A small molecule inhibitor MCC950 ameliorates kidney injury in diabetic nephropathy by inhibiting NLRP3 inflammasome activation

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Background: Diabetic nephropathy (DN) is a lethal diabetic microvascular complication characterized by chronic low-grade inflammation. The NOD-like receptor pyrin domain-containing protein 3 (NLRP3) inflammasome is implicated in the progression of DN. MCC950 is a selective and potent inhibitor of NLRP3; however, its efficacy in DN requires further investigation.

Methods: To investigate the efficacy of MCC950 in DN, eight-week-old type 2 diabetic db/db mice received injections of MCC950 intraperitoneally (10 mg/kg) twice per week for 12 weeks. Urinary albumin-to-creatinine ratio (ACR) and neutrophil gelatinase-associated lipocalin (NGAL), renal function, pathological changes, markers of podocyte and fibrosis and NLRP3/caspase-1/IL-1β expression in the renal cortices of db/db mice were evaluated.

Results: The NLRP3 inflammasome was activated in db/db mice and HG-induced mesangial cells by upregulating NLRP3/caspase-1/IL-1β pathway. Inhibition of the NLRP3 inflammasome with MCC950 reduced the production of active caspase-1 and active IL-1β in db/db mice and HG-induced mesangial cells. MCC950 reduced serum creatinine, urinary ACR and NGAL, attenuated mesangial expansion with increased matrix and tubular dilatation, alleviated thickened glomerular basement membrane (GBM) and foot process fusion without affecting body weight and blood glucose levels in db/db mice. MCC950 increased the expression of podocin in db/db mice, and decreased the expression of TGF-β1, fibronectin, collagen I and α-smooth muscle actin (α-SMA) in renal cortices of db/db mice and HG-induced mesangial cells.

Conclusion: MCC950 ameliorated renal function, thickened GBM, podocyte injury and renal fibrosis in db/db mice, and decreased the production of fibrosis markers in HG-induced mesangial cells. MCC950 effectively ameliorated diabetic kidney injury by inhibiting NLRP3/caspase-1/IL-1β pathway, which may be a potential therapeutic strategy to prevent the progression of DN.

Keywords: diabetic nephropathy, MCC950, NLRP3 inflammasome, db/db mice, mesangial cells

Introduction
Diabetic nephropathy (DN), a common diabetic microvascular complication, is the primary cause of end-stage renal disease (ESRD) worldwide.1,2 Type 2 diabetes mellitus (T2DM) patients occupy almost 50% of new cases of ESRD in the US...
population. Patients with ESRD as a result of diabetes require renal replacement therapy, contributing to considerable individual and socioeconomic costs worldwide.

Recent studies have reported persistent inflammation in circulatory and renal tissues; in particular, NLRP3 inflammasome-mediated inflammation is crucial to the pathogenesis of DN. NLRP3 inflammasome activation is upregulated in type 2 diabetic patients. NLRP3 is a useful biomarker discriminating DN patients from type 2 diabetic patients. In the innate immune system, nucleotide-binding oligomerization domain (NOD)-like receptors (NLRs) are pattern recognition receptors that recognize pathogen-associated molecular patterns and danger-associated molecular patterns (DAMPs). Upon detection of cellular stress, intracellular NOD-like receptor pyrin domain-containing protein 3 (NLRP3) combines with apoptosis-associated speck-like protein (ASC) that contains a caspase recruitment domain (CARD) and binds pro-caspase-1 to form the NLRP3 inflammasome. The NLRP3 inflammasome promotes pro-caspase-1 self-cleavage to generate the active caspase-1 p10/p20 tetramer that induces pro-IL-1β maturation and activates IL-1β p17 secretion. NLRP3 inflammasome activation occurs both in innate immune cells and in nonimmune cells intrinsic renal cells (mesangial cells, podocytes and epithelial cells); therefore, targeting the NLRP3 inflammasome may be a promising therapeutic strategy for DN.

