Piperazine ferulate prevents high-glucose-induced filtration barrier injury of glomerular endothelial cells

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Abstract. Filtration barrier injury induced by high glucose (HG) levels leads to the development of diabetic nephropathy. The endothelial glycocalyx plays a critical role in glomerular barrier function. In the present study, the effects of pipera‑zine ferulate (PF) on HG‑induced filtration barrier injury of glomerular endothelial cells (GEnCs) were investigated and the underlying mechanism was assessed. Immunofluorescence was used to observe the distribution of the glycocalyx as well as the expression levels of syndecan‑1 and Zonula occludens‑1 (ZO‑1). Endothelial permeability assays were performed to assess the effects of PF on the integrity of the filtration barrier. Protein and mRNA expression levels were measured by western blotting and reverse transcription‑quantitative PCR analyses, respectively. In vitro experiments revealed that adenosine monophosphate‑activated protein kinase (AMPK) mediated HG‑induced glycocalyx degradation and endothelial barrier injury. PF inhibited the HG‑induced endothelial barrier injury and restored the expression levels of heparanase‑1 (Hpa‑1), ZO‑1 and occludin‑1 by AMPK. In vivo assays demonstrated that PF reduced the expression levels of Hpa‑1, increased the expression levels of ZO‑1 and attenuated glycocalyx degradation in the glomerulus. These data suggested that PF attenuated HG‑induced filtration barrier injury of GEnC by regulating AMPK expression.

Introduction

Diabetic nephropathy (DN) is associated with damage to the glomerular filtration barrier (GFB), which consists of the three following layers: The endothelial layer, the glomerular basement membrane and the podocytes (1,2). GFB is responsible for filtering water and small molecules from circulating plasma. Accumulating evidence has demonstrated that glomerular endothelial cells (GEnCs) act as the first barrier to prevent macromolecules from passing the GFB (2,3). Glycocalyx is the main component of the endothelial surface layer, which constitutes a molecular size- and charge-specific selective barrier together with the fenestrations (4,5). The degradation of the glycocalyx and the decreased number of fenestrations are the most important characteristics of filtration barrier injury to GEnCs during DN (6). Since glycocalyx dysfunction exerts an important role in the development of DN, the prevention of glycocalyx degradation may be an alternative approach to treat DN.

Adenosine monophosphate‑activated protein kinase (AMPK) is a ubiquitous heterotrimeric protein composed of a catalytic subunit (α) and two regulatory subunits (β and γ) (7). A previous study demonstrated that AMPK is an important molecule that regulates the progression of DN (8). Renal hypertrophy (9) and podocyte apoptosis (10) in diabetes are regulated by AMPK, whereas metformin, an activator of AMPK, has been demonstrated to decrease albuminuria in diabetic rats as well as in patients with type 2 diabetes mellitus (11,12). Furthermore, AMPK has been shown to mediate low shear stress‑induced glycocalyx impairment (13). This evidence indicates that AMPK may be a possible drug target to prevent glomerular filtration injury.

Piperazine ferulate (Piperazine 3-methoxy-4-hydroxy-cinnamate; PF; Fig. 1A) is a derivative of ferulic acid that exhibits anti‑hypertensive effects (14), reduces IgA‑mediated nephropathy (15) and prevents the development of DN (16). A systematic meta‑analysis indicated that PF combined with irbesartan could improve the efficiency of treatment...
of DN (17). Our previous study also demonstrated that PF retarded the progression of DN (16) and attenuated the high glucose (HG)-induced mesangial cell injury (18). However, it remains unclear whether the mechanism of PF against DN is associated with the regulation of GEnC filtration barrier injury. In the present study, the effects of PF on the HG-induced filtration barrier injury of GEnCs were investigated. Moreover, the potential mechanism of its action was assessed.

