

Physcion Protects Rats Against Cerebral Ischemia-Reperfusion Injury via Inhibition of TLR4/NF- κ B Signaling Pathway

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Background: Ischemic stroke (IS) is characterized by the rapid loss of brain function due to ischemia. Physcion has been found to have a neuroprotective effect against cerebral ischemia-reperfusion (I/R) injury. However, the mechanism by which physcion regulates cerebral I/R injury remains largely unknown.

Methods: An oxygen-glucose deprivation/reperfusion (OGD/R) model in SH-SY5Y cells and a rat cerebral ischemia-reperfusion (I/R) model were established, respectively. CCK-8 and flow cytometry assays were used to detect the viability and apoptosis of SH-SY5Y cells. Moreover, enzyme-linked immunosorbent assay (ELISA) was used to measure the levels of SOD, MDA, GSH-Px, TNF- α , IL-1 β , IL-6 and IL-10 in the supernatant of SH-SY5Y cells. Meanwhile, Western blot assay was used to detect the expressions of TLR4, p-p65 and p-I κ B in SH-SY5Y cells and I/R rats.

Results: In this study, physcion treatment significantly rescued OGD/R-induced neuronal injury. In addition, physcion decreased inflammatory response in SH-SY5Y cells after OGD/R insult, as shown by the decreased levels of the pro-inflammatory factors TNF- α , IL-1 β , IL-6 and IL-10. Moreover, physcion attenuated the oxidative stress in OGD/R-treated SY-SY5Y cells, as evidenced by the increased SOD and GSH levels and the decreased ROS and MDA levels. Meanwhile, physcion significantly reduced cerebral infarction, attenuated neuronal injury and apoptosis in I/R rats. Furthermore, physcion markedly decreased the expressions of TLR4, p-NF- κ B p65 and p-I κ B in the brain tissues of rats subjected to I/R and in SH-SY5Y cells exposed to OGD/R.

Conclusion: In conclusion, our study indicated that physcion protected neuron cells against I/R injury in vitro and in vivo by inhibition of the TLR4/NF- κ B pathway; thus, physcion might serve as a promising therapeutic candidate for IS.

Keywords: stroke, ischemia-reperfusion injury, physcion, NF- κ B

Introduction

Stroke is the third leading cause of mortality worldwide, which is an important cause of long-term disability.¹ Ischemic stroke (IS) is a common neurovascular disease in the central nervous system (CNS), which commonly accounts for about 80% of all stroke cases.² Recanalization therapies are known as the primary treatments for IS, which can replenish oxygen, nutrients, and remove toxic metabolites.³ However, restoration of blood flow unavoidably generates pathological damage in the ischemic brain tissue, ultimately leads to the cerebral ischemia/reperfusion (I/R) injury.⁴ Since numerous mechanisms, such as cell toxicity, apoptosis, oxidative stress and inflammation are found to be involved in cerebral I/R injury,⁵ the pathogenesis of stroke is extremely

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complex.⁶ Therefore, the development of new drugs may be beneficial for the treatment of IS.

Nuclear factor-kappa B (NF- κ B), a transcription factor, has been found to participate in several cell functions including hematogenesis, inflammation, immune response, cell apoptosis.⁷ In addition, NF- κ B has been shown to regulate the production of multiple pro-inflammatory factors, including TNF- α , IL-6 and IL-1 β in neuronal cells.⁸ Liu et al indicated that pterostilbene could attenuate neuronal oxidative damage following I/R injury by inactivation of NF- κ B.⁹

Physcion has an anthraquinone chemical structure deriving from the rhizome of the Chinese herbs medicine *Radix et Rhizoma Rhei*.¹⁰ In addition, physcion has been found to exhibit strong anti-inflammatory, anti-bacterial, and anti-tumor properties.^{11,12} Li et al indicated that physcion exerted a potential anti-ischemic effect in cerebral ischemia reperfusion (I/R) model in rats.¹³ Meanwhile, feng et al indicated that rhubarb anthraquinones were easier to be absorbed in I/R rats, which proves the rationality of using it in stroke.¹⁴ However, the function and regulatory mechanism of physcion in IS remains poorly unclear. Therefore, this study aimed to investigate the role of physcion during the neuronal damage after cerebral I/R.

Materials and Methods

Cell Culture

SH-SY5Y cell line was obtained from the Type Culture Collection of the Chinese Academy of Sciences (Shanghai, China). The cells were cultured in Dulbecco's Modified Eagle Medium (DMEM, Thermo Fisher Scientific, Inc., Waltham, MA, USA) supplemented with 10% heat-inactivated fetal bovine serum (Thermo Fisher Scientific) and penicillin/streptomycin, and incubated with 5% CO₂ at 37°C.

Oxygen-Glucose Deprivation/Reoxygenation (OGD/R) Model

For OGD treatment, the normal culture medium was replaced with glucose-free DMEM medium, and cells were exposed to hypoxic conditions (5% CO₂, 95% N₂) at 37°C for 4 h.¹⁵ After that, the cells were reoxygenated under normoxic conditions (5% CO₂ and 95% air) and cultured in a glucose-containing culture medium plus 10% FBS for 24 h. There are three groups: control group, an OGD/R group and an OGD/R + physcion group (physcion intervention group; 10, 20, or 40 μ M

physcion were added). In the control group, the cells were incubated in DMEM supplemented with serum and glucose in a normoxic atmosphere. In the OGD/R group, cells were subjected to OGD/R. In the OGD/R + physcion group, cells were treated with OGD for 4 h and then returned to normal culture conditions with physcion (10, