

Scutellaria barbata D. Don Polysaccharides Inhibit High Glucose-Induced Proliferation and Angiogenesis of Retinal Vascular Endothelial Cells

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Purpose: The traditional Chinese medicine *Scutellaria barbata* D. Don (*S. barbata*) has been reported to exhibit anti-cancer and anti-inflammation activities. The ethanol extract of *S. barbata* has been confirmed to attenuate diabetic retinopathy (DR). This study aimed to investigate the effects and underlying mechanisms of the polysaccharides isolated from *S. barbata* (PSB) on the proliferation and angiogenesis of retinal vascular endothelial cells (RVECs) in DR.

Methods: Human RVECs (HRVECs) were cultured in normal glucose (NG, 5.5 mM), mannitol (MA, 30 mM), high glucose (HG, 30 mM) and HG plus 40 µg/mL PSB, respectively. Then, cell proliferation, migration and angiogenesis were evaluated. The cell proliferation was also estimated in the presence of SLIGKV, which was used to induce the phosphorylation of ERK (p-ERK).

Results: PSB reduced normal and HG-induced HRVECs cell viability in a concentration-dependent manner. The protein expression of proliferating cell nuclear antigen (PCNA) and proliferating antigen Ki67 (Ki67), the migration rate and tube formation ability, which were increased by HG treatment, were significantly decreased by PSB. PSB also inhibited the phosphorylation of Raf, MEK and ERK in HG-stimulated HRVECs. Moreover, the application of SLIGKV recovered cell viability and the expression of p-ERK, PCNA and Ki67, in HG plus PSB-treated cells. Finally, the HG-enhanced expression of VE-cadherin, Frizzled, β-catenin, MMP-2 and MMP-9 was all reversed by PSB.

Conclusion: PSB could inhibit HG-induced HRVECs proliferation, migration and neovascularization, and these effects might work through blocking the activation of MEK/ERK pathway and VEGF/VE-cadherin axis.

Keywords: *Scutellaria barbata*, polysaccharides, retinal vascular endothelial cells, diabetic retinopathy

Introduction

Diabetic retinopathy (DR) is caused by abnormal insulin metabolism in diabetic patients, which leads to changes in ocular tissues, nerves and blood vessels and microcirculation, resulting in poor ocular nutrition and damage of visual function.¹ It is one of the common complications of diabetes and the primary cause of blindness.² Many different biological processes are involved in the pathological changes of DR, such as oxidative stress, mitochondrial dysfunction, retinal cell proliferation and angiogenesis, but the specific pathogenesis of DR remains to be elucidated.^{3–5} At present, the main therapeutic approaches for DR are anti-vascular

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endothelial growth factor (VEGF) drug therapy (such as oral angiotensin-converting enzyme inhibitors), cold therapy, photocoagulation therapy, and vitrectomy. However, these therapies have not achieved satisfactory therapeutic effects.⁶ Therefore, it is urgent to develop novel and effective drugs for the prevention and treatment of DR.

In the late stage of DR, new blood vessels grow along the retinal interface, which in turn causes complications such as vitreous hemorrhage, traction retinal detachment, and neovascular glaucoma. Angiogenesis is a complex process involving the migration, proliferation and formation of endothelial cell tubes.⁷ Increased VEGF content in the vitreous is the key to retinal neovascularization. VEGF can induce the proliferation and migration of retinal vascular endothelial cells (RVECs), the formation of vascular lumen and the formation of abnormal new blood vessels.⁸ It has been suggested that VEGF could activate MEK/ERK pathway and VEGF/vascular endothelial (VE)-cadherin axis to regulate RVECs proliferation and angiogenesis.⁹ Although research is ongoing in the field of angiogenesis, the underlying molecular mechanism remains to be totally clarified.

Scutellaria barbata D. Don (*S. barbata*), also known as “Ban Zhi Lian” in China, is an important species belonging to the Lamiaceae family and has been applied in traditional Chinese medicine (TCM). *S. barbata* has long been utilized for heat-clearing and detoxification in the TCM system.¹⁰ Modern pharmacological studies have shown that *S. barbata* exhibits a wide range of biological activities, including anti-cancer, anti-inflammation, and anti-microbial.^{11,12} *S. barbata* contains complex chemical composition including flavonoids, lactones and polysaccharide, among them, each specific component has been reported to possess different degrees of anti-inflammatory and anti-tumor properties. The polysaccharides isolated from *S. barbata* (PSB) was illustrated to inhibit tumor angiogenesis in lung cancer.¹³ In a previous study, the ethanol extract of *S. barbata* has been confirmed to attenuate DR by preventing retinal inflammation and downregulating tight junction protein expression,¹⁴ whereas no attention was devoted to investigate whether PSB could relieve the proliferation, migration and angiogenesis of DR.

