Geniposide Attenuates Hyperglycemia-Induced Oxidative Stress and Inflammation by Activating the Nrf2 Signaling Pathway in Experimental Diabetic Retinopathy

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Geniposide (GEN) is a natural antioxidant and anti-inflammatory product and plays an important role in the treatment of diabetes and diabetic complications. To explore the biological functions and mechanism of GEN in diabetic retinopathy (DR), we constructed the in vitro and in vivo model of DR by using primary cultured mouse retinal Müller cells and C57BL/6 mice, respectively. We found that GEN inhibited ROS accumulation, NF-κB activation, Müller cell activation, and inflammatory cytokine secretion both in vitro and in vivo, which is probably mediated through the Nrf2 pathway. Exendin (9-39) (EX-9), an antagonist of glucagon-like peptide-1 receptor (GLP-1R), abolished the protective effect of GEN on high glucose- (HG-) induced Müller cells. Additionally, GEN decreased hyperglycemia-induced damage to Müller cells and blood-retinal barrier in the retinas of mice with DR. We demonstrated that GEN was capable of protecting Müller cells and mice from HG-induced oxidative stress and inflammation, which is mostly dependent on the Nrf2 signaling pathway through GLP-1R. GEN may be an effective approach for the treatment of DR.

1. Introduction

Diabetes is one of the most important and serious global health problems worldwide. Epidemiological studies have shown that the prevalence of diabetes will increase to 7.7% and affect 439 million adults by 2030 [1]. Diabetes causes serious damage to microvessels and macrovessels and leads to vascular complications in the human body [2]. Among microvascular complications, diabetic retinopathy (DR) remains one of the most serious and common diabetes-associated complications. As the main glial cells in the retina, Müller cells play a crucial role in the progression of DR. Our previous studies showed that Müller cells become activated and secrete several inflammatory cytokines in experimental DR model. Inhibition of Müller cell gliosis may decrease damage to the blood-retinal barrier (BRB) and reduce the loss of retinal ganglion cells (RGCs) [3, 4]. Thus, preventing Müller cell gliosis and subsequent inflammatory factor production may be an effective therapeutic strategy for DR treatment.

Chronic hyperglycemia-induced oxidative stress and low-grade inflammation are thought to play crucial roles in the onset and development of DR [5]. When there is an imbalance between excessive reactive oxygen species (ROS) production and the ability of endogenous antioxidant factors to clear ROS, oxidative stress occurs [6]. Oxidative stress is characterized by ROS-induced overexpression of proinflammatory and proangiogenic factors, which damage glial cells, vascular cells, and neurons [7, 8]. Cumulative evidence has indicated that ROS plays an important role in activating transcription factor nuclear factor-kappa B (NF-κB) [9, 10]. NF-κB is triggered and translocated to the nucleus
where it activates the transcription of proinflammatory cytokines, such as tumor necrosis factor-α (TNF-α), interleukin-6 (IL-6), and interleukin-1β (IL-1β) [11, 12].

The interaction of retinal glial cells and blood vessels is important for maintaining the homeostasis and survival of retinal tissue [13]. Müller cells are the main glial cells in the retina, and under hyperglycemic conditions, they undergo oxidative damage and exhibit a reactive phenotype, which is manifested by the upregulation of glial fibrillary acidic protein (GFAP) expression and the subsequent production of proinflammatory factors [14, 15]. In addition, Müller cells span the entire thickness of the retina, and the anatomical association of Müller cells with neurons and microvessels means that damage to Müller cells will lead to severe injury to neurons and blood vessels [16–18]. Therefore, inhibiting oxidative stress and inflammation in Müller cells in a hyperglycemic environment may improve retinal vascular and nerve damage to alleviate DR progression.

Geniposide (GEN) is a natural product extracted from gardenia fruit that has a variety of biological properties, such as antioxidant and anti-inflammatory activities [19, 20]. Studies have revealed that GEN protects against myocardial ischemia reperfusion injury in diabetic rats by suppressing oxidative stress through the nuclear factor erythroid 2-related factor (Nrf2)/heme oxygenase-1 (HO-1) signaling pathway [3]. In ApoE-/- mice and RAW264.7 cells, GEN treatment decreased the areas of atherosclerotic plaques and the production of inflammatory cytokines, and the anti-inflammatory mechanism was related to the miR-101/mitogen-activated protein kinase phosphatase-1/P38 signaling pathway [21]. The metabolite of GEN, genipin, leads to HO-1 upregulation and participates in the anti-inflammatory response, which is mediated by PI3 kinase and activation of the downstream targets JNK1/2 and Nrf2 [22]. Studies have suggested that GEN plays a vital antioxidant and anti-inflammatory role and is closely associated with the Nrf2 signaling pathway. Moreover, as a novel agonist of glucagon-like peptide-1 receptor (GLP-1R), GEN also has a role in the treatment of diabetes and diabetic complications [23–25]. However, the effect of GEN on DR pathogenesis and whether Nrf2 is involved remain unknown. Therefore, the purpose of our study was to explore the role of GEN in DR development and the underlying mechanisms.

2. Materials and Methods

2.1. Cell Culture. Mouse primary retinal Müller cells were isolated from 3-day-old newborn C57BL/6 pups. The mice were obtained from the Experimental Animal Center of Soochow University. Müller cells were extracted and identified as previously described [26]. Retinal Müller cells were cultured in DMEM supplemented with 10% fetal bovine serum (FBS, Gibco), streptomycin (100 mg/ml), and penicillin (100 U/ml) (Gibco). Then, the cells were cultured in a humidification incubator (5% CO₂) at 37°C, and the medium was replaced every two days.

2.2. Cell Treatments. The cells were treated with 5 mM D-glucose (normal glucose, NG) or 30 mM D-glucose (HG) (#310808; Sigma, USA) for 24 h to mimic the diabetic environment before or after certain experiments. Different concentrations of GEN (#SML0153; Sigma, USA) (25, 50, 100, and 200 μg/ml) were used to treat Müller cells for 24 h before HG was added. The dose used of GEN was based on a previous study [27]. The ROS scavenger N-acetylcysteine (NAC) (1 mM, 24 h) (#A7250, Merck, USA) [26] and GLP-1R antagonist exendin (9–39) (EX-9) (200 nmol/l, 1 h) (#ab141101, Abcam, UK) [28] were used to treat the cells before HG or GEN administration. After certain treatments, the cell culture medium was collected and stored at -80°C for cytokine analysis.

2.3. Cell Viability Assay. Müller cells were seeded in 96-well plates at a density of 5 × 10³ cells/well. After 24 h, the cells were treated with different concentrations of GEN (25, 50, 100, and 200 μg/ml) with or without HG for 24 h. Then, cell viability was measured by a Cell Counting Kit-8 (CCK-8, #CK04, Dojindo, Japan) according to the manufacturer’s instructions. Briefly, after treatment, Müller cells were washed with Hank’s solution, and 100 μl of medium supplemented with 10 μl of CCK-8 solution was added to each well. After being incubated for 2 h in a CO₂ incubator, the absorbance at 450 nm was measured by a Thermo MultiSkans GO microplate reader (Thermo Fisher, USA).

2.4. Immunofluorescence Analysis. Müller cells were seeded in the slide chamber at a density of 1 × 10⁵ cells per well. After the indicated treatments, the cells were fixed with 4% paraformaldehyde (PFA) at room temperature for 30 min. The retinal patch was first isolated from the mouse retina and fixed at room temperature with PFA for 1 h. After being blocked and permeabilized, the slides were incubated with nuclear factor kappa B (NF-κB) p65 antibodies (host species: rabbit; species reactivity: mouse, rat; dilution: 1: 1000; #ab16502, Abcam), and the retinas were incubated with lectin B4 (1: 1000, #121411, Invitrogen) and GFAP (host species: rabbit; species reactivity: mouse, human; dilution: 1:1000; #CK04, Dojindo, Japan) according to the manufacturer’s instructions.