Materials and methods

**Materials.** PF was provided by Hunan QianJinXiangJiang Pharmaceutical Industry Co., Ltd. Fluorescein isothiocyanate (FITC)-dextran and FITC conjugated wheat germ agglutinin (FITC-WGA) were obtained from Sigma-Aldrich (Merck KGaA). Rabbit polyclonal anti-occludin-1 (cat. no. DF7504) and anti-heparanase (Hpa-1) antibodies (cat. no. DF12411) were purchased from Affinity Biosciences. Rabbit polyclonal anti-AMPK (cat. no. 2532) and anti-phosphorylated (p)-AMPKα (Thr172) antibodies (cat. no. 2535) were obtained from Cell Signaling Technology, Inc. Rabbit polyclonal anti-Zonula occludens-1 (ZO-1; cat. no. PB9234) and mouse monoclonal anti-β-actin antibody (cat. no. BM0627) as well as the syndecan-1 (Sdc-1) ELISA kits (cat. no. EK1339 and EK1554) were purchased from Boster Biological Technology, Ltd. SYBR® Premix Ex Taq™ and PrimeScript reverse transcription reagent kit were obtained from Takara Bio, Inc. TRizol® was obtained from Thermo Fisher Scientific, Inc. RIPA lysis buffer, 5-aminoimidazole-4-carboxamide riboside (AICAR), the bicinonic acid (BCA) assay kit and a BeyoECL Plus Western blot analysis kit were obtained from Beyotime Institute of Biotechnology. Glucose and mannitol were purchased from Guoyao Chemical Reagent Co. Compound c was obtained from Selleck Chemicals.

**Cell culture and treatment.** Human GEnCs were purchased from the ScienCell Research Laboratories, Inc. and cultured under standard cell culture conditions at 37°C with 5% CO₂ in a humidified incubator using endothelial cell medium (ECM; Shanghai Zhong Qiao Xin Zhou Biotechnology Co., Ltd.), supplemented with 100 U/ml penicillin, 100 µg/ml streptomycin, 10% (v/v) FBS and 1% endothelial cell growth factor (Zhongqiaoxinzhou Biotech). Cells between passages 3 and 8 were used for the experiments. To study the role of AMPK in the protective effect of PF on HG-induced GEnCs injury, GEnCs were preincubated with 20 µM Compound c for 1 h and subsequently treated with HG (30 mM) in the presence or absence of PF for 48 h. Next, the permeability of GEnCs to FITC-dextran was determined.

**Endothelial permeability assays.** FITC-dextran was used to measure permeability of GEnC monolayers. GEnC monolayers were cultured on the apical chamber of the Transwell inserts (0.4 µm pore size; Corning, Inc.) until they reached 100% confluence and were subsequently treated with 5.5 mM glucose (control), 30 mM HG or mannitol (5.5 mM glucose + 24.5 mM mannitol) with or without PF (25-200 µM) for the indicated time periods. Dextran was dissolved in ECM and added to the apical chamber at a concentration of 2.0 g/l following different experimental treatments. Following 1 h of incubation, the amount of fluorescence of FITC-dextran in the basolateral chamber was measured using fluorescence spectrophotometry with an excitation and emission wavelength of 490 and 520 nm, respectively. The results are presented as fold-change following normalization to the control group.

**Small interfering RNA (siRNA)-mediated knockdown.** Specific and negative control siRNAs were purchased from Guangzhou Ribobio Co., Ltd. siRNA-AMPK (40 nM; cat. no. siB1456165958-1-5) was transfected into GEnCs using Dharma FECT transfection reagent (Thermo Fisher Scientific, Inc.) according to the manufacturer's protocol. The negative control group were GEnCs transfected with scrambled RNA (40 nM, cat. no. siN0000001-1-5). Following transfection, the cells were incubated for 24 h and used for further experiments.

**Reverse transcription-quantitative PCR (RT-qPCR).** Total RNA was extracted from renal tissues using TRIzol® reagent and cDNA was synthesized using a PrimeScript reverse transcription reagent kit according to the manufacturer's protocol. RT-qPCR was performed using a LightCycler96® (Roche Diagnostics GmbH) and SYBR-Green master mix. The RT-qPCR cycling conditions were as follows: Initial denaturation at 95°C for 30 sec followed by 45 cycles of denaturation at 95°C for 5 sec and annealing 60°C for 30 sec. The species-specific primer sequences were as follows: Mouse occludin-1 forward, 5’-CTACGGAGGTGCTATGGG-3’ and reverse, 5’-AAGGAAGCGTGAAACAGA-3’; mouse ZO-1 forward, 5’-ATGACTCTGACGTTGGTC-3’ and reverse, 5’-GGCTCAAACAGGTAATTGC-3’; mouse β-actin forward, 5’-ACTGCTCTGCTCTAGC-3’ and reverse, 5’-ACATCTGCTGAAGTGAC-3’. Gene expression was calculated using the 2^ΔΔCq method (19). The data are expressed as the fold-change normalized to β-actin.