In the present study, we aimed to explore the effects and underlying mechanisms of PSB on high glucose (HG)-induced human retinal vascular endothelial cells (HRVECs) proliferation and angiogenesis. First, the effect of different concentrations of PSB on normal and

HG-induced HRVECs viability was evaluated. Second, cell proliferation, cell migration, tubular network formation and the expression of VEGF, MEK/ERK pathway and VEGF/VE-cadherin axis-related proteins were investigated in the presence of normal glucose (NG, 5.5 mM), mannitol (MA, 30 mM), HG (30 mM)¹⁵ and HG plus 40 µg/mL PSB. Finally, the effect of phosphorylated (p)-ERK agonist-SLIGKV on the functions of PSB in HG-stimulated HRVECs was determined. The results showed that PSB could inhibit HG-induced HRVECs proliferation, migration and tube formation through blocking the activation of MEK/ERK pathway and VEGF/VE-cadherin axis. The findings presented in this work may contribute to promising new approaches for DR therapy.

Methods

Cell Culture and Treatment

Human retinal vascular endothelial cells (HRVECs; Procell Life Science&Technology Co., Ltd., Wuhan, China; Cat NO. CP-H130) were cultured in Dulbecco's modified Eagle's medium (DMEM) (Thermo Fisher Scientific, USA) with 10% fetal bovine serum (FBS) (Gibco, USA), 100 U/mL penicillin and streptomycin (Sigma-Aldrich, USA) in a humidified atmosphere with 5% CO₂ at 37°C.

The cells in normal glucose (NG), mannitol (MA), high glucose (HG) groups were cultured with 5.5 mmol/L glucose, 30 mM MA and 30 mM glucose (Sigma-Aldrich) for 48 h, respectively. For PSB (Nanjing Ze Lang Pharmaceutical Technology, Ltd., cat. no. ZL20131124BD; purity>95%; dissolved in dimethyl sulphoxide) treatment, cells were treated with different concentrations of PSB for 24h. For HG+PSB group, cells were cultured in HG medium for 48 h and then treated with PSB for 24 h. SLIGKV (100 µM, Sigma-Aldrich) was used to stimulate cells for 24 h to induce the phosphorylation of ERK1/2.¹⁶

Cell Counting Kit-8 (CCK-8)

HRVECs were seeded in 96-well plates with density of 2×10^4 cells per well, then subsequently maintained in indicated medium for 48 h, followed by treated with or without different concentrations of PSB for 48 h. Thereafter, 10 µL CCK-8 (Beyotime Biotechnology Co., LTD, China) solution was added to each well and incubated at 37°C for 2 h, and OD values were measured at 450 nm through a microplate reader (Bio-Rad Model 550, USA).

Wound Healing Assay

HRVECs (2×10^5 cells/well) were seeded in 6-well plates and being starved for 24 h, after which, 10 μ L tip was used to scratch the cells. Subsequently, cells were washed with PBS and treated with NG, MA, HG and HG + 40 μ g/mL PSB at 37°C for 24 h. Images of the scratches were captured at 0 h and 24 h to observe cell confluence. The average widths of the scratch wounds were analyzed using Image J software. The migrated distances were calculated as the width of the scratch at 24 h minus the width of the scratch at 0 h. The relative migration distance was calculated by normalizing to NG group.

Transwell Assay

Transwell assays were performed using 24-well culture plates inserted with 8-mm pore size (Transwell; Falcon, BD Biosciences). The lower chamber was filled with 600 μ L DMEM containing 10% FBS. Cells (1×10^5 cell/well) were seeded into the upper chamber. After 24 h of incubation, the migrated cells were fixed with 5% glutaraldehyde and then stained with crystal violet, and the number of cells in the bottom well was counted by counting chamber.

Tube Formation Assay

Cells (3×10^4 cells/well) were seeded into the 96-well plate, which was precoated with 30 μ L matrigel and then immediately placed at 37°C for 2 h for solidification. The plate was kept at 37°C for 24 h. The pictures of cells were captured using a microscope (Olympus, Japan; magnification, x40) and the number of formed capillaries was counted.

Immunofluorescence Staining

HRVECs were fixed with 4% paraformaldehyde, blocked with 3% FBS albumin (Sigma-Aldrich), and next incubated with polyclonal rabbit anti-proliferating antigen KI67 (Ki67) (Santa Cruz Biotechnology, CA, USA) primary antibody (1:200) overnight at 4°C. Secondary antibody Alexa Fluor 488 donkey anti-rabbit (1:200) was employed for 1 h at 37°C. Nuclei were visualized by staining with 4', 6-diamidino-2-phenylindole (DAPI; Sigma-Aldrich). An Olympus FV1000 confocal laser scanning microscope (Olympus, Japan; magnification, x200) was used to acquire images.

Terminal Deoxynucleotidyl Transferase-Mediated Deoxyuridine Triphosphate (dUTP)-Biotin Nick End Labeling (TUNEL) Apoptosis Staining

Apoptotic cells were examined by an apoptosis detection kit (Roche, Basel, Switzerland) according to standard protocol. Briefly, cells were stained by a TUNEL kit, followed by the counterstaining with DAPI for the nuclei. The staining was observed under a fluorescence microscope (Olympus, Japan; magnification, x200). Cell apoptosis rate was captured under a fluorescence microscope (Olympus Corporation, magnification, x200).

