Irbesartan ameliorates myocardial fibrosis in diabetic cardiomyopathy rats by inhibiting the TGFβ1/Smad2/3 pathway

MIN ZONG, HUA ZHAO, QIANG LI, YANBING LI and JIANJUN ZHANG

Department of Cardiology, Beijing Chaoyang Hospital, Capital Medical University, Beijing 100043, P.R. China

Received July 27, 2019; Accepted May 15, 2020

DOI: 10.3892/etm.2020.9245

Abstract. Myocardial fibrosis (MF) is an important pathological change in diabetic cardiomyopathy. The aim of the present study was to investigate whether irbesartan serves a role in improving MF in a diabetic rat model. Fasting blood glucose (FBG), total cholesterol (TC), triglyceride (TG), high-density lipoprotein cholesterol (HDL-C) and low-density lipoprotein cholesterol (LDL-C) levels were measured in rats using biochemical methods. Heart weight index (HWI), left ventricular weight index (LVWI), left ventricular systolic pressure (LVSP) and left ventricular end-diastolic pressure (LVEDP) were also measured, whilst type I collagen and hydroxyproline content in myocardial tissue was quantified. Western blotting was used to measure the expression of transforming growth factor β1 (TGFβ1), phosphorylated (p)-Smad2/3 and collagen type I α 1 chain (COL1A1) in myocardial tissues or rat cardiac fibroblast (RCF) cells. Cell proliferation was measured using EdU staining. Procollagen type III N-terminal peptide (PIIINP) content, FBG, TC, TG and LDL-C levels were found to be significantly higher, whilst HDL-C levels were found to be significantly lower in rats in the diabetic group. Those in the diabetic group also exhibited significantly elevated HWI, LVWI, LVEDP, myocardial tissue type I collagen content and hydroxyproline content values, but significantly reduced LVSP. Changes in the aforementioned indicators were reversed after treatment with irbesartan, where the protein expression levels of TGFβ1 and p-Smad2/3 in myocardial tissue were also significantly reduced. In RCF cells, irbesartan significantly reversed high glucose-induced upregulation of TGFβ1 expression, Smad2/3 phosphorylation and COL1A1 expression, as well as reducing cell proliferation and rat type I PICP and PIIINP levels. Application of pirfenidone produced additive effects on reducing the expression levels of the proteins aforementioned when combined with irbesartan. Therefore, the present results demonstrated that irbesartan reduced the activity of the TGFβ1/Smad2/3 pathway and ameliorated diabetic MF by downregulating the expression of TGFβ1.

Introduction

Diabetic mellitus (DM), which describes a series of metabolic disorders affecting protein, fat and electrolytes, is caused by the absolute or relative deficiency of insulin secretion or decreased insulin sensitivity in target tissue cells, which is mainly characterized by abnormally elevated blood sugar content (1,2). The metabolic disorder caused by DM directly affects cardiac myocytes and cardiac fibroblasts, changing the function of cardiac myocytes and the deposition of collagen in the cardiac interstitium, leading to decreases in myocardial compliance and diastolic function (3). As the disease progresses, the systolic function of the heart also decreases, causing congestive heart failure, cardiogenic shock and sudden mortality, which seriously affect the life and health of patients with DM (4). Myocardial fibrosis (MF) is an important factor in cardiac diastolic, contractile function impairment and congestive heart failure during the progression of diabetic cardiomyopathy (DCM) (5-7).