2.5. Animals. Male C57BL/6 mice (8-weeks old) were purchased from the Laboratory Animal Center of Soochow University. The mice were housed in standard pathogen-free conditions and were randomly divided into 6 groups: normal, normal+PBS, normal+GEN, DM, DM+PBS, and DM+GEN. The diabetic mouse model was established as previously described [3]. Briefly, the mice were fasted for 12 hours before streptozotocin (STZ) injection. Then, the mice received intraperitoneal injections of 50 mg of STZ once per day for 5 consecutive days. Mice with a blood glucose greater than 16.7 mmol/l were regarded as diabetic and were used in follow-up experiments. The mice in normal group were given an intraperitoneal injection of the same amount of citrate buffer. Four weeks after successful
modeling, the mice in the GEN treatment group were given
tail vein injections of GEN (50 mg/kg/day) for one week.
GEN was dissolved in phosphate-buffered saline (PBS),
and equal volumes of PBS without GEN were injected as
the solvent control. Finally, the mice were sacrificed for
certain experiments. All animal experiments were approved
by the Animal Research Ethics Committee of Soochow
University and were in accordance with the Chinese
National Standard.

2.6. Intracellular ROS Detection. The generation of ROS was
measured by an ROS assay kit (#S0033M, Beyotime, China).
In vitro, Müller cells were seeded on 24-well plates and
measured by an ROS assay kit (#S0033M, Beyotime, China).
The level of ROS in the retinas of mice was measured as pre-
viously described [29]. ROS levels were measured by an
immunofluorescence microscope (Leica).

2.7. Western Blotting. Total protein was extracted from cul-
tured Müller cells and mouse retinas with protein lysis buffer.
Cytoplasmic and nuclear proteins were extracted from
cells using a PARIS Kit (#AM1556, Life Technologies,
USA) according to the manufacturer's instructions. The protein concentration was measured by a BCA assay kit
(#P0012S, Beyotime, China). Equal amounts of proteins in
each sample were separated by SDS-PAGE and then trans-
ferred to PVDF membranes. After being blocked, the
membranes were hybridized with primary antibodies against
IκBα (1:1000 dilution, #ab7217, Abcam), p-IκBα (1:1000
dilution, #2859, Cell Signaling Technology, USA), NF-κB
P65 (1:2000 dilution, #ab16502, Abcam), p-P65 (1:1000
dilution, #3031, Cell Signaling Technology), GFAP (1:3000
dilution, #ab7260, Abcam), Nrf2 (1:1000 dilution, #ab92946,
Abcam), β-actin (1:1000 dilution, #ab8226, Abcam), histone
H3 (H3) (1:1000 dilution, #ab6147, Abcam), HO-1 (1:2000
dilution, #ab189491, Abcam), NAD(P)H quinone dehydro-
genase 1 (NQO1) (1:1000 dilution, #ab34173, Abcam), and
GAPDH (1:5000 dilution, #ab8245, Abcam) at 4°C overnight.
β-Actin and H3 were used as cytoplasmic and nuclear internal
controls, respectively. The membranes were then washed and
hybridized with horseradish peroxidase-conjugated secondary
antibodies (Cell Signaling Technology) at room temperature
for 1 h. The protein bands were detected with a chemilumines-
cence reagent and visualized with a Bio-Rad imaging system
(Bio-Rad Laboratories, Hercules, CA, USA). ImageJ software
was used to quantify the band intensities, and GAPDH was
used as the loading control.

2.8. Quantitative Real-Time PCR (qRT-PCR). Total RNA
was extracted with TRIzol reagent (#15596018, Invitrogen,
USA) according to the manufacturer’s instructions. Total RNA (1 μg)
was synthesized into cDNA using a Revert Aid
First Strand cDNA Synthesis Kit (#K1622, Thermo Scien-
tific, USA). The sequences of the qRT-PCR primers were as
follows: VE-cadherin: forward primer 5′-TGGAAAGGTCT
TGACCTGCTA-3′, reverse primer 5′-TCTGCTGGTG
TGCCCAACAT-3′, reverse primer 5′-CGGCACCCCTGC
GATCA-3′; occludin: forward primer 5′-TGCTGGATA
AGGAACATTTATGA-3′, reverse primer 5′-CAGACA
CATTITTAACCCACTCTCA-3′; and ZO-1: forward
primer 5′-TGAACGCTCTATAAGCTTCGTAA-3′, reverse
primer 5′-ACCGTACCAACCATCATTTAGT-3′. PowerUP
SYBR Green Master Mix (#A25742, Thermo Scientific) was used
to detect the transcription products of the cDNA samples
on an ABI 7500 Real-Time PCR system (Foster City, USA).
GAPDH was used as the internal control, and the 2−ΔΔCT
method was used to calculate the relative expression of the
target genes.

2.9. Enzyme-Linked Immunosorbent Assay (ELISA). The concentrations of TNF-α (#BMS603-7, Invitrogen), IL-1β
(#PI301, Beyotime, China), and IL-6 (#BMS603-2, Invitrogen)
in the cell culture supernatant and retinas were measured by commercial ELISA kits according to the manu-
facturers’ instructions. By measuring the optical density of
each well at 450 nm, the concentrations of these cytokines
were quantified with reference to the standard curve.

2.10. Cell Transfection. Nrf2 siRNA and scramble siRNA
were purchased from RIBO Biology Company (Nrf2 siRNA,
5′-UGAAAGCACAGCAAGAUUT-3′). According to the manu-
facturers’ instructions, Lipofectamine 2000 transfec-
tion reagent (#11668019, Invitrogen, USA) was used to per-
form the cell transfections.

2.11. Statistical Analysis. All data are presented as the mean ± SEM. GraphPad Prism version 7 software (GraphPad,
USA) was used for statistical analysis. All the experiments in
our study were repeated at least three times. Student’s t-test
(2-group comparisons) and one-way ANOVA followed by
Tukey’s multiple comparison posttest (multiple-group com-
parisons) were used to assess whether there was a signif-
ant difference between the groups.

3. Results

3.1. ROS Accumulation Promotes NF-κB Activation,
Followed by Glial Activation and Inflammatory Cytokine
Secretion in HG-Stimulated Müller Cells. As shown in
Figure 1(a), intracellular ROS levels were increased under
HG conditions and decreased by the ROS scavenger NAC
(Figure 1(a)). After HG stimulation, the phosphorylation of
the NF-κB-related proteins IκBα and P65 was significantly
upregulated, and this effect was reversed by NAC pretreat-
ment (Figures 1(b)–1(d)). We also used western blot and
immunofluorescent staining to determine the subcellular
localization of NF-κB P65 in Müller cells. We found that
NF-κB P65-positive staining was mainly localized in the
cytoplasm in the control group and was translocated to the
nucleus by HG stimulation. Interestingly, the HG-induced
translocation of NF-κB P65 was inhibited in the presence
of NAC (Figure 1(g)). Additionally, HG promoted the expres-
sion of the glial activation marker GFAP (Figures 1(b) and
1(e)) and inflammatory factors, including TNF-α, IL-1β, and
Figure 1: Continued.
IL-6 (Figures 1(h)–1(j)), while NAC treatment counteracted the effect of HG. These data suggest that HG promotes ROS accumulation and induces the NF-κB signaling pathway activation, thereby increasing glial activation and inflammatory cytokine secretion by HG-stimulated Müller cells.

3.2. GEN Is Not Cytotoxic to Müller Cells within a Range of Concentrations. We next examined the cytotoxicity of GEN on Müller cells under normal or HG conditions. As shown in Figures 2(a) and 2(b), 25, 50, and 100 μg/ml GEN did not inhibit Müller cell viability, while 200 μg/ml GEN suppressed cell viability under both normal and HG conditions (Figures 2(a) and 2(b)).