Western Blot Analysis

The HRVECs were lysed with ice-cold RIPA buffer and the concentration of protein was determined using a BCA protein assay reagent kit (Beyotime, China). The proteins (15 μ g) were separated by dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and transferred onto a polyvinylidene fluoride (PVDF) membrane. Then, the membrane was blocked with 5% skimmed milk for 2 h at room temperature. The PVDF membranes were then incubated overnight at 4°C with the following primary antibodies to: proliferating cell nuclear antigen (PCNA; Cell Signaling Technology, Boston, MA, USA), Ki67 (Santa Cruz Biotechnology, CA, USA), (p-) Raf (Cell Signaling Technology, Boston, MA, USA), (p-) MEK (Cell Signaling Technology, Boston, MA, USA), (p-) ERK (Cell Signaling Technology, Boston, MA, USA), VE-cadherin (Cell Signaling Technology, Boston, MA, USA), Frizzled (Abcam Company, Cambridge, UK), β -catenin (Cell Signaling Technology, Boston, MA, USA), MMP-2 (Abcam Company, Cambridge, UK), MMP-9 (Abcam Company, Cambridge, UK) and GAPDH (Cell Signaling Technology, Boston, MA, USA). Followed by being incubated with a horseradish peroxidase (HRP)-labeled secondary antibody (Abcam, Cambridge, MA, USA). The results were visualized with an enhanced chemiluminescence (ECL) detection kit. ImageJ software (National Institutes of Health, Bethesda, MD, USA) was used to quantify the density of the bands.

Reverse Transcription-Quantitative PCR (RT-qPCR)

Total RNA was isolated from HRVECs using TRIzol reagent (Invitrogen), and then reverse-transcribed (RT)

into cDNA using the PrimeScript™ RT reagent kit (TaKaRa). qPCR was performed using TB Green Fast qPCR Mix (TaKaRa) according to instructions. The PCR reaction procedure was as follows: 5 min at 95°C, with 40 cycles of 30 sec at 95°C and 45 sec at 65°C. The relative quantification of mRNA levels was determined using the $2^{-\Delta\Delta C_t}$ method. The real-time PCR primers were as followed: VEGF, forward 5'-ATGCG GATCAAACCTCACCA-3', reverse 5'-CACCAACG TACACGCTCCAG-3'; VEGF-receptor 2 (R2), forward 5'-GCAGGAAGTAGCCGCATTTG-3', reverse 5'-ACTCGAAGAACACGCAACCT-3'. GAPDH, forward 5'-GCAACCGGGAAGGAAATGAATG-3', reverse 5'-CCCAATACGACCAAATCAGAGA-3'. GAPDH was used as a reference gene.

Statistical Analysis

All experiments were repeated three times. All data are represented as mean \pm standard deviation (SD) from 3 independent experiments. One-way ANOVA followed by Bonferroni's test was utilized to analyze the difference among multiple groups by GraphPad Prism 5 software. A value of $P < 0.05$ was considered statistically significant.

Results

PSB Inhibits HG-Induced HRVECs

Proliferation

Firstly, different concentrations (0, 10, 20, 40 and 80 $\mu\text{g}/\text{mL}$) PSB was used to treat cells for 24 h to observe the effect of PSB on HRVECs viability. Results showed that PSB decreased HRVECs viability in a concentration-dependent manner, and 80 $\mu\text{g}/\text{mL}$ PSB led to more than 50% decrease in cell viability (Figure 1A). Additionally, 10, 20 and 40 $\mu\text{g}/\text{mL}$ PSB had no effects on apoptosis of HRVECs, but 80 $\mu\text{g}/\text{mL}$ PSB caused cell apoptosis as comparison to the untreated group (Figure 1B and C). Therefore, 10, 20 and 40 $\mu\text{g}/\text{mL}$ PSB were used to perform the subsequent experiments. Then, cells were treated with NG, MA, HG and HG plus PSB (10, 20 and 40 $\mu\text{g}/\text{mL}$). As shown in Figure 1D, compared with NG and MA group, HG resulted in a markedly increase in cell viability. However, the presence of PSB gradually canceled this elevation in cell viability and the cell viability of HRVECs in HG + 40 $\mu\text{g}/\text{mL}$ PSB group was lower than NG group. Therefore, 40 $\mu\text{g}/\text{mL}$ PSB was chosen for the following experiments.

HRVECs were subjected to NG, MA, HG and HG + 40 $\mu\text{g}/\text{mL}$ PSB, then the expression of proteins involved in cell proliferation was assessed. We found that HG significantly enhanced the protein expression of PCNA and Ki67, which as reversely reduced by additional treatment of 40 $\mu\text{g}/\text{mL}$ PSB (Figure 1E). The same results are observed in Figure 1F, which presented the immunofluorescence staining for Ki67. The above data demonstrated the inhibitory effect of PSB on HG-caused HRVECs proliferation enhancement.

PSB Suppresses HG-Stimulated HRVECs Migration and Angiogenesis

Next, we performed wound healing and transwell assay to investigate cell migration. Figure 2A reveals that HG-induced enhancement in migration distance was remarkably reversed by 40 $\mu\text{g}/\text{mL}$ PSB. Similarly, PSB canceled the HG-increased number of migrated cells in transwell assay (Figure 2B).