Previous studies have shown that transforming growth factor β1 (TGFβ1) regulates the transcription of related extracellular matrix (ECM) components by activating downstream Smad proteins and serves an important regulatory role in fibrosis in various tissues and organs (8-10). In the condition of diabetic pathological hyperglycaemia, the renin angiotensin aldosterone system (RAAS) is activated, which increases blood circulation and angiotensin II (Ang II) content in the myocardium (11). Furthermore, Ang II increases the expression of a variety of fibrotic factors, including transforming growth factor β (TGF-β), connective tissue growth factor (CTGF) and collagen in the myocardial interstitium whilst reducing the degradation of collagen, thus promoting the development of diabetic MF (12,13). Irbesartan is an Ang II receptor antagonist that can effectively block the binding of Ang II to its receptor and inhibit its downstream biological functions (14). Currently, irbesartan is widely used in the clinic, mainly for the treatment of hypertension and type II diabetic nephropathy with hypertension (15). In addition, previous studies have reported that irbesartan can improve MF induced by hypertension (16-18); however, to the best of our knowledge, there have been an insufficient number of studies on the effects of irbesartan on...
MF in patients with DM. Therefore, the aim of the present study was to investigate whether irbesartan serves a role in improving MF in a diabetic rat model.

**Materials and methods**

**Ethical statement.** The present study was approved by the Animal Ethics Committee and was conducted in accordance to the relevant agreements with the Beijing Chao yang Hospital of the Capital Medical University (approval no. 20170523). All procedures were performed in accordance with the Guidance Suggestions for the Care and Use of Laboratory Animals (19).

**Main reagents and materials.** DMEM, FBS and penicillin-streptomycin were purchased from Gibco (Thermo Fisher Scientific, Inc.), and a hydroxyproline assay kit (cat.no. A030-1-1) was purchased from Nanjing Jiancheng Bioengineering Institute. Irbesartan was purchased from Sanofi (Hangzhou) Pharmaceutical Co., Ltd. Healthy adult male Sprague Dawley (SD) rats (n=30; age, 6 weeks; weight, 220-240 g) were purchased from the Guangdong Medical Laboratory Animal Center. A high-fat and high-sugar diet was purchased from Jiangsu Synergy Pharmaceutical Bioengineering Co., Ltd. (https://www.jsxtsw.com/) and streptozotocin (STZ) citrate buffer was purchased from Sigma-Aldrich (Merck KGaA). A rat collagen I ELISA kit (cat. no. 20180302A) was purchased from Suzhou Calvin Biotechnology Ltd. (https://bio.calvin.biogeo.net/) and rat cardiac fibroblasts (RCFs) were purchased from Cloud-Clone Corp. (cat. no. CSI095Ra01). The TGFβ1 inhibitor pirfenidone was purchased from Selleck Chemicals. Rabbit anti-human TGFβ1 (cat. no. ab92486; 1:2,000), anti-phosphorylated (p)-Smad2/3 (cat. no. ab272332; 1:2,000), Smad2/3 (cat. no. ab63672, 1:2,000), and anti-collagen type I α1 (COL1A1; cat. no. ab34710; 1:2,000) polyclonal antibodies were purchased from Abcam. ECL chemiluminescence solution and bicinechonic acid protein quantitative detection kits were purchased from Beyotime Institute of Biotechnology. Rat type I procollagen carboxy-terminal peptide (PICP; cat. no. CSB-E08081r) and Type III procollagen amino terminal pro-peptide (PIIINP) ELISA kits (cat. no. CSB-E08096r) were purchased from CusabioTechnology LLC. Cholesterol and PBS were obtained from Sangon Biotech Co., Ltd. Biological function was assessed with a BL-420L biological signal collection system (Chengdu Tai meng Software Co., Ltd.), and flow cytometry was performed using a Beckman Coulter FC500 MCL flow cytometer (Beckman Coulter, Inc.).

**Establishment of the DCM rat model.** Rats were allowed free access to water and food under 22±2°C, 55±10% humidity and 12-h light/dark cycle. A rat model of type 2 DM was established by feeding freely with a high-fat and high-sugar diet combined with an intraperitoneal injection of a low dose of STZ (40 mg/kg). After 1 week of adaptive feeding of SD rats, the model was established. DM model rats were fed a high-fat and high-sugar diet for 2 months. After the rats were fasted for 12 h, 1% STZ citrate buffer was injected into the abdominal cavity (40 mg/kg). The model was considered successful when the fasting blood glucose (FBG) concentration was >16.7 mmol/l for 3 consecutive days. Blood was obtained by using the blood glucose meter blood sampling needle to pierce the tip of the tail of the rat and then squeezed a drop of blood onto the blood glucose test paper to test the blood glucose level. Food intake, drinking water intake and urine output were significantly higher in model rats compared with control rats, indicating that the DM model was successfully established. The rats remained on the high-fat and high-sugar diet for a further 10 weeks.