**Western blot analysis.** Total protein was extracted from cells using RIPA lysis buffer and the protein concentration was determined using the BCA method according to the manufacturer's protocol. A total of 35 µg protein/lane was loaded onto a SDS-PAGE (10%) and separated at 120 V. The proteins were transferred to PVDF membranes and blocked in 5% non-fat milk for 1 h at room temperature. Subsequently, the blots were incubated with primary antibodies against ZO-1 (1:500), occludin-1 (1:500), AMPK (1:500), hpa-1 (1:500), p-AMPKα (Thr172, 1:300) and β-actin (1:500) at 4°C overnight. The blots were incubated with HRP-conjugated goat anti-mouse (Boster Biological Technology; cat. no. BA1050) or HRP-conjugated goat anti-rabbit (Boster Biological Technology; cat. no. BA1054) diluted with secondary antibody dilution buffer (Beyotime Institute of Biotechnology, cat. no. P0023D) for 1 h at room temperature. The bands were visualized using a BeyoECL Plus kit. Densitometry analysis was performed using ImageJ version 1.45s (National Institutes of Health). The protein expression levels were normalized to those of β-actin.

**Animal model.** The animal experimental protocol was approved by the Ethics Committee of Animal Experiments of the Central South University and was performed in accordance with the Guidelines for the Care and Use of Laboratory Animals.
Animals (20). Male C57BL/6J mice, aged 8 weeks (weight, 19-22 g), were purchased from the Experimental Animal Center of SiLaiKeJingDa. The animals were maintained under a constant 12 h light/dark cycle at 22±2˚C and allowed free access to food and water at least for 1 week. Type 1 diabetic mice were established using a streptozotocin (STZ) injection as previously described (16). Briefly, the mice were intraperitoneally injected for 5 consecutive days with STZ (65 mg/kg body weight). After 2 days, the levels of blood glucose were measured by the mouse tail snipping method using a glucometer (Sannuo Biotech Ltd.). Mice with blood glucose levels ≥16.7 mmol/l were considered diabetic. Normal control mice were injected with citrate buffer solution (0.1 M, pH 4.0). Then, 1 week after blood glucose stabilization, diabetic mice were divided into the two following groups (n=15 per group): The diabetic mice group were treated with vehicle (DM) and the PF group of mice treated with PF (100 mg/kg, DM+PF). PF was provided by intragastric administration daily for 12 weeks. All mice were anaesthetized using sodium pentobarbital (50 mg/kg body weight). Normal control mice were injected with citrate buffer solution (0.1 M, pH 4.0). Then, 1 week after blood glucose stabilization, diabetic mice were divided into the two following groups (n=15 per group): The diabetic mice group were treated with vehicle (DM) and the PF group of mice treated with PF (100 mg/kg, DM+PF). PF was provided by intragastric administration daily for 12 weeks. All mice were anaesthetized using sodium pentobarbital (50 mg/kg body weight) and blood samples (1-1.2 ml) were collected from the retro-orbital sinus after eyeball removal. Finally, the mice were sacrificed by exsanguination under anesthesia (50 mg/kg sodium pentobarbital) and the kidney tissues were collected. Collected blood was left to stand at room temperature for 1 h, and then centrifuged at 2,000 x g at 4˚C for 15 min. The supernatant was transferred to a new tube and stored at -80˚C.

Immunofluorescence analysis. Renal tissues were fixed in 4% paraformaldehyde at 4˚C overnight, embedded in paraffin and cut into 5 µm-thick slices. The slices were processed by dewaxing, gradient alcohol dehydration, antigen repair with 10 mM sodium citrate (pH 6.0, microwaved for 10 min) and finally washed in PBS three times. The sections were incubated with FITC-WGA (10 µg/ml) at room temperature for 2 h, stained with DAPI at room temperature for 10 min and examined using fluorescence microscopy (Zeiss AG; magnification, x400). Immunofluorescence was performed by incubating the slides with ZO-1 (1:200)/CD31 (1:200) or Sdc-1 (1:200)/CD31 (1:200) overnight at 4˚C. The slides were incubated with the corresponding anti-FITC-conjugated secondary antibody (Beyotime Institute of Biotechnology; cat. no. A0562; 1:500) or Cy3-conjugated secondary antibody (Beyotime Institute of Biotechnology; cat. no. A0516; 1:500) diluted with secondary antibody dilution buffer for 1 h at room temperature in the dark. The cell nuclei were stained with DAPI at room temperature for 5 min. Finally, the images were captured using fluorescence microscopy (magnification, x400).