MCC950 is a highly selective, potent small molecule inhibitor of NLRP3; it has been associated with the treatment of several diseases. Early NLRP3 inhibition by MCC950 prevented kidney fibrosis in a murine model of crystal nephropathy. MCC950 reduced renal inflammation, fibrosis and dysfunction in mice with established hypertension. MCC950 inflammasome inhibition by MCC950 reduced atherosclerotic lesion development. MCC950 attenuated severe, steroid-resistant asthma. The cytotoxicity of MCC950 was tested using human kidney and liver cell lines HEK293 or HepG2, and no evidence of toxicity was found. Nevertheless, the efficacy of MCC950 for DN requires further study. The db/db mouse was identified initially in 1966 at Jackson Labs and is widely used as a mouse model of type 2 diabetic nephropathy, because db/db mice and patients with DN share similar renal pathological changes. To demonstrate whether MCC950 protects against progression of DN, we used type 2 diabetic db/db mice in vivo and glomerular mesangial cells in vitro to investigate NLRP3/caspase-1/IL-1β expression and the efficacy of MCC950.

Materials and methods

Animals

Seven-week-old diabetic male db/db (C57BLKS/J-leprdb/leprdb) mice and non-diabetic male db/+ littermates (C57BLKS/J-leprdb/) were purchased from the Model Animal Research Center of Nanjing University (Nanjing, China). The mice were maintained in a room with controlled environment (23±3°C, 50%±20% humidity, 12-hr light/dark cycle) with free access to water and food. All procedures were carried out according to the approved Institutional Animal Care and Use Committee protocol of China Medical University (Number: 16095M). The mice were randomly divided into three groups (n=10 per group). For the MCC950 treated group (db/db+MCC950), db/db mice were treated with 10 mg/kg of MCC950 (Selleck Chemicals, Houston, USA) twice per week intraperitoneally from 8 weeks of age to 20 weeks of age. The control (db/+ ) and untreated groups (db/db) received an equal volume of vehicle (saline). The treatment lasted 12 weeks. Body weight (BW) was measured weekly, and fasting blood glucose levels were measured every 4 weeks. All mice were sacrificed at 20 weeks of age. All mice were anesthetized with pentobarbital, and blood samples were collected by orbital vein bleeding. Kidney weight (KW) was measured after cardiac perfusion as described previously. Renal cortical samples were harvested for subsequent studies.

Cell culture

The rat mesangial cell line HBZY-1 was purchased from the China Center for Type Culture Collection (Wuhan, China) and were cultured in low-glucose (5.5 mmol/L) DMEM (Hyclone, SH30021, USA) with 10% FBS (Bioind, 04-001-1A, USA), 100 µg/mL streptomycin and 100 U/mL penicillin (Gibco, 15140-122, USA). The cells were routinely cultured at 5% CO2 and 37°C with saturated humidity, digested and passaged after 85% cell confluence. At 24 hrs after cell passaging and attachment, the cells were divided into five groups and cultured for 48 hrs: 1) the normal-glucose group (NG) that received 5.5 mmol/L glucose (Sigma, A24940-01,USA); 2) the high-glucose group (HG) that received 30 mmol/L glucose with vehicle (DMSO); 3) the 0.01 µM MCC950 group (0.01) that received 30 mmol/L glucose with 0.01 µmol/L MCC950; 4) the 0.1 µM MCC950 group (0.1) that received 30 mmol/L glucose with 0.1 µmol/L MCC950 and 5) the 1 µM MCC950 group (1) that
received 30 mmol/L glucose with 1 μmol/L MCC950. We tested cell viability at 0 and 48 hrs by light microscopy and trypan blue exclusion.

Blood examination
Serum samples were used to measure serum creatinine and blood urea nitrogen (BUN) on a VITROS 950 automatic biochemical analyzer (Johnson & Johnson, New Brunswick, NJ).

ELISA assay
Mice were housed in individual metabolic cages, and 24-hr urine samples were collected every 4 weeks. Urinary albumin and neutrophil gelatinase-associated lipocalin (NGAL) were determined with a mouse ELISA kit (Cloud-Clone Corp, CEB028Mu, SEB388Mu, China) in accordance with the manufacturer’s instructions. Urinary creatinine concentrations were determined with a creatinine assay kit (NJJCBO, C011-2, China). Urinary albumin excretion was expressed as urinary albumin concentration versus creatinine concentration ratio (ACR, μg/mg). ELISA kits were also used for the measurement of levels of IL-1β in renal cortical tissue homogenates and cell culture supernatants (CUSABIO, E08054m, E08055r, China) in accordance with the manufacturer’s instructions.