The eight rats in the control group were fed on a standard diet, given intraperitoneal injections of citrate buffer (10 ml/kg, no STZ) after 2 months of feeding and continued on the standard feed until the end of the experiment.

**Irbesartan treatment of DCM rats.** In total, 30 rats were randomly divided into the blank control group (n=8) and the DCM1 group (n=22). The remaining rats in which the DCM model was successfully established were divided into two groups: DCM group (n=6) and the DCM + irbesartan group (n=6). Irbesartan was intragastrically administered at 50 mg/kg (20) once daily for 10 weeks. In the DCM and blank control groups, a corresponding volume of saline (10 ml/kg) was administered by gavage. DCM rats were fed the high-fat and high-sugar diet, whilst rats in the blank control group were fed on the standard maintenance diet for 10 weeks.

**Blood glucose and blood lipid detection.** Rats were anaesthetized with an intraperitoneal injection of chloral hydrate (300 mg/kg), before 2 ml blood was obtained from the orbital venous plexus and a sterile cotton ball was used to stop the bleeding immediately after pressing the orbit, following which the rats resumed normal diet and activities. FBG, total cholesterol (TC), triglyceride (TG), high-density lipoprotein cholesterol (HDL-C) and low-density lipoprotein cholesterol (LDL-C) levels were measured with a 7180 Automatic Biochemical Analyzer (Hitachi, Ltd.).

**Heart weight index (HWI) and left ventricular weight index (LVWI) determination.** After the rats were weighed, they were anaesthetized with an intraperitoneal injection of chloral hydrate (300 mg/kg) and the hearts were removed. After the heart tissue was washed with saline and filter paper was used to absorb excess water, each heart was weighed. Subsequently, the left and right atrium and the right ventricle were removed, and the left ventricle was weighed to provide the LVW. HWI and LVWI were calculated using the following equations:

\[
\text{HWI} = \frac{\text{HW}}{\text{BW}}, \quad \text{LVWI} = \frac{\text{LVW}}{\text{BW}}
\]

The heart tissue was then stored at -80°C for further experimentation.

**Cardiac function evaluation.** Cardiac haemodynamic indexes of each group were measured by cannulation of the carotid artery, and cardiac function was evaluated after intraperitoneal injection of 10% chloral hydrate (300 mg/kg). The right cervical artery was isolated, exposed and cannulated. The left ventricle was cannulated through the right common carotid artery. The left ventricular systolic pressure (LVSP) and left ventricular end-diastolic pressure (LVEDP) were detected using the BL-420L biological signal collection system.

**Detection of myocardial hydroxyproline.** In total, 100 mg myocardial tissue and 400 μl pre-cooled PBS were added to
a manual glass homogenizer, which was then evenly grinded it in an ice box and poured into a 1.5 ml tube. This mixture was centrifuged at 4,000 x g 5 min at 4°C, before the supernatant was collected. The hydroxyproline content was detected by ELISA according to manufacturer's protocol. The plates included blank, standard and sample wells. A myocardial hydroxyproline standard sample (10 µl) was first diluted in a 40 µl solution. The plate was sealed and incubated at 37°C for 30 min. The sealing film was then carefully removed, the liquid was discarded and the plate was dried and washed with PBS five times for 30 sec each. Enzyme-labelling reagent (50 µl) was added to each well (except the blank well), and the plate was sealed and incubated at 37°C for 30 min. The sealing membrane was carefully removed from the plate, and the wells were filled with washing liquid. This process was repeated five times. Chromogenic agent A (50 µl) was added to each well and then 50 µl colour agent B was added. Subsequently, the plates were lightly shaken at 37°C for 15 min in the dark. Terminating solution (50 µl) was added to each well to terminate the reaction. With the blank well set to zero, the absorbance of each well was measured at 450 nm.