Transmission electron microscopy. The kidney tissues were fixed in 2.5% glutaraldehyde at 4˚C overnight and post-fixed in 1% osmium tetroxide at 4˚C for 2 h. Following dehydration, the samples were embedded in EMBed 812 resin (Thermo Fisher Scientific, Inc.) and polymerized at 60˚C for 48 h. The
ultrathin sections (60-80 nm) were cut and stained with 1% uranyl acetate at room temperature for 10 min, followed by 2% lead citrate buffer for 2 min at 37°C. The structure of the glomerulus was observed using a Hitachi High-Tech 7700 electron microscope (magnification, x5,000).

**Determination of the expression levels of soluble Sdc-1.** The levels of soluble Sdc-1 in the cell culture supernatant or serum were measured using the Sdc-1 ELISA kits according to the manufacturer’s protocols.

**Determination of glycocalyx via endothelial surface analysis.** Following treatment, GEnCs were washed with PBS and fixed in 4% paraformaldehyde at room temperature for 10 min. Subsequently, the cells were incubated with FITC-WGA (10 µg/ml) at room temperature for 30 min. Finally, the images were captured using fluorescence microscopy (magnification, x400).

**Statistical analysis.** Data are presented as the mean ± standard error of at least three independent experiments. Data were analyzed using SPSS software (version 17.0; SPSS, Inc.). A one-way ANOVA followed by a post-hoc Tukey’s test was performed to compare difference between multiple groups. P<0.05 was considered to indicate a statistically significant difference.

**Results**

**AMPK mediates HG-induced filtration barrier injury of GEnCs.** The effects of HG on the permeability of the GEnC barrier to FITC-dextran were examined. HG (30 mM) increased the permeability of FITC-dextran after 48 h, whereas mannitol did not have an effect for 48 h (Fig. 1B). Subsequently, the effects of HG on the expression levels of AMPK and p-AMPKα (Thr-172) were measured by western blot analysis. GEnCs treated with HG exhibited decreased levels of p-AMPKα (Thr-172), although the total expression levels of AMPK were not altered notably (Fig. 1C and D). The effects of AMPK on the permeability of GEnC to FITC-dextran were further examined using AICAR, a selective agonist of AMPK. As shown in Fig. 1E, GEnCs pretreated with AICAR (1 mM) decreased the HG-induced permeability of GEnC to FITC-dextran. Similarly, the results of FITC-WGA staining demonstrated that AICAR significantly inhibited the decrease of the glycocalyx content in GEnC induced by HG (Fig. 1F). These data suggested that AMPK mediated HG-induced filtration barrier injury of GEnCs.

**AMPK regulates the expression of Hpa-1 and tight junction (TJ) proteins.** Sdc-1 is the main constituent of the glycocalyx. This protein can be cleaved by Hpa-1, an enzyme capable of degrading polymeric heparan sulfate molecules of Sdc-1 into shorter chain length oligosaccharides (21). HG increased the expression levels of Hpa-1 without affecting the expression levels of Sdc-1 in GEnCs (Fig. 2A). Therefore, the expression levels of Sdc-1 in the cell culture supernatant were determined. The results demonstrated that the expression levels of Sdc-1 were increased in the HG group. These changes were significantly inhibited by co-incubation of the cells with AICAR (Fig. 2A-D). Furthermore, the effects of AICAR on the expression of TJ proteins, ZO-1 and occludin-1 were investigated. AICAR significantly inhibited the HG-induced downregulation of the expression of occludin-1 and ZO-1 at the protein level (Fig. 2A, E and F). These data suggested that AMPK mediated the HG-induced filtration barrier injury of GEnCs by regulation of the expression levels of Hpa-1 and TJ proteins.

**PF attenuates HG-induced filtration barrier damage.** The effects of PF on the permeability defects of GEnCs...
induced by HG were assessed. Incubation of GEnCs with PF (100 and 200 µM) significantly inhibited the increase noted in the HG-induced permeability of GEnCs to FITC-dextran (Fig. 3A). The specific dose of PF used for further investigations was 100 µM. Preincubation of GEnCs with PF attenuated the increase of soluble Sdc-1 expression levels in the cell culture supernatant (Fig. 3B). Moreover, it increased the glyco- claxyx content of GEnCs (Fig. 3C and D). PF ameliorated the HG-induced increase in the expression levels of Hpa-1 protein in GEnCs (Fig. 3E and F). Similarly, PF inhibited the decrease in the expression levels of ZO-1 and occludin-1 in GEnCs exposed to HG (Fig. 3E, G and H). GEnCs incubated with PF exhibited a significant increase in the p-AMPK/AMPK ratio compared with those noted in the HG group (Fig. 3E and I). These finding suggested that PF inhibited HG-induced filtration barrier injury in GEnCs.

