs and drive liver lipogenesis and fibrosis. And iron elevated collagen gene expression and increased the proliferation of rat HSCs [84,85] and promoted cirrhosis in mice [86]. Similarly, studies with murine HSCs showed transferrin-induced iron overload and ferroptosis in lung epithelial cells in the early inflammatory phase. Moderate intracellular labile ferrous accumulation regulated by TFRC via the TGF-β-TAZ-TEAD signaling pathway promotes fibroblast-to-myofibroblast transition in the late fibrotic phase. These findings provide compelling evidence that therapeutic strategies for targeting ferroptosis and/or iron homeostasis disorders can be developed to treat iron overload-associated diseases, including pulmonary damage and fibrosis.

Ethics statement

The animal experiment had been reviewed and approved by the Laboratory Animal Welfare and Ethics Committee of the Air Force Medical University (approval certificate IACUC-20211102).

Declaration of competing interest

The authors declare no competing interests.

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