3.3. GEN Induces Nrf2 Nuclear Translocation and Inhibits ROS Accumulation in HG-Stimulated Müller Cells. We also found that GEN treatment dose-dependently induced Nrf2 nuclear translocation (Figures 2(c) and 2(d)). Moreover, the Nrf2-targeted antioxidant genes HO-1 and NQO1 were also assessed. As shown in Figures 2(e) and 2(f), the expression of HO-1 and NQO1 in Müller cells was markedly decreased under HG conditions and dose-dependently increased by GEN treatment. HG stimulation exacerbated ROS accumulation. Conversely, GEN significantly reduced HG-induced ROS levels (Figure 2(g)).

3.4. GEN Inhibits NF-κB Activation and HG-Induced Glial Activation and Inflammatory Cytokine Secretion in Müller Cells. The inhibitory effect of GEN on ROS accumulation prompted us to investigate whether GEN was involved in HG-induced NF-κB activation, glial activation, and inflammatory cytokine secretion. We found that the phosphorylated forms of the NF-κB-related proteins IκBa and P65 were significantly upregulated after HG stimulation, and this effect was dose-dependently reversed by GEN pretreatment (Figures 3(a)–3(c)). Additionally, HG promoted the expression of GFAP and the secretion of inflammatory cytokines, including TNF-α, IL-1β, and IL-6, while GEN treatment reversed HG-mediated promotion of gliosis (Figures 3(a) and 3(d)) and inflammatory cytokine secretion (Figures 3(e)–3(g)).

3.5. GEN Inhibits NF-κB Activation, Müller Cell Activation, and Inflammatory Cytokine Secretion through the Nrf2 Antioxidant Pathway. To confirm whether the anti-inflammatory effect of GEN was mediated by activating the Nrf2 antioxidant pathway, we investigated the effect of Nrf2 knockdown on oxidative stress and inflammation in HG-stimulated Müller cells. GEN was used at a concentration of 100 μg/ml in the subsequent experiments due to its improved protective effects. The results suggested that GEN treatment significantly promoted the expression of HO-1 and NQO1, and this effect was reversed by downregulating Nrf2 (Figures 4(a)–4(c)). Conversely, GEN decreased ROS accumulation in HG-stimulated Müller cells, and Nrf2 knockdown blocked the inhibitory effect of GEN on ROS accumulation (Figure 4(d)). In addition to cellular oxidative stress, Nrf2 knockdown also suppressed the effect of GEN on the NF-κB pathway activation (Figures 4(e)–4(g)), expression of the glial activation marker GFAP (Figures 4(e) and 4(h)), and inflammation-associated cytokine secretion (Figures 4(i)–4(k)).

3.6. The Protective Effects of GEN on HG-Stimulated Müller Cells Were Abolished by a GLP-1R Antagonist. To further investigate the mechanism of GEN, we evaluated the effects of the GLP-1R antagonist EX-9 on oxidative stress and inflammation in Müller cells. GEN increased the nuclear translocation of Nrf2, and EX-9 inhibited the effect of GEN (Figures 5(a) and 5(b)). Similarly, EX-9 eliminated the GEN-induced increases in HO-1 and NQO1 (Figures 5(c)–5(e)). In addition, GEN suppressed ROS accumulation (Figure 5(f)), NF-κB pathway activation (Figures 5(g)–5(i)), expression of the glial activation marker GFAP (Figure 5(j)), and inflammation-associated cytokine secretion (Figures 5(k)–5(m)), and these effects were reversed by EX-9.
3.7. GEN-Mediated Activation of the Nrf2 Signaling Pathway Reduces NF-κB Activation and Decreases GFAP Production and Inflammatory Cytokine Secretion in the Retinas of Diabetic Mice.

To investigate whether the Nrf2 antioxidant signaling pathway and the NF-κB inflammatory pathway were involved in diabetic mice in vivo, we treated the DR mouse model with GEN. As shown in Figures 6(a) and 6(b), GEN increased the nuclear translocation of Nrf2 in the retinas of diabetic mice. In addition, the Nrf2-targeted antioxidant genes HO-1 and NQO1 were decreased in the retinas of mice with diabetes, and GEN treatment increased Nrf2 antioxidant pathway activation in diabetic mice without affecting normal mice (Figures 6(c)–6(e)). Consistent with the in vitro experiments, ROS accumulation and the expression of p-IκBα and p-P65 were upregulated in mice with DR and were reversed by GEN treatment (Figures 6(f)–6(i)). Analyses of glial reactivity and inflammatory cytokine production are shown in Figure 6(g) and