The ability of capillary formation in HRVECs was also estimated. As shown in Figure 2C, compared with NG and MA group, the number of capillary-like structure was considerably boosted in HG group. However, the addition of PSB also obviously resulted in a decrease in the number of capillary-like structure. These data implicated that PSB could block HG-induced HRVECs migration and angiogenesis.

The Agonist of p-ERK, SLIGKV Recovers PSB-Inhibited HRVECs Proliferation

Subsequently, we observed that, consistent with previous studies,^{17,18} the mRNA level of VEGF and VEGF-Receptor 2 (R2) was significantly increased upon HG stimulation (Figure 3A and B). At the same time, HG treatment promoted the phosphorylation Raf, MEK and ERK proteins, which were the crucial components of MEK/ERK pathway. However, PSB also reduced the expression of p-Raf, p-MEK and p-ERK, indicating that the activation of MEK/ERK pathway might contribute to the effects of PSB on HG-treated HRVECs (Figure 3C).

Therefore, we utilized SLIGKV, which has been confirmed to result in significantly greater induction of ERK phosphorylation,¹⁶ and in order to stimulate HG + PSB-treated HRVECs. As illustrated in Figure 3D, the expression of p-ERK was indeed enhanced by SLIGKV, as compared to HG + PSB group. In addition, the decreased cell viability and blunted expression of PCNA and Ki67,