**RCF cell culture and treatment.** RCF cells were inoculated into culture dishes (1x10^4/well), and DMEM supplemented with 10% FBS and 1% penicillin-streptomycin was added. Cells were cultured in a CO₂ culture box with 5% CO₂ at 37°C. After the cells were grown to 80-90% confluency, cells were then cultured at a 1:5 ratio and tested within four generations.

According to the different treatment concentrations of D-glucose (Dalian Meilun Biotechnology Co., Ltd.), the RCF cells were divided into the normal-glucose concentration group (5 mM D-glucose; control group) and the high-glucose (HG) treatment group (25 mM D-glucose). RCF cells treated with HG were further divided into three groups: HG group, the HG + irbesartan (10 µM) group and the HG + irbesartan (10 µM) + pirfenidone (5 mM) group. Cells in each treatment group were cultured (5% CO₂, 37°C) for 48 h in vitro and collected for mRNA and protein detection.

**EdU detection of cell proliferation.** After RCF cells were incubated in complete culture medium for 2 h at 37°C with 10 µl EdU, the cells were cultured for an additional 48 h in accordance with the aforementioned groups and digested with 0.2% trypsin at 25°C for 60 sec. After PBS washing and centrifugation (500 x g, 25°C, 5 min), cells were incubated in 10% neutral formaldehyde at room temperature for 15 min. PBS was then added to the cells, which were then centrifuged at 300 x g for 5 min at 25°C. Subsequently, 100 µl 0.5% Triton X-100 in PBS was added to the cells at room temperature and 500 µl reaction liquid was added for detection. Cell proliferation was detected after incubation at room temperature for 30 min, followed by the addition of 3 ml washing liquid and centrifugation of the washing liquid. The cell pellets were then resuspended in 500 µl washing liquid and flow cytometry was performed using a Beckman Coulter PC 500 MCL flow cytometer, which was analyzed using the FlowJo software (FlowJo v7.6.5; FlowJo LLC).

**ELISA detection of type I collagen in rat tissues and PICP and PIIINP in RCF cell culture medium.** PBS was added to the myocardial tissue at 1:5 (myocardial tissue:PBS), which was then homogenized and centrifuged at 1,000 x g 10 min at 4°C before the supernatant was obtained. Type I collagen content was then measured in the supernatant according to the manufacturer's instructions.

**Western blotting of protein expression.** After SDS lysis buffer (cat. no. P0013G; Beyotime Institute of Biotechnology) was added to the cells or myocardial homogenate, each sample was boiled for 5 min and the bicinchoninic acid method was used to detect the protein concentration. In total, 60 µg sample was then separated by 5% SDS-PAGE and transferred to a PVDF membrane. The membrane was blocked with 5% skimmed milk powder for 1 h at room temperature and then incubated with primary antibodies (cat. no. G-21234; Thermo Fisher Scientific, Inc.) for 1 h at room temperature. The membranes were subsequently detected with ECL and analyzed using the Quantity One image analysis software (version 4.5.0; Bio-Rad Laboratories, Inc.).

**Statistical analysis.** Statistical analysis was performed using SPSS 18.0 software (SPSS, Inc.). Data are presented as the mean ± standard deviation from three experimental repeats. Differences among >2 experimental groups were evaluated by one-way ANOVA followed by Tukey's test. P<0.05 considered to indicate a statistically significant difference.

**Results**

**Abnormal blood glucose and blood lipid content in DCM model rats.** The blood glucose test results indicated that the FBG content in the DCM group was significantly higher compared with the control group (P<0.001). Moreover, the levels of TC, TG and LDL-C in the DCM group were significantly higher compared with the control group (P<0.05), while the levels of HDL-C were significantly lower (P<0.05), as presented in Table I.