**PF alleviates HG-induced filtration barrier injury via AMPK.**

The regulatory mechanism of PF on HG-induced filtration barrier injury was further investigated. GEnCs were preincubated with Compound c (AMPK inhibitor, 20 µM) for 1 h and subsequently treated with HG (30 mM) in the presence or absence of PF for 48 h. Compound c abolished the protective effects of PF on the HG-induced increased of the permeability of GEnCs to FITC-dextran and increased the Sdc-1 levels in the cell culture supernatant (Fig. 4A and B). Following knockdown of AMPK in GEnCs by siRNA (Fig. S1), the effects of PF on the HG-induced upregulation of Hpa-1 expression and the downregulation of ZO-1 and occludin-1 were inhibited (Fig. 4C-F). Taken together, the data revealed that AMPK mediated the protective effects of PF on the HG-induced filtration barrier injury.

**PF alleviates endothelial glycocalyx injury in vivo.**

*In vitro* experiments demonstrated that diabetic mice treated with PF for 12 weeks exhibited an increase in the number of GEnC fenestrations compared with that of the model group (Fig. 5A and B). Immunofluorescence analysis indicated that the glycocalyx content was increased in the glomerular cells of the PF-treated group compared with that of diabetic mice (Fig. 5C and D). Immunohistochemical analysis indicated that PF significantly increased p-AMPKα (Thr-172) protein expression (Fig. 5E and F) and inhibited the increase in Hpa-1 protein levels in the glomerular cells of diabetic mice (Fig. 5G and H). Immunofluorescence analysis indicated that the expression levels of Sdc-1 (Fig. 6A) and ZO-1 (Fig. 6B) in GEnCs were increased in PF-treated mice compared with those noted in diabetic mice. In addition, treatment of diabetic mice with PF inhibited the increase noted in the expression levels of Sdc-1 in the serum.
YANG et al.: PIPERAZINE FERULATE PREVENTS FILTRATION BARRIER INJURY

(6 Fig. 6C). It concomitantly increased the expression levels of ZO-1 and occludin-1 in kidney tissues (Fig. 6D and E). Taken together, these finding indicated that PF reduced GFB damage.

Discussion

Endothelial barrier injury is characterized by increased endothelial permeability and is associated with acute and chronic nephropathy (22). Emerging evidence has revealed that damage to the GEnC barrier plays an important role in the onset and progression of DN development (4). Our previous study demonstrated that PF attenuated the levels of 24 h-albuminuria, blood urea nitrogen and serum creatinine in STZ-induced DN mice (16), and it also reduced HG-induced mesangial cell injury (18). The results of the present study indicated that PF exerted a protective effect on GFB injury. The effects of PF were investigated on HG-induced filtration barrier injury of GEnCs, and the results suggested that this compound could reduce the damage to the glycocalyx degradation noted in GEnCs. These findings highlight a novel mechanism by which PF retards the progression of DN. This mechanism of action may restore endothelial filtration barrier injury.

Endothelial glycocalyx maintains the vascular function and plays an important role in vascular permeability, leucocyte, platelet adhesion as well as transduction of fluid shearing...
forces (23,24). FITC-WGA has been shown to bind to the glyco-
protein of the glycocalyx (25). This marker was used to evaluate
the effects of PF on glomerular endothelial glycocalyx. The data
indicated that PF prevented HG-induced glycocalyx degrada-
tion in vivo and in vitro. Hpa-1 is an endo-β-D-glucuronidase
that releases 5-7 kDa fragments of heparan sulfate from intact
heparan sulfate chains of proteoglycans (26). In addition, Hpa-1
regulates the expression levels and turnover of syndecans (27).
The results of the present study showed that HG increased the
expression levels of Hpa-1 in vitro and in vivo. PF attenuated
the increased expression of Hpa-1 in HG-induced GEnCs as
well as in the glomeruli of diabetic mice. However, the expres-
sion levels of Sdc-1 did not decrease in HG-stimulated GEnCs.
The effect of HG on the expression of Sdc-1 is similar to those
of Singh et al (28). Immunofluorescence analysis indicated that
Sdc-1 levels in the glomeruli of diabetic mice were decreased,
whereas PF inhibited these changes. The possible reason for
the discrepancy noted between the in vivo and in vitro experi-
ments is the inconsistent exposure factors used for the different
models. The degradation of glycocalyx in vivo may be affected
by blood glucose levels, inflammatory factors and the course of
diabetes (29). The exposure of GEnCs to HG involved a short
term period (24 h), while the course of diabetes was completed
following 12 weeks.