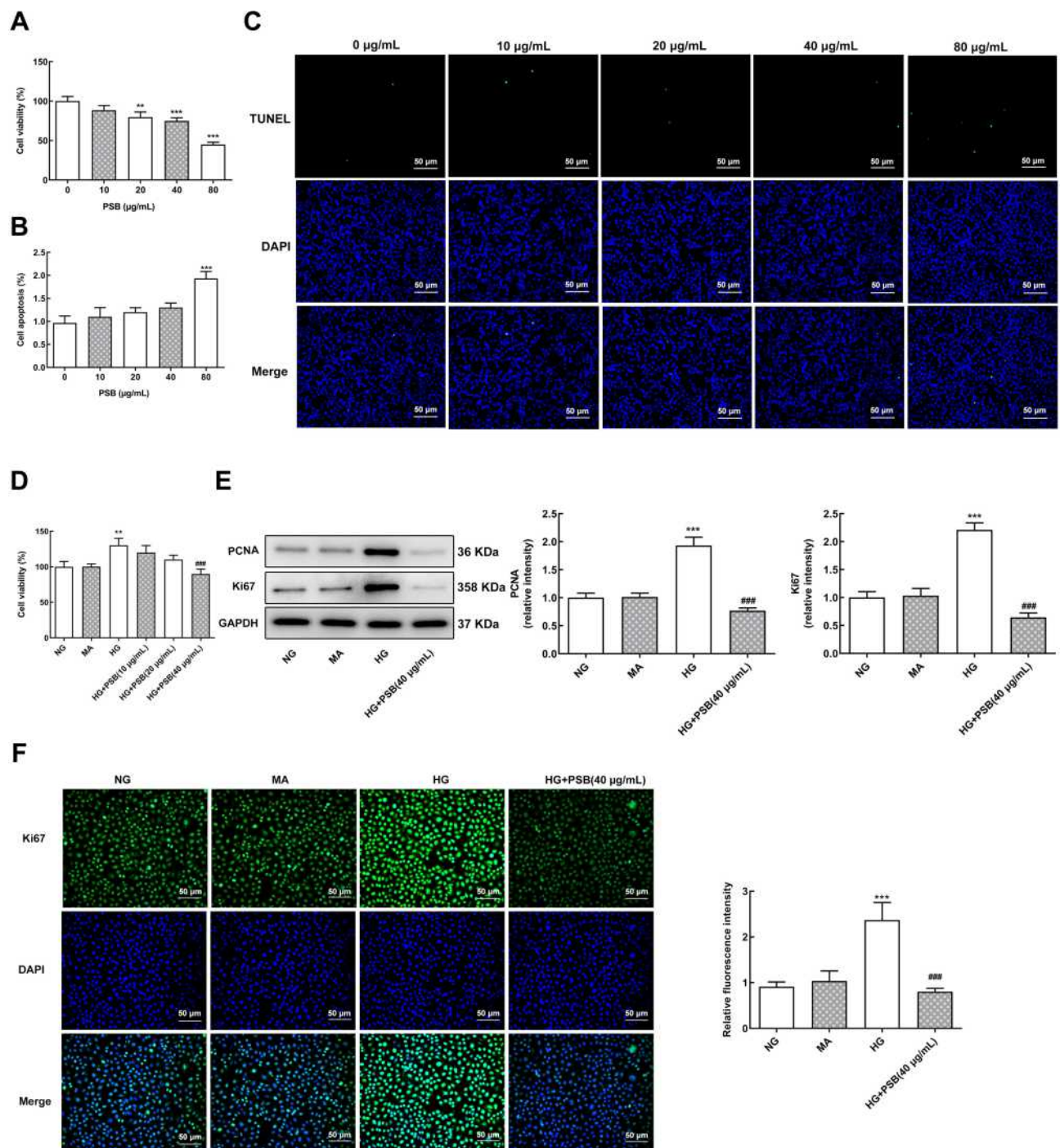


Figure 1 The effect of PSB on HG-enhanced HRVECs proliferation. **(A)** The cell viability of HRVECs that exposed to 0, 10, 20, 40 and 80 µg/mL PSB was determined using a CCK-8 kit. **(B and C)** Cell apoptosis was evaluated with TUNEL staining. ** $P < 0.01$ and *** $P < 0.001$ vs 0 µg/mL PSB. **(D)** The cell viability of HRVECs that cultured in NG, MA, HG or HG plus PSB (10, 20, 40 µg/mL) was examined by a CCK-8 kit. **(E)** The protein expression of PCNA and Ki67 in HRVECs that cultured in NG, MA, HG, and HG plus PSB (40 µg/mL) was assessed by Western blot analysis. **(F)** Representative images of immunofluorescence staining for Ki67 (green), blue represents the nuclei which were stained with DAPI (x200). ** $P < 0.01$ and *** $P < 0.001$ vs NG; #### $P < 0.001$ vs HG.

caused by PSB treatment, were also partially rescued by the addition of SLIGKV (Figure 3E and F). The above results suggested that PSB may inhibit HRVECs proliferation through inactivating MEK/ERK pathway.

PSB Inactivates VEGF/VE-Cadherin Axis in HG-Induced HRVECs

Finally, the expression of proteins associated with VEGF/VE-cadherin axis was detected. As demonstrated in Figure

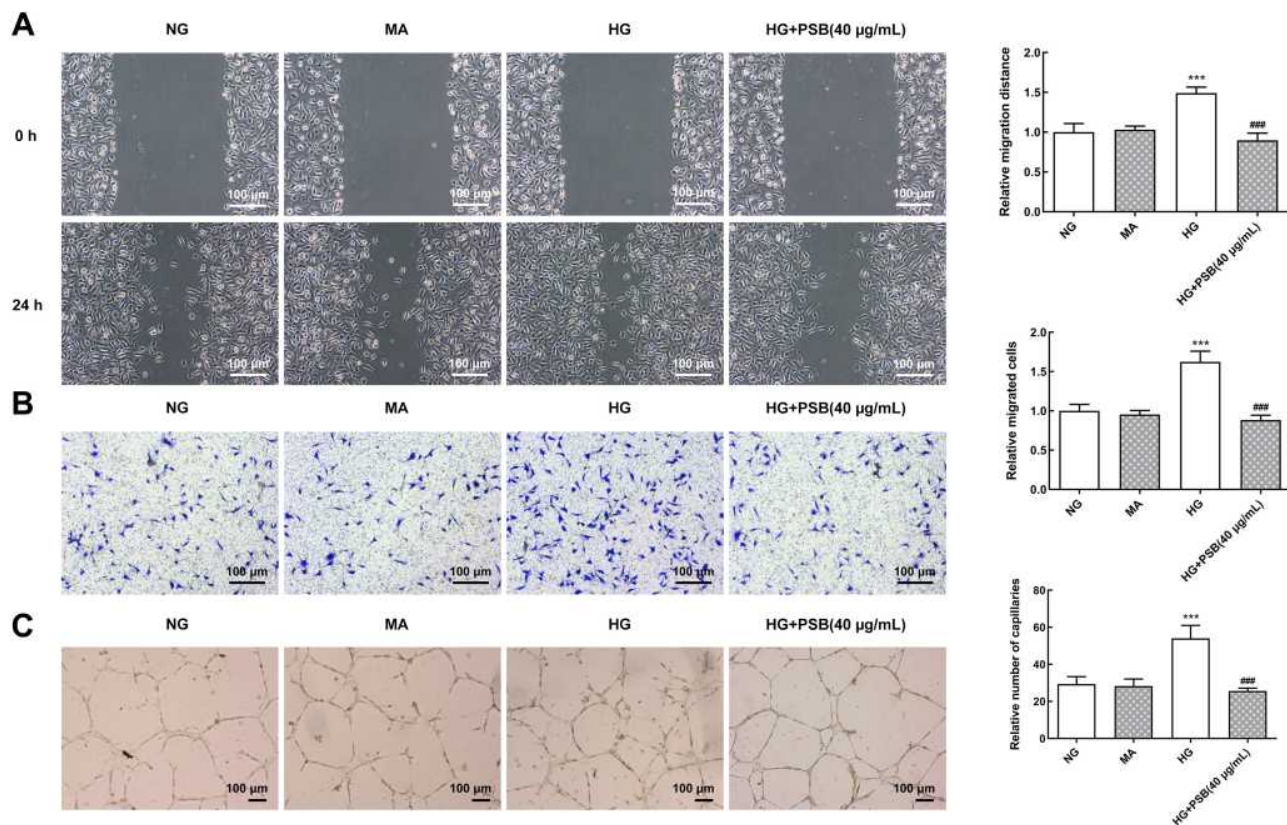


Figure 2 The effect of PSB on HG-stimulated HRVECs migration and angiogenesis. **(A)** Representative images for wound healing assay. The migration distance was calculated as the width of the scratch at 24 h minus the width of the scratch at 0 h. The relative migration distance was calculated by normalizing to NG group. **(B)** Representative images for transwell assay. **(C)** Representative images for tube formation assay. *** $P < 0.001$ vs NG, #### $P < 0.001$ vs HG.

4, compared with NG and MA group, the expression of VE-cadherin, Frizzled, β -catenin, MMP-2 and MMP-9 were also remarkably increased upon HG stimulation. However, the addition of PSB in HG-induced HRVECs, led to a considerably decrease in the expression of these proteins expression, implicating the involvement of VEGF/VE-cadherin axis in the functions of PSB in HG-stimulated HRVECs.

Discussion

DR is the most common cause of diabetic blindness, and the pathological process of DR is retinal microvascular complications. New blood vessel formation is the main feature of DR. This study aims to investigate the beneficial effects of PSB on DR and reveal the underlying molecular mechanisms.

In the present study, the diabetic cell model was established by culturing HRVECs in medium containing high concentration of glucose (HG, 30 mM). HG is known to impair the physiology of normal human endothelial cells and to induce hyperosmolarity, which can promote the development of DR.^{19–21} Moreover, hypoxia induced by

HG in retinal vascular tissues can stimulate angiogenesis by regulating the balance of pro-angiogenic and anti-angiogenic mediators, which results in retinal neovascularization, which is a hallmark of DR.²² To exclude the influence of osmotic pressure, MA group was also set up in this study. In accordance with previous studies, we observed that cell proliferation, migration and angiogenesis were remarkably enhanced in response to HG.^{21,23,24} Furthermore, the mRNA level of VEGF and VEGF-R2 was also increased by HG. VEGF is recognized as one of the most effective angiogenesis factors and is closely related to DR.²⁵ Under normal physical circumstances, VEGF level in retinal tissues is low, thus playing a key role in the formation and stability of retinal blood vessels. While in a state of hyperglycemia, VEGF level is increased and participates in the progression of DR. VEGF-R2 is one of the high-affinity receptors of VEGF in retinal endothelial cells. Previous studies have shown that induction of the VEGF expression and its receptors can promote the proliferation of endothelial cells, stimulate angiogenesis, increase the permeability of small blood