**DCM model rats exhibit notable MF.** The cardiac test results demonstrated that HWI and LVWI in the DCM group were significantly higher compared with the control group (P<0.001; Table II). ELISA results suggested that the myocardial concentrations of type I collagen and hydroxyproline in the DCM group were significantly increased compared with that in the control group (P<0.001; Table II). The LVSP were significantly decreased (P<0.05; Table II) and LVEDP were significantly increased (P<0.001; Table II) in the DCM group compared with those in the control group (P<0.001; Table II).
Irbesartan inhibits the TGFβ1/Smad2/3 pathway and reduces MF in DCM rats. The blood glucose test indicated that there was no significant difference in FBG content between the DCM group and the irbesartan group. Cardiac measurements identified that, compared with the DCM group, the DCM + irbesartan group had a significantly lower HWI and LVWI (P<0.001). Furthermore, compared with the DCM group, the DCM + irbesartan group had significantly reduced levels of collagen I and hydroxyproline in myocardial tissue (P<0.001), as presented in Table III.

Irbesartan alleviates DM-induced cardiac dysfunction. Compared with the DCM group, the DCM + irbesartan group had a significantly higher LVSP (P=0.016) but significantly lower LVEDP (P<0.001; Table IV). Thus, these data indicated that irbesartan improved myocardial contractile function and compliance in DM rats.
Irbesartan inhibits the TGFβ1/Smad2/3 pathway and HG-induced fibrosis in RCF cells. Western blot analysis results demonstrated that the expression levels of TGFβ1 and p-Smad2/3 in RCF cells were significantly higher in the HG treatment group compared with the control group. In addition, HG treatment significantly increased the expression of COL1A1 in RCF cells, indicating that the ability of RCF cells to synthesize collagen was enhanced and that fibrosis was significantly changed.

After treatment of HG-exposed RCF cells with irbesartan, TGFβ1 expression was significantly decreased, TGFβ1/Smad2/3 pathway activity was significantly inhibited, COL1A1 expression was reduced. When combined with irbesartan, treatment with the TGFβ1 inhibitor pirfenidone reduced TGFβ1/Smad2/3 pathway activity, COL1A1 expression further in RCF cells under HG treatment conditions (Fig. 2).

Irbesartan inhibits RCF cell proliferation and collagen synthesis. The proliferative ability of RCF cells in the HG treatment group was significantly increased compared with the control group. However, irbesartan treatment reduced the RCF proliferative ability, which was further decreased by combined treatment with irbesartan + pirfenidone (Fig. 3).

The ELISA results demonstrated that PICP and PIIINP contents in HG RCF cell culture conditions were significantly higher compared with the control culture conditions. It was found that irbesartan treatment significantly reduced the PICP and PIIINP content in the culture solution, while combined treatment with irbesartan + pirfenidone resulted in the lowest PICP and PIIINP content in the culture solution (Table V).

Discussion

Previous studies have reported that TGFβ1 plays an important role in regulating fibrosis in various tissues and organs, such as cardiac (21), liver (22), lung (23) and kidney (24) tissue, by activating downstream Smad proteins. TGFβ1 must bind with specific cell-surface receptors to exert its corresponding
biological effects. In mammalian cells, three types of TGFβ receptors (TβRs) are expressed: TβR-I, TβR-II and TβR-III (25). Moreover, TβR-III lacks intrinsic activity and has no signal transduction function itself. The active forms of TβR-II and TβR-III respond to stimuli in the body and function in signal transduction (26). Smads have signal transduction functions and act in the cytoplasm with TβR-Ⅰ kinase, which plays a key role in the regulation of the TGFβ1/Smad pathway; within this pathway, Smad2 and Smad3 mediate TGFβ1 protein signal transduction (27,28). Under the stimulation of various factors, TGFβ1 combines with TβR-I and TβR-II binds to TβR-I in response to phosphorylation (25,26). Phosphorylation-activated TβR-I then combines with Smad2/3. Moreover, phosphorylation-activated Smad2/3 combines with Smad4 to form a protein polymer, which is transferred from the cytoplasm to the nucleus and regulates the transcription and expression of a variety of downstream target genes, including Col-I, Col-III and α-SMA (29-31). In a positive feedback mechanism, the protein complex acts on the promoter region of the TGFβ1 gene to promote its expression and TGFβ1 autocrine signalling. In addition, TGFβ1 plays a positive feedback role by promoting the expression of the TβR-I and TβR-II receptor genes and amplifying the TGFβ1/Smad2/3 signalling pathway (29,30).