Histologic analysis
Renal cortices were fixed in an alcohol-formalin-acetic acid solution (AFA) and were embedded in paraffin. The sections (3 μm) were stained with HE, periodic acid-Schiff (PAS) and Masson trichrome. The semi-quantitative index was used to evaluate the degree of glomerular sclerosis by PAS staining and tubulointerstitial damage by Masson staining, based on an average of 20 glomeruli per mouse. Glomerular sclerosis was scored from 0 to 5 as follows: 0, normal; 1, the mesangial matrix and slight glomerular damage and/or hyalinosis involving <10% of the glomerulus; 2, 10–20%; 3, 20–30%; 4, 30–40%; and 5, >40%. Tubular damage was scored by the percentage of injured tubules (tubular dilation, interstitial inflammation and fibrosis) from 0 to 5 as follows: 0, normal; 1, tubular lesion <10%; 2, 10–20%; 3, 20–30%; 4, 30–40%; and 5, >40%. Two pathologists performed a blinded analysis of all sections.

Transmission electron microscopy (TEM)
Renal cortices (1 mm³) were fixed with 2.5% glutaraldehyde at 4°C and were examined using a transmission electron microscope (H-7650, Olympus, Japan). Electron micrographs were used to determine the glomerular basement membrane (GBM) thickness and podocyte foot process width as previously described.23

Immunohistochemistry (IHC)
The deparaffinized and rehydrated sections (3 μm) of paraffin-embedded renal cortices were subjected to heat-mediated antigen retrieval and incubated with 3% H₂O₂ for 10 mins. Sections were then incubated overnight at 4°C with primary antibodies against proliferating cell nuclear antigen (PCNA) (Wanleibio, w102208, 1:200), phosphor-nuclear factor-κB p65 (p-NF-κB) (Wanleibio, w102169, 1:100), podocin (Abcam, ab181143, 1:500), fibronectin (Proteintech, 15613-1-AP, 1:200), α-SMA (Abcam, ab5694, 1:50), NLRP3 (Cell Signaling Technology, 15101, 1:200), caspase-1 (Abcam, ab1872, 1:25) or IL-1β (Cell Signaling Technology, 12242, 1:100). For PCNA, p-NF-κB and podocin, the percentage of a positively stained area over the whole glomerular area was calculated. For other antibodies, the micrographs were quantified blindly using Image-Pro Plus 6.0 software and were expressed as the ratio of integrated optical density to the whole area. Each renal section was counted for at least 10 images.

Western blot (WB) analysis
Total proteins from renal cortices were extracted with ice-cold RIPA lysis buffer (Beyotime, China), and protein concentrations were measured by BCA Protein Assay Kit (Beyotime, China). Equal amount of proteins was separated by SDS-PAGE and transferred to PVDF membranes (Millipore, USA). Membranes were blocked in 5% nonfat milk at room temperature for 1 hr and then incubated overnight at 4°C with primary antibodies to podocin (Abcam, ab181143, 1:2000), TGF-β1 (Proteintech, 18978, 1:1000), collagen I (Proteintech, 14695, 1:1000), α-SMA (Abcam, ab5694, 1:1000), NLRP3 (CST, 15101, 1:500), caspase-1 (Abcam, ab1872, 1:400), IL-1β (CST, 12242, 1:500) and β-actin (ZSGB-BIO, TA-09,1:3000). After incubation with the appropriate secondary antibodies (Abbkine, 1: 10,000) at room temperature for 1 hr, target bands were detected by enhanced chemiluminescence substrate (Abbkine, USA). Each Western blot was performed at least three times. Semi-quantitative analysis was measured using ImageJ software.