In the present study, the effects of PF on the TJ of GEnCs
were also evaluated. The expression levels of ZO-1 were
decreased in GEnCs of STZ-induced diabetic mice and in
HG-treated GEnCs, respectively. PF treatment increased
the expression levels of ZO-1. TJs between endothelial cells
regulate the passage of ions and small molecules through the
paracellular pathway and are involved in the pathogenesis of
DN (30). The integrity of glomerular GFB is also maintained
by TJ proteins that consist of at least 40 different proteins
including transmembrane proteins, cytoplasm accessory ZO
Figure 6. PF restores GFB injury. (A) Immunofluorescence images of Sdc-1 in the glomerulus. Magnification, x400. (B) Immunofluorescence image of ZO-1 in the glomerulus. Magnification, x400. (C) Expression levels of Sdc-1 in the serum. (D) ZO-1 mRNA expression levels in the kidney tissues. (E) Occludin-1 mRNA expression levels in the kidney tissues. Data are presented as the mean ± standard deviation of three repeats. **P<0.01, ***P<0.001 vs. the control group; #P<0.05, ##P<0.01 vs. model group. PF, piperazine ferulate; GFB, glomerular filtration barrier injury; Sdc-1, syndecan-1; ZO-1, Zonula occludens-1.

Figure 7. Schematic of pathways involved in the protective effects of PF on filtration barrier injury of GEnCs in diabetic mice. HG induces glycocalyx degradation in GEnCs and TJ damage between GEnCs in diabetic mice. PF inhibits glycocalyx degradation by regulation of AMPK. PF, piperazine ferulate; GEnCs, glomerular endothelial cells; HG, high glucose; TJ, tight junction; AMPK, adenosine monophosphate-activated protein kinase.

proteins and cytoskeletal proteins (30). HG influences the expression levels and/or redistribution of ZO-1, which leads to TJ destruction and filtration barrier injury (31,32). Thus, cell junctions may serve as a therapeutic target of DN.
AMPK is an important energy-sensing enzyme that controls cellular energy metabolism and is considered a potential pharmacological target for the treatment of DN (33,34). Phosphorylation of the α subunit at Thr-172 is essential for AMPK activity (35). Pharmacological activation of AMPK by AICAR results in restoration of the permeability of podocytes to albumin (36). It also increases the epithelial TJ assembly following calcium switch (37). In addition, ampkione is an AMPK activator, which can be used to attenuate endothelial glycocalyx impairment induced by low shear stress (13). The results of the present study demonstrated that AMPK mediated HG-induced glycocalyx impairment and TJ injury in GEnCs. Based on western bloting, p-AMPK expression was decreased under HG conditions; however AMPK expression was not altered; whereas, PF increased the phosphorylation of AMPK following HG treatment. It is worth noting that treatment of GEnCs with the AMPK siRNA or inhibitor (compound c) abolished the protective effects of PF on the HG-induced filtration barrier injury. Specific molecular mechanisms may be involved in the phosphorylation of AMPK regulated by PF. A previous report indicated that ferulic acid, which is one of the components of PF, regulated the phosphorylation of liver kinase B1 and AMPK in C2C12 myoblasts via activation of sirtuin 1 (Sirt1) (38). An additional study demonstrated that ferulic acid stimulated the Sirt1/p-AMPK pathway in human cervical carcinoma cells (39). Therefore, PF may regulate AMPK phosphorylation via the Sirt1/LKB1 pathway.

In conclusion, the present study indicated that PF improved filtration barrier injury of GEnCs in a model of early DN, and that this action was mediated by AMPK (Fig. 7). Therefore, restoration of GFB injury via modulation of AMPK activity may serve an alternative approach for future treatment of DN.

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Availability of data and materials
The datasets used and/or analyzed during the present study are available from the corresponding author on reasonable request.

Authors’ contributions
LYY and DXX designed the study. YYY, ZC and XDY performed the experiments. RRD and LXS contributed to data analysis. YYY wrote and revised the manuscript. All authors have read and approved the final manuscript. YYY and LXS confirm the authenticity of all the raw data.

Ethics approval and consent to participate
The animal experimental protocol used in the present study was approved by the Ethics Committee of Animal Experiments of the Central South University and was performed in accordance with the Guidelines for the Care and Use of Laboratory Animals.

Patient consent for publication
Not applicable.

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

References


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