TGFβ1 is the regulatory factor most closely related to MF (21). Increased expression and activation of TGFβ1 occurs within a common pathway and acts as an intermediate bridge for MF caused by several factors. Thus, TGFβ1 is considered to be a direct MF factor (32). Previous studies have revealed that Ang II signalling can lead to collagen deposition via a variety of pathways, including the 5’adenosine monophosphate-activated protein kinase-angiotensin converting enzyme 2 and CTGF-fractalkine pathways, leading to fibrosis (33,34). Irbesartan, an Ang II receptor antagonist, affects the biological function of Ang II by blocking the binding of Ang II to its receptor (14). In a previous study examining ventricular fibrosis induced by hypertension, irbesartan regulated the degradation and synthesis of ECM by downregulating the expression of matrix metalloproteinase-2 and TIMP metallopeptidase inhibitor 2 to reduce the degree of ventricular MF (35). Irbesartan is widely used in the treatment of hypertension and type II diabetic nephropathy with hypertension and has been shown to alleviate MF caused by renal vascular hypertension (15). However, to the best of our knowledge, there have been few reports in China or elsewhere on whether irbesartan can reverse DM-induced MF. Therefore, the aim of the present study was to investigate whether irbesartan can improve MF in DM model rats.

Table V. PICP and PIIINP levels in RCF cell culture medium from each treatment group.

| Group                        | Control | HG            | HG + Irbesartan | HG + Irbesartan + Pirfenidone |
|------------------------------|---------|---------------|----------------|-----------------------------|
| PICP (pg/ml)                 | 615.8±31.9 | 1186.6±43.8<sup>a</sup> | 926.5±36.9<sup>a</sup> | 811.8±35.1<sup>a</sup> |
| PIIINP (pg/ml)               | 236.5±18.7 | 458.4±37.5<sup>a</sup> | 364.3±26.5<sup>a</sup> | 306.7±24.7<sup>a</sup> |

<sup>a</sup>P<0.05 vs. control group. HG, high glucose; PICP, procollagen carboxyl-terminal peptide; PIIINP, procollagen type III N-terminal peptide.
Lipid metabolism dysfunction is an important factor in type 2 DM pathogenesis (36). Therefore, the present study examined the serum TC level in STZ-induced DM model to determine whether the model was successfully established. Compared with the control group, the DMC group had significantly higher blood glucose levels and abnormal blood lipid levels, indicating the successful establishment of the DM rat model. Furthermore, the HWI and LVWI of DCM rats were significantly increased. The myocardial content of type I collagen and hydroxyproline in DCM rats was significantly higher compared with control rats, suggesting that the myocardial tissue of DCM rats had obvious fibrotic changes. In line with other studies, such as that by Shen et al (37), significantly increased levels of the type I collagen COL1A1 have been shown in cardiac muscle tissue in DM rats compared with control rats.