RNA extraction and real-time qPCR
Total RNA from renal cortices was extracted using TRIzol (Invitrogen, USA). Reverse transcription was performed...
using a Prime Script reverse transcription reagent kit (9108, TaKaRa Bio, Japan), and RNA (1 μg) was reverse transcribed into cDNA. Real-time qPCR was performed using TB Green reagent on an ABI 7500 real-time PCR System (RR820A, TaKaRa Bio, Japan) in accordance with the manufacturer’s instructions. In vivo experiments, the primer sequences of mice used for mRNA detection were as follows: TGF-β1, forward 5'-CAACAACTTCCGCTAACCTTG-3' and reverse 5'-GAAGCCTGTTAATTGCTCCTCTCTTG-3'; fibronectin, forward 5'-AGAGAGGAGGAGTCCCAAGAAG-3' and reverse 5'-GAAGGCGGCTGAAAGAGG-3'; Collagen I, forward 5'-TGCGGATCTCCAAGATGTG-3' and reverse 5'-CACACAAGGTTGCCTAGGTGA-3'; β-actin, forward 5'-GAGACCTTCAACCCCAGC-3' and reverse 5'-ATGTCAACGCAGATTCCC-3'. In vitro experiment, the primer sequences of cells were generated for mRNA detection: TGF-β1, forward 5'-ATGGTGAGCCGAAACAACGCTTG-3' and reverse 5'-GAGCCTGAAACCGCAGCGCCTTG-3'; fibronectin, forward 5'-ACGGGCCTGACCTAACCCAGC-3' and reverse 5'-GGGAAGTGCCACAAGGCCT-3'; Collagen I, forward 5'-AAACGGGAGGGCGAGTG-3' and reverse 5'-CATAGGACATCCTGCCCTGGCCT-3'. PCR was carried out at 95°C for 30 s, followed by 40 cycles each of 95°C for 5 s, 60°C for 30 s, and 72°C for 34 s. The relative quantitative expression was calculated by 2^{-ΔΔCt} method. Each group was examined in triplicate.

Immunofluorescence

The cells were fixed in 4% paraformaldehyde for 15 mins at room temperature after washing with PBS, permeabilized for 30 mins with 0.5% Triton X-100 in PBS and blocked for 1 h at 37°C with 5% BSA. The cells were incubated with caspase-1 (Abcam, ab1872, 1:25) overnight at 4°C in a humidified chamber. After washing with PBS, cells were incubated with secondary antibody (Abbkine, A23420, 1:500) for 1 hr at 37°C in a darkened humidified chamber. Cell nuclei were stained with DAPI. Fluorescence was observed using a confocal laser scanning microscope (Nikon A1R, Japan).

Statistical analysis

Quantitative data were presented as mean ± standard error of mean from three independent experiments. Statistical analysis was performed using GraphPad Prism 7.0 software. Differences between several groups were evaluated using ANOVA. Differences between two groups were assessed using the Student's t-test. A value of P<0.05 was regarded as statistically significant.

Results

MCC950 decreased the urinary albumin-to-creatinine ratio and improved renal function and pathological changes in db/db mice

BW and blood glucose were higher in db/db mice than in nondiabetic db/m mice; however, BW and blood glucose in db/db mice were not affected by MCC950 treatment (P>0.05, Figure 1A and B). The ratio of kidney weight to body weight (KW/BW) in db/db mice was lower than that of db/m mice; however, the KW/BW ratio showed no statistical difference between the db/db and db/db+MCC950 groups (P>0.05, Figure 1C). Serum creatinine of db/db mice was higher than that of db/m mice, and MCC950 significantly blunted the diabetes-induced increase in serum creatinine (P<0.05, Figure 1D). There was no significant difference in BUN between the db/m, db/db and db/db+MCC950 groups (P>0.05, Figure 1E). Compared to db/m mice, urinary albumin to creatinine ratio (ACR) of db/db mice was greater and was effectively reduced by MCC950 treatment (P<0.05, Figure 1F). The degree of decreasing urinary ACR increased with the prolongation of treatment time (F=13.95; P<0.01). Urinary NGAL as a marker of tubular injury was also reduced by MCC950 treatment in db/db mice (P<0.05, Figure 1G). HE, PAS and Masson’s staining showed mesangial matrix expansion, glomerular hypertrophy, tubular dilation and interstitial inflammation in db/db mice (Figure 1H). The glomerular sclerosis index and tubulointerstitial damage index were higher in db/db mice than in db/m mice (P<0.05, Figure 1I and J). However, both were significantly attenuated by MCC950 treatment, with MCC950 ameliorating the glomerular sclerosis and tubular damage of DN. Expression levels of PCNA and p-NF-κB were greater in the nucleus of glomeruli in db/db mice than in db/m mice, which confirmed that mesangial cells were proliferated and activated, and were inhibited by MCC950 treatment as measured by IHC (P<0.05, Figure 1K-N).

MCC950 attenuated podocyte injury in db/db mice

Electron microscope images showed that the increased GBM thickness and foot process width were significantly attenuated by MCC950 treatment (P<0.05, Figure 2A–C). Lower expression of podocin as a marker of podocyte in renal cortical tissue...
of db/db mice was increased by MCC950 treatment with IHC and WB (P<0.05, Figure 2D–G). MCC950 prevented diabetes-induced podocytes injury.

**MCC950 alleviated renal fibrosis in db/db mice**

Expression levels of fibronectin and α-smooth muscle actin (α-SMA) (both of which are markers of fibrosis) were greater in db/db mice than in db/m mice, and were inhibited by MCC950 treatment as measured by IHC (P<0.01, Figure 3A–D). MCC950 treatment also suppressed the increase in fibrosis marker protein levels caused by TGF-β1, collagen I and α-SMA in db/db mice, as measured by WB (P<0.05, Figure 3E–H). Furthermore, mRNA expression levels of TGF-β1, fibronectin and collagen I were significantly decreased by MCC950 treatment in db/db mice (P<0.05, Figure 3I–K). MCC950 effectively prevented the progression of renal fibrosis in db/db mice.

**MCC950 inhibited diabetes-induced renal NLRP3 inflammasome activation in db/db mice**

The renal expressions of NLRP3, caspase-1 and IL-1β were higher in db/db mice than in db/m mice as shown by IHC and were reduced by MCC950 treatment (P<0.01, Figure 4A–F). MCC950 also decreased NLRP3 protein expression in db/db mice as measured by WB (P<0.05, Figure 4G–H). The activities of caspase-1 and IL-1β were greater in db/db mice and were inhibited by MCC950.
treatment. The activities of caspase-1 and IL-1β were determined by WB according to their protein expression levels from the immature form to active form (P<0.05, Figure 4G, I and J). Higher expression of IL-1β in renal cortical tissue of db/db mice was decreased by MCC950 treatment by ELISA (F=6.331, P<0.05, Figure 4K). Therefore, MCC950 inhibited NLRP3 inflammasome activation and the activities of caspase-1 and IL-1β.

MCC950 inhibited HG-induced NLRP3 inflammasome activation in mesangial cells  
Mesangial cells showed higher NLRP3 protein expression after 48-hr stimulation with HG, as well as higher levels of mature caspase-1 and IL-1β than in the NG group, that were inhibited by 0.1 μM or 1 μM MCC950 treatment. MCC950 inhibited NLRP3 inflammasome activation, accordingly disturbing the transformation of pro-caspase-1 into mature caspase-1 and the production of mature IL-1β p17 (P<0.05, Figure 5A–D). Immunofluorescent images showed higher caspase-1 expression in the HG group that was diminished by 0.1 μM or 1 μM MCC950 treatment (P<0.05, Figure 5E). Higher expression of IL-1β in the supernatant of the HG group was decreased by 0.1 μM or 1 μM MCC950 treatment by ELISA (F=7.218, P<0.05, Figure 5F).

MCC950 decreased expression of fibrosis markers by inhibiting NLRP3 inflammasome activation in mesangial cells  
Protein levels of fibrosis markers TGF-β1, collagen I and α-SMA were higher in the HG group, as demonstrated on
WB; these levels were significantly inhibited by 0.1 μM or 1 μM MCC950 treatment (P<0.05, Figure 6A–D). The similar results were obtained by RT-qPCR. MCC950 prevented HG-induced increases mRNA expression levels of TGF-β1, collagen I and fibronectin in mesangial cells (P<0.05, Figure 6E–G).