**Figure 2:** The effects of geniposide (GEN) on viability, Nrf2 nuclear translocation, and ROS accumulation in HG-stimulated Müller cells. Müller cells were exposed to different concentrations of GEN with or without HG. (a, b) The viability of Müller cells was measured by CCK-8 assays. **P < 0.01 vs. the control group. (c, d) The protein expression of Nrf2 in the nucleus and cytoplasm was measured by western blotting. H3 and β-actin were used as nuclear and cytoplasmic loading controls, respectively. *P < 0.05 and **P < 0.01 vs. the NG group and #P < 0.05 vs. the HG group. (e, f) The protein expression of HO-1 and NQO1 was measured by western blotting. **P < 0.01 vs. the NG group and #P < 0.05 vs. the HG group. (g) ROS production was measured by an ROS assay kit. n = 4/group.
Figure 3: GEN inhibits NF-κB activation and HG-induced glial activation and inflammatory cytokine secretion in Müller cells. (a–d) The protein expression of IκBα, p-IκBα, P65, p-P65, and GFAP was measured by western blotting. "*" P < 0.01 vs. the NG group and "#" P < 0.05 and "##" P < 0.01 vs. the HG group. (e–g) ELISA was used to measure the protein levels of TNF-α, IL-1β, and IL-6 in Müller cells. "###" P < 0.001 vs. the NG group and "*" P < 0.05 and "**" P < 0.01 vs. the HG group. n = 4/group.
Figure 4: GEN inhibits NF-κB activation, Müller cell activation, and inflammatory secretion through the Nrf2 antioxidant pathway. Nrf2 siRNA transfection was used to knockdown Nrf2 expression, and scramble siRNA (Scr siRNA) was used as the negative control. GEN was administered to Müller cells 24 h prior to stimulation with HG. (a–c) The protein expression of HO-1 and NQO1 was measured by western blotting. ∗∗P < 0.01 vs. the HG group and #P < 0.05 vs. the HG+GEN group. (d) ROS production was measured by an ROS assay kit. (e–h) The protein expression of IκBα, p-IκBα, P65, p-P65, and GFAP was measured by western blotting. ∗∗P < 0.01 vs. the HG group and #P < 0.05 vs. the HG+GEN group. (i–k) ELISA was used to measure the protein levels of TNF-α, IL-1β, and IL-6 in Müller cells. ∗∗P < 0.01 vs. the HG group and ##P < 0.01 vs. the HG+GEN group. n = 4/group.
Figure 5: The protective effects of GEN on HG-stimulated Müller cell activation were abolished by a GLP-1R antagonist. Müller cells were pretreated with exendin (9-39) (EX-9) before HG or GEN administration. (a, b) The protein expression of Nrf2 in the nucleus and cytoplasm was measured by western blotting. **P < 0.01 vs. the HG group and #P < 0.05 vs. the HG+GEN group. (c–e) The protein expression of HO-1 and NQO1 was measured by western blotting. **P < 0.01 vs. the HG group and #P < 0.05 vs. the HG+GEN group. (f) ROS production was measured by an ROS assay kit. (g–j) The protein expression of IκBα, p-IκBα, P65, p-P65, and GFAP was measured by western blotting. **P < 0.01 vs. the HG group and ##P < 0.01 vs. the HG+GEN group. (k–m) ELISA was used to measure the protein levels of TNF-α, IL-1β, and IL-6 in Müller cells. **P < 0.01 and ***P < 0.001 vs. the HG group and ##P < 0.01 vs. the HG+GEN group. n = 4/group.
Figure 6: Continued.
Figures 6(j)–6(m). The levels of GFAP and the secretion of TNF-α, IL-1β, and IL-6 were significantly increased in the retinas of mice with diabetes compared with those of the mice in the normal group. However, GEN decreased glial reactivity (Figures 6(g) and 6(j)) and inflammatory cytokine secretion (Figures 6(k)–6(m)) in DR mice but not in normal mice.