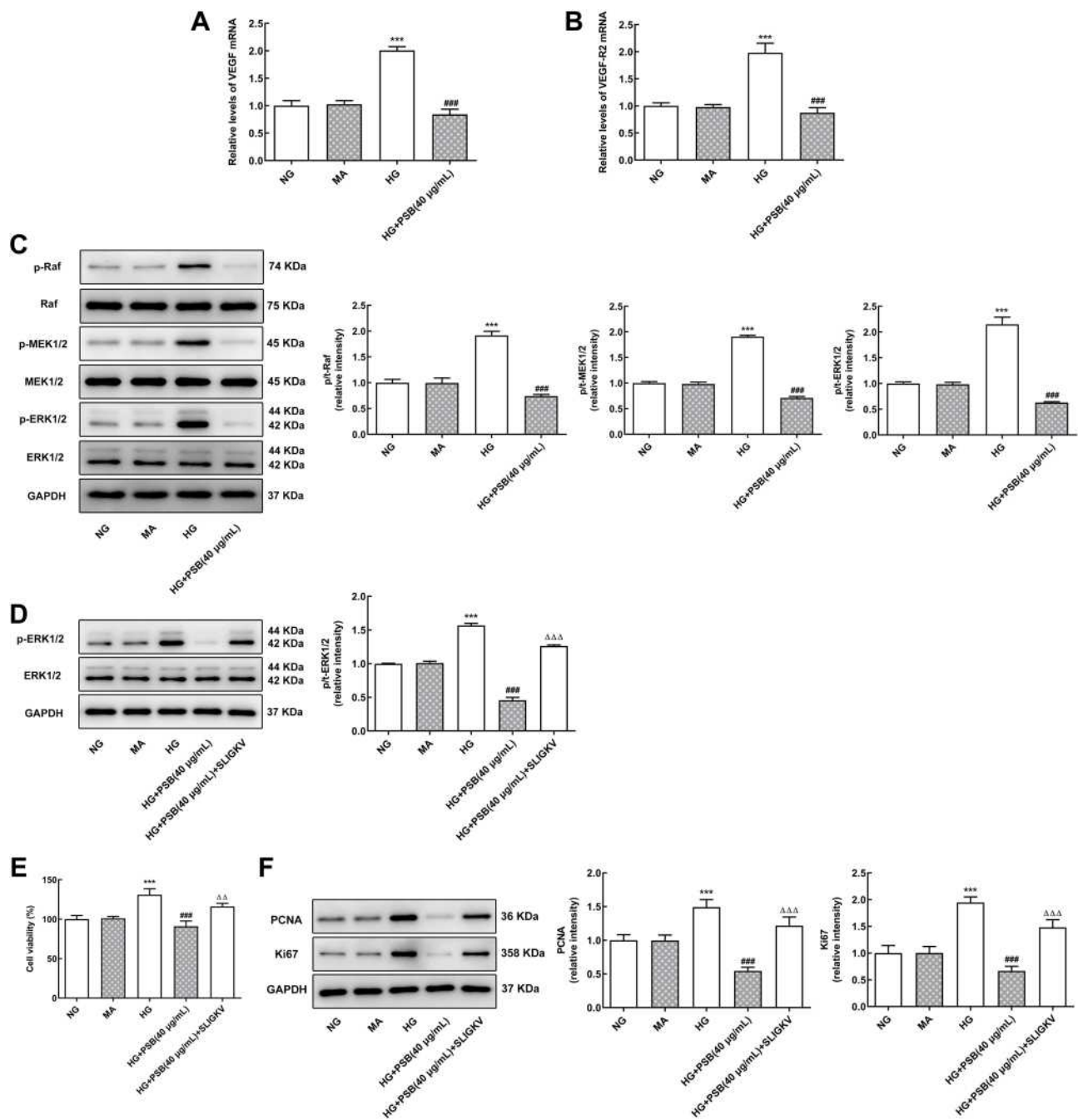


Figure 3 The effect of MEK/ERK signaling on PSB-inhibited HRVECs proliferation. **(A and B)** The expression of VEGF mRNA and VEGF-R2 mRNA in HRVECs that cultured in NG, MA, HG and HG plus PSB (40 μg/mL) was assessed by RT-qPCR. **(C)** The protein expression of p/t-Raf, p/t-MEK1/2 and p/t-ERK in HRVECs that cultured in NG, MA, HG and HG plus PSB (40 μg/mL) was tested by means of Western blot analysis. **(D)** The protein expression of p/t-ERK1/2 in HRVECs that cultured in NG, MA, HG, HG plus PSB (40 μg/mL) and HG plus PSB (40 μg/mL) plus p-ERK inhibitor SLIGKV was measured using Western blotting. **(E)** The cell viability of HRVECs that cultured in NG, MA, HG, HG plus PSB (40 μg/mL) and HG plus PSB (40 μg/mL) plus p-ERK inhibitor SLIGKV was examined by a CCK-8 kit. **(F)** The protein expression of PCNA and Ki67 in HRVECs that cultured in NG, MA, HG, HG plus PSB (40 μg/mL) and HG plus PSB (40 μg/mL) plus p-ERK inhibitor SLIGKV was detected with Western blot analysis. ***P<0.001 vs NG; ####P<0.001 vs HG; ΔΔP<0.01 and ΔΔΔP<0.001 vs HG + PSB (40 μg/mL).

vessels, and thereby exacerbate DR.²⁶ Our results verified the successful establishment of DR cell model induced by HG, and further confirmed the effects HG on DR.

In China, TCM has been widely used to prevent and relieve a great diversity of diseases for more than 1000

years, and certain agents utilized to treat cancer have been derived from TCM. However, the exact anti-disease mechanisms of TCM remain to be fully clarified, which hinders the application of TCM in international clinical treatment. PSB is the polysaccharides