Discussion

Chronic low-grade sterile inflammation and proinflammatory cytokines release are thought to be crucial mechanisms in the progression of DN. 24,25 In DN, the NLRP3 inflammasome is an intracellular platform that converts pro-caspase-1 and pro-IL-1β into active forms (caspase-1 p10/p20 and IL-1β p17) responding to danger signals and triggers inflammatory programmed cell death called pyroptosis. 26,27 Pyroptosis features cell swelling and subsequently rupturing the cell membrane, causing massive leakage of cytosolic contents, further promoting killing by a secondary phagocyte, contributing to the expansion of the inflammatory response. 28–30 MCC950 is a selective small molecule inhibitor of NLRP3 that blocks canonical and non-canonical NLRP3 activation at nanomolar concentrations.13 In recent studies, MCC950 reduced liver inflammation and fibrosis by suppression collagen I, α-SMA and hepatic connective tissue growth factor expression in a mouse model of non-alcoholic steatohepatitis.31 MCC950 also exerted protective effects in cholestatic liver injury and liver fibrosis by suppression hepatic expression of the pro-fibrotic markers (TGF-β1, α-SMA and Col1a1) in a mouse model of bile duct ligation.32 MCC950 protected human retinal endothelial cells against

Figure 3 MCC950 alleviated renal fibrosis in db/db mice.

Notes: (A–D) Representative immunohistochemical and immunohistochemical scores of renal sections for fibronectin and α-SMA staining at 20 weeks of age. Bar =50 μm. (E–H) Western blot analysis of TGF-β1, fibronectin, collagen I and α-SMA in renal cortical tissue. (I–K) Relative mRNA expression levels of TGF-β1, fibronectin and collagen I in renal cortical tissue by real-time qPCR. Data represent means ± SEM (n=10). *P<0.01 versus db/m mice; **P<0.05 and ***P<0.01 versus db/db mice.

Abbreviations: α-SMA, α-smooth muscle actin; PCR, polymerase chain reaction; SEM, standard error of mean.
HG-induced dysfunction. MCC950 ameliorated proteinuria, pathological changes and podocyte foot process effacement in lupus-prone mice. Based on these studies, we proposed a model of the possible mechanism of action of MCC950 in DN (Figure 7). In our study, we used MCC950 to intervene in type 2 diabetic db/db mice in vivo and HG-induced mesangial cells in vitro. The efficacy of MCC950 for the treatment of DN in db/db mice has not been reported previously. MCC950 effectively inhibited expression of NLRP3 and the conversion to active forms of caspase-1 and IL-1β in both renal cortices of db/db mice and in HG-induced mesangial cells. MCC950 suppressed renal inflammation by inhibiting NLRP3 inflammasome activation, the mechanism of by which MCC950 ameliorated kidney injury in DN.

Albuminuria is the most important clinical feature of DN, primarily caused by an impaired glomerular filtration barrier. Urinary ACR is the most important marker for evaluating early renal decline and diagnostic basis in diabetic nephropathy. Podocytes are the weakest link in the glomerular filtration barrier. Podocyte damage gives rise to albuminuria in DN and NLRP3 inflammasome activation in podocytes contributes to podocyte loss. Podocytes are highly differentiated epithelial cells. Podocyte damage leads to irreversible renal decline. In this study, MCC950 treatment ameliorated podocyte foot process effacement and thickened GBM in db/db mice. Podocin is a glomerular slit diaphragm protein between neighboring podocytes, and a critical component of glomerular filtration barrier, that plays
MCC950 inhibited HG-induced NLRP3 inflammasome activation in mesangial cells.

**Notes:** (A–D) Western blot analysis of NLRP3 (F=3.499), caspase-1 (F=3.508) and IL-1β (F=4.671) in mesangial cells. (E) Immunofluorescent localization of caspase-1 (red) in mesangial cells with nuclei staining by DAPI (blue). Bar = 50 µm. (F) Expression of IL-1β in cell culture supernatants by ELISA (F=7.218). Data represent means ± SEM. *P<0.05 and **P<0.01 versus NG; *P<0.05 and ***P<0.01 versus HG.

**Abbreviations:** NLRP3, NOD-like receptor pyrin domain-containing protein 3; NG, normal-glucose group; HG, high-glucose group; SEM, standard error of mean.

a key role in maintaining the normal structure and function of podocytes. Decreased expression of podocin suggests that podocyte foot processes fuse, destroying the glomerular filtration barrier, resulting in production of albuminuria. In our study, the expression of podocin was markedly elevated by MCC950 treatment in db/db mice. Therefore, MCC950 protected against podocyte injury to reduce urinary ACR and improved renal function in db/db mice.