In the present study, irbesartan-treated DCM rats exhibited significantly milder fibrosis and lower TGFβ1 and p-Smad2/3 expression in the myocardium compared with DCM rats, suggesting that irbesartan may reduce TGFβ1/Smad2/3 pathway activation to alleviate MF in DM rats by reducing the expression of TGFβ1. Lv et al (38) investigated irbesartan and visceral fibrosis in diabetic nephropathy model animals, and found that irbesartan significantly reduced the expression of TGFβ1 in the glomerular tissue of diabetic nephropathy rats. Moreover, Tunçdemir et al (39) revealed that irbesartan treatment significantly reduced TGFβ1 expression in the glomeruli of diabetic nephropathy rats. Furthermore, irbesartan has been shown to significantly inhibit remodelling and fibrosis of the myocardium in hypertensive rats by downregulating the expression of TGFβ1 (40). In a study by Kataoka et al (41), irbesartan was revealed to serve a role in downregulating the expression of TGFβ1 and inhibiting MF in atrial tachycardia models. In addition, Tanaka et al (42) demonstrated that irbesartan significantly inhibited bleomycin-induced pulmonary fibrosis and reduced the expression of TGFβ1 in lung tissue. It has also been shown that the degree of renal tubulointerstitial fibrosis is significantly lower in irbesartan-treated animals compared with untreated animals, and that the expression of TGFβ1 and the phosphorylation of Smad2 are also significantly inhibited in irbesartan-treated animals (43). Using a rat model of hepatic fibrosis induced by bile duct ligation, Kim et al (44) reported that irbesartan significantly inhibited the expression of TGFβ1 in the liver and reduced the severity of liver fibrosis. Thus, these previous findings suggest that irbesartan can inhibit fibrosis of the viscera by regulating the expression of TGFβ1 and downregulating the activity of the TGFβ1/Smad2/3 pathway, which is consistent with the present results.

Through further in vitro experiments, the present study also demonstrated that irbesartan decreased TGFβ1 expression in RCF cells, inhibiting Smad2/3 activity. Enhancement and overactivation of cardiac fibroblast proliferation play important roles in the development of MF (45). Cardiac fibroblast activation mainly manifests as increased amounts of ECM around cardiac fibroblasts (46). Moreover, ratio of type I collagen to type III collagen synthesis is an important metric for cardiac fibroblast activation (47). Therefore, detection of the amounts of the PICP before the formation of type I collagen and PIIINP before the formation of type III collagen has become a common method that reflects the content of type I and type III collagen proteins (48). The present study found that irbesartan significantly inhibited the proliferation of RCF cells under HG conditions and reduced the content of PICP and PIIINP. Combined treatment with pirfenidone further enhanced the inhibitory effect of irbesartan on RCF cell proliferation, collagen synthesis and fibroblast activation. As there is no RAAS in cultured cells in vitro, irbesartan may regulate the expression of TGFβ1 directly or indirectly via other mechanisms and affect the activity of the TGFβ1/Smad2/3 pathway, thus inhibiting cardiac myocyte fibrosis. Similarly, in vitro studies by Lv et al (38) have revealed that irbesartan significantly downregulated the expression of TGFβ1, collagen IV and fibronectin in HG-treated glomerular mesangial cells and inhibited fibrosis of HG-treated mesangial cells, indicating that irbesartan can inhibit fibrosis by directly downregulating TGFβ1 without relying on the RAAS system. Thus, it was indicated that lowering the expression of TGFβ1 and the activity of the TGFβ1/Smad2/3 pathway may be one of the mechanisms via which irbesartan reduces MF in DM rats. However, irbesartan may also regulate MF in DM rats via other mechanisms; for instance, Liu et al (49) have shown that irbesartan alleviated MF in DM rats by regulating protein kinase D and the endoplasmic reticulum stress system. As this study was limited in its examination of other possible mechanisms, future studies will investigate the other possible mechanisms, including those of oxidative stress or immunomodulation, via which irbesartan affects diabetic MF.

In conclusion, the present results suggested that irbesartan may downregulate TGFβ1 expression, inhibit TGFβ1/Smad2/3 pathway activity and promote collagen synthesis to reduce MF lesions and improve myocardial function.

Acknowledgements

Not applicable.

Funding

No funding was received.

Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

Authors’ contributions

MZ and HZ designed the experiments. QL, YL and JZ performed the experiments. YL and JZ collected and analyzed the data. MZ and HZ drafted the manuscript. All authors read and approved the final manuscript.

Ethics approval and consent to participate

The present study was approved by the Animal Ethics Committee and was conducted in accordance to the relevant agreements with the Beijing Chaoyang Hospital of the Capital Medical University (approval no. 201705231). All procedures were performed in accordance with the Guidance Suggestion
for the Care and Use of Laboratory Animals, formulated by the Ministry of Science and Technology of China.

Patient consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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