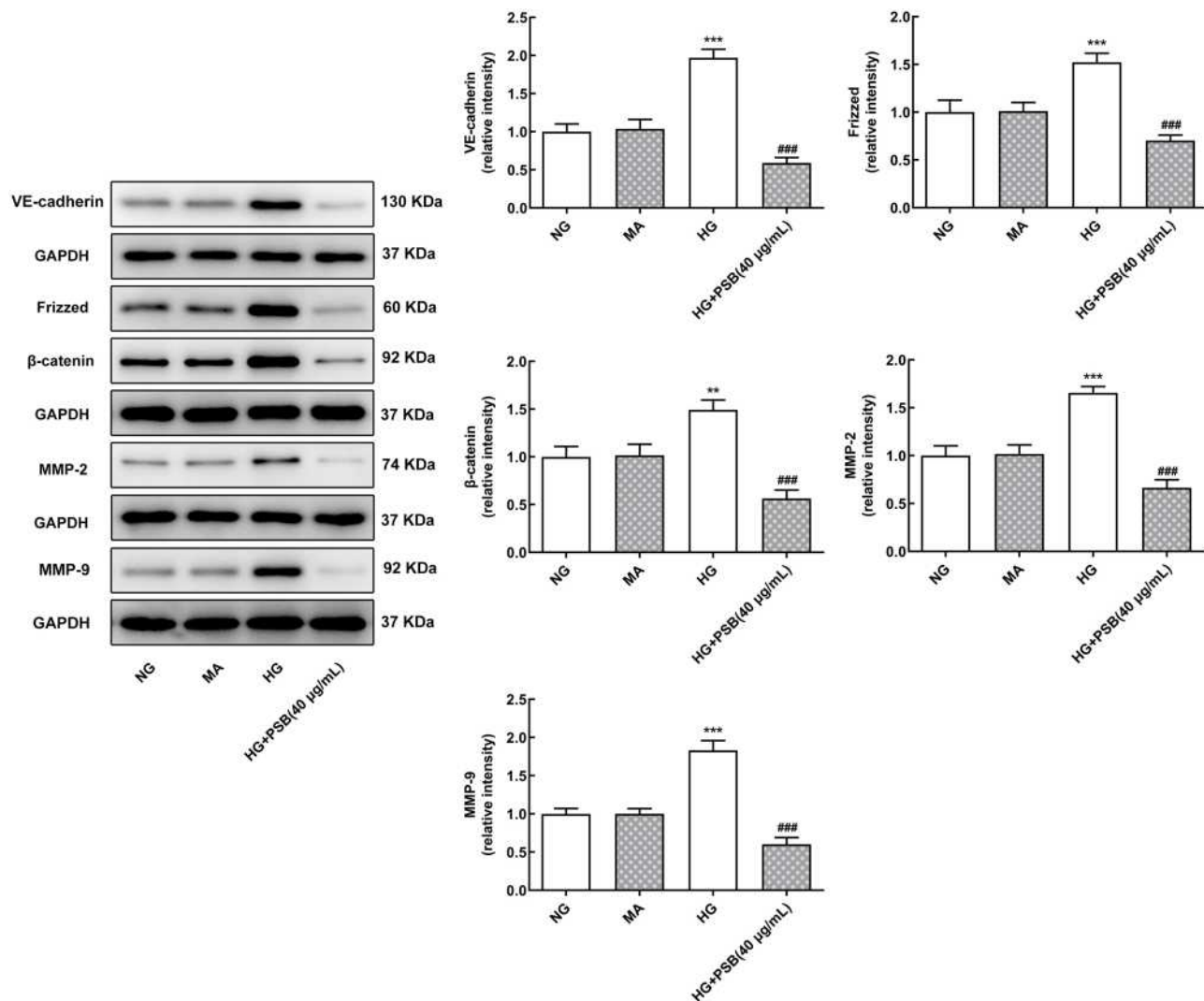


Figure 4 The effect of PSB on the expression of proteins involved in VEGF/VE-cadherin axis. The protein expression of VE-cadherin, Frizzled, β -catenin, MMP2 and MMP9 in HRVECs that cultured in NG, MA, HG, HG plus PSB (40 μ g/mL) was detected by Western blot assay. ** $P < 0.01$ and *** $P < 0.001$ vs NG; #### $P < 0.001$ vs HG.

extracted from a Chinese medicine *S. barbata*. Herein, we demonstrated that PSB effectively inhibited HRVECs proliferation, migration and angiogenesis upon HG stimulation, indicating the potential role of PSB in alleviating DR. Moreover, we found that the HG-increased mRNA level of VEGF and VEGF-R2 was significantly reduced by PSB treatment, suggesting that PSB may exert its beneficial role in DR via lowering VEGF and VEGF-R2 expression. Similarly, the anti-angiogenic effect of *S. barbata* through inhibiting VEGF expression has been proposed by previous studies, although these studies were mainly focused on the anti-tumor effect of *S. barbata*.^{27–29} Our study for the first time shed light on the anti-angiogenic effect of *S. barbata* in retina.

The MEK/ERK pathway has been reported to function as an important regulator in proliferation, differentiation and survival of numerous different cells. It has been demonstrated that Raf/MEK/ERK pathway could be activated in the retina and its endothelial cells in diabetes, and this cascade might promote angiogenesis in the pathogenesis of diabetic retinopathy.^{30,31} In addition, VEGF has been confirmed to activate downstream MEK/ERK pathway thereby contribute to retinopathy.^{9,32} In the current study, we also detected the activation of Raf/MEK/ERK pathway in response to HG, which was markedly prevented by the application of PSB. Inactivation of MEK/ERK has been implicated to restrain the proliferation, migration, invasion and angiogenesis of retinoblastoma cells.³³ Therefore, we speculated that PSB inhibited HG-

induced HRVECs proliferation via inactivating MEK/ERK pathway. Subsequently, the agonist of p-ERK, SLIGKV was applied to treat HRVECs in the presence of HG plus PSB. Results showed that SLIGKV re-activated the expression of p-ERK and also blunted the inhibitory effect of PSB on cell proliferation in HG-induced HRVECs. We inferred that PSB inhibited HG-induced HRVECs proliferation via inactivating MEK/ERK pathway.

VE-Cadherin is a type II endothelial-restricted cadherin, which is the main transmembrane adhesion molecule of endothelial adhesion connection.³⁴ The evidence indicated that silence of the VE-cadherin gene led to severe angiogenesis defects, causing abnormal VEGF signaling. VEGF has recently been recognized as an important angiogenic factor in DR. In addition, angiogenic factors include MMP2 and MMP9, which are downstream of and activated by VEGF.³⁵ In this study, we found that HG stimulation also increased the protein expression of VEGF/VE-cadherin axis including VE-cadherin, Frizzled and β -catenin, together with the expression of MMP2 and MMP9. However, PSB treatment cancelled the enhancement of these proteins induced by HG, indicating that PSB may inhibit HRVECs angiogenesis through blocking VEGF/VE-cadherin and VEGF/MMP axis.

Conclusion

Collectively, we have demonstrated for the first time that PSB restrained DR by suppressing retinal cells proliferation through inactivating MEK/ERK pathway, inhibiting angiogenesis via inhibition of VEGF/VE-cadherin axis and MMP expressions in HG-stimulated HRVECs. Although specific functions of PSB in DR remain to be verified by performing in vivo experiments, the molecular mechanisms of PSB on DR need to be further studied. Our results suggested the potential value of PSB in treating DR.

Ethics Approval

This article does not contain any studies with human participants or animals performed by any of the authors.

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Disclosure

The authors declare that they have no competing interests.

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