The typical pathological features of early DN are glomerular hypertrophy, thickened glomerular basement membrane and mesangial expansion with increased matrix deposition. Renal fibrosis of both glomerular and tubulointerstitial compartments, characterized by extracellular matrix (ECM) accumulation, is recognized as a basic mechanism leading to diabetic glomerular disease. Mesangial cells as intrinsic renal cells secrete a large number of proinflammatory and profibrotic cytokines responding to injury in DN. IL-1β has been demonstrated to stimulate production of TGF-β1, fibronectin, collagen I and mesangial proliferation. The positive PCNA and p-NF-κB immunohistochemistry staining also confirmed the proliferation and activation of mesangial cells in db/db mice. Therefore, we chose mesangial cells in vitro to study the mechanism of MCC950 in DN. TGF-β1 is a major cytokine secreted by mesangial cells that mediates the development of DN. It has many biological activities such as regulating cell proliferation, differentiation and migration. TGF-β1 is also a key cytokine mediating collagen deposition in kidney, including promoting the production of ECM, inhibiting the degradation of ECM and participating in renal fibrosis. ECM in patients with DN is produced by mesangial cells and is primarily composed of fibronectin, collagen IV and a few collagen I. The expression of collagen I is low level in normal kidney tissue; however, it is significantly increased under high glucose stimulation. The expression of α-SMA in normal mesangial cells is very weak. However, the
expression of α-SMA in mesangial cells is increased under high glucose stimulation, suggesting that mesangial cells undergo phenotypic transformation from resting state to secretion/proliferative state. In our study, MCC950 significantly decreased mesangial cells proliferation and the expression levels of fibrosis markers (TGF-β1, fibronectin, collagen I and α-SMA) both in renal cortices of db/db mice and in HG-induced mesangial cells.

Urinary NGAL as a marker of renal tubular injury is also an important marker for evaluating early progressive renal decline in type 2 diabetes mellitus. Inflammatory cell infiltration in the tubules and interstitium is also one of
the pathological changes of DN, leading to renal fibrosis. In glomerular diseases, the development of glomerular inflammation precedes interstitial fibrosis.41 Furthermore, NLRP3 inflammasome activation also occurs in intrinsic renal cells such as tubular epithelial cells.11,12 Knockdown of NLRP3 alleviated high glucose-induced epithelial-to-mesenchymal transition in human renal tubular cells.46 MCC950 inhibited NLRP3 inflammasome activation in tubular epithelial cells of db/db mice. These also can explain how MCC950 significantly decreased urinary NGAL as a marker of renal tubular injury and fibrosis markers in db/db mice. MCC950 also ameliorated pathologic changes, including glomerular sclerosis and tubular damage in db/db mice. These data suggested that MCC950 prevents renal fibrosis from renal decline by inhibiting NLRP3 inflammasome activation in DN. Nevertheless, one limitation of this study is that we only used mesangial cells in vitro to study the mechanism of MCC950. In the future, we will further investigate the role of MCC950 in podocytes. However, there are always some differences between mice model and human, and so, this efficacy of MCC950 in DN requires further clinical investigation.

We found that MCC950 treatment improved renal function and pathologic changes in db/db mice without affecting BW and blood glucose levels. H.L. Kammoun et al confirmed the same results that MCC950 did not impact BW and blood glucose levels in db/db mice.47 Inhibition of the NLRP3 inflammasome by MCC950 ameliorated kidney injury in DN; the renoprotection of MCC950 for DN occurred independently of BW and blood glucose levels.

**Conclusion**

The data presented in this study suggested that MCC950 effectively ameliorates kidney injury in DN, independent of BW and blood glucose levels. MCC950 inhibited NLRP3 inflammasome activation to reduce the production of active caspase-1 and IL-1β. These phenomena might result in reducing urinary ACR and urinary NGAL, improving renal function and alleviating podocyte injury and renal fibrosis. Therefore, MCC950 may be a promising therapeutic strategy to prevent the progression of DN.

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**Disclosure**

The authors report no conflicts of interest in this work.

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