3.8. GEN Decreases Hyperglycemia-Induced Damage to the BRB. Previous studies have shown that glial activation followed by inflammatory cytokine secretion are important factors that damage the BRB in the diabetic retinas [30]. We then investigated the effect of GEN on the BRB in diabetic mice. The results showed that the retinal vessels (marked with isolectin B4) in the diabetic group were tortuous and the exudates were increased, which were reversed by GEN. Moreover, GFAP expression was increased in the retinas of DR mice, and this effect was inhibited by GEN treatment (Figure 7(a)). In DR mice, the expressions of junction proteins such as VE-cadherin, claudin-5, occludin, and ZO-1 were decreased, while this decline was abolished by GEN treatment (Figures 7(b)–7(e)).

4. Discussion

Chronic inflammation and oxidative stress are considered to be the key components of DR pathogenesis, which is characterized by neuronal and vascular degeneration. Hyperglycemia leads to massive ROS production in DR [31]. ROS promotes the production and activation of NF-κB, which in turn translocates to the nucleus and promotes the expression of inflammatory cytokines (such as IL-1β and IL-6) [32]. Müller cells are the main glial cells in the retina and play a central role in retinal metabolism. These cells are highly sensitive to metabolic changes, such as those associated with diabetes [33]. Moreover, Müller cells have been widely used in the investigation of DR pathogenesis, and so we used HG-stimulated Müller cells as an in vitro model to explore the mechanism of DR. We found that under HG conditions, ROS production and NF-κB pathway activation were increased in Müller cells and were significantly attenuated by the ROS scavenger NAC. Simultaneously, the expressions of the NF-κB downstream gene GFAP (a glial activation marker) and inflammatory cytokines, including TNF-α, IL-1β, and IL-6, were enhanced by HG stimulation, whereas NAC obviously reversed the effect of HG, suggesting that ROS exert proinflammatory effects on Müller cells under HG conditions. Based on these findings, we conclude that HG promotes ROS production and induces NF-κB signaling pathway activation, thereby increasing glial activation and inflammatory cytokine secretion in HG-stimulated Müller cells.

It has been reported that GEN exerts potent antioxidant effects to combat various oxidative stress-related diseases, such as osteoblast diseases [34], nonalcoholic fatty liver [35], and myocardial ischemia reperfusion in diabetic rats [36]. According to previous studies, GEN prevents oxidative stress-induced damage by activating the Nrf2 antioxidant pathway [20, 37]. Under oxidative stress conditions, the transcription and synthesis of Nrf2 are increased. Moreover, oxidative stress can also facilitate the dissociation of Nrf2 from the Keap1-Nrf2 complex, allowing Nrf2 to then bind with antioxidant response elements, which in turn promotes Nrf2-mediated regulation of antioxidant genes, such as HO-1 and NQO1 [38, 39]. In the present study, we explored the biological function of GEN in DR and the potential mechanisms. We found that GEN pretreatment increased Nrf2 nuclear translocation and the expression of the downstream genes HO-1 and NQO1. In addition, ROS accumulation was dose-dependently alleviated by GEN. To further verify that GEN plays a role via the Nrf2 pathway, Nrf2 was knocked down by siRNA. The results showed that the GEN-mediated promotion of HO-1 and NQO1 expression and the inhibition of ROS production was reversed by Nrf2 knockdown. Based on these results, GEN protects Müller cells from HG-induced oxidative stress by activating the Nrf2 antioxidant signaling pathway.

Previous studies have shown that a lack of Nrf2 is associated with augmented cytokine production in experimental models of brain injury [40]. The Nrf2 activator dh404 prevented an increase in diabetes-induced inflammatory
mediators, including TNF-α, IL-6, ICAM-1, and MCP-1, in Müller cells [41]. Moreover, studies on Nrf2-/- mouse embryonic fibroblasts (MEFs) showed that IKKβ activity was increased, IκBα phosphorylation was enhanced, and IκBα was subsequently degraded [42]. The NF-κB inflammatory signaling pathway can be regulated by Nrf2. Wang et al. confirmed that genipin, a metabolite of GEN, can activate Nrf2 and thereby inhibit NF-κB activation and inflammatory mediator production in BV2 microglial cells [43]. We revealed that GEN decreased ROS accumulation by activating Nrf2. However, whether GEN is involved in inhibiting NF-κB activation and inflammatory mediator production in Müller cells remains unknown. In this study, we found that GEN inhibited the activation of NF-κB and the downstream gene GFAP and inflammatory cytokines (TNF-α, IL-1β, and IL-6) in a concentration-dependent manner. Furthermore, Nrf2 knockdown reversed the anti-glial and anti-inflammatory effects of GEN on HG-stimulated Müller cells, suggesting that GEN could also inhibit the activation of NF-κB and the downstream gene GFAP and inflammatory cytokines (TNF-α, IL-1β, and IL-6), which are mediated through the Nrf2 antioxidant pathway.

In recent years, GLP-1R agonists have been shown to be effective and safe treatments for diabetes and diabetic complications [44]. GLP-1R activation exerts both neuroprotective and vasculotropic effects to prevent vascular leakage in the context of DR [44]. Moreover, GLP-1R is distributed diffusely in the retina [45]. Therefore, we next explored

**Figure 7:** GEN decreased hyperglycemia-induced damage to the blood-retinal barrier (BRB). The mice in the normal and DM groups were treated with GEN. (a) Immunofluorescence was used to detect isolectin B4 (green) and GFAP (red) in the whole-mount retinas. White arrows show the exudate. (b–e) qRT-PCR was used to measure the mRNA expression of junction proteins, including VE-cadherin, claudin-5, occludin, and ZO-1, in the retinas of mice. n = 4/group.
whether GEN functioned in a GLP-1R-dependent manner, since GEN is a novel agonist of GLP-1R. We confirmed that GEN treatment was beneficial and inhibited HG-induced oxidative stress and inflammation. GEN promoted Nrf2 nuclear translocation and the expression of the downstream genes HO-1 and NQO1, while EX-9 (GLP-1R antagonist) attenuated the inhibitory effect of GEN on oxidative stress. Additionally, GEN suppressed NF-κB pathway activation, GFAP expression, and inflammation-associated cytokine secretion, and these effects were reversed by EX-9. To the best of our knowledge, this is the first report to confirm that the protective effects of GEN against DR are mediated through GLP-1R.

In conclusion, we demonstrated that GEN could protect Müller cells and mice from HG-induced oxidative stress and inflammation, and the effects were mostly dependent on upregulating the Nrf2 signaling pathway through GLP-1R. The activation of Nrf2 inhibited ROS accumulation, thus decreasing NF-κB activation and the subsequent gliosis and inflammatory response (Figure 8). Moreover, GEN treatment alleviated the decrease in the expression of junction proteins and may be an effective approach for the treatment of DR.

Data Availability

The data used to support the findings of this study are available from the corresponding author upon request.

Conflicts of Interest

The authors declare that there are no conflicts of interest.

Authors’ Contributions

M.H.Z. and Y.Y.T. are responsible for the conceptualization. M.H.Z. is responsible for the methodology. L.L.L., L.L.Z., and Y.Y.T. are responsible for the software. M.H.Z. and Y.Y.T. are responsible for the validation. Y.G. is responsible for the formal analysis. S.D. and Y.Y.T. are responsible for the investigation. M.H.Z. and Y.Y.T. are responsible for the resources. L.L.L. and Y.X.Z. are responsible for the data curation. Y.Y.T. is responsible for the writing of the original draft preparation. L.L.L. and M.H.Z. are responsible for the writing of review and editing. L.L.L. and Y.T.Z. are responsible for the visualization. L.L.L. and Z.Z.W. are responsible for the supervision. M.H.Z. is responsible for the project administration. M.H.Z. and Y.Y.T. are responsible for the funding acquisition. All authors have read and agreed to the published version of the manuscript. Yuanyuan Tu and Lele Li contributed equally to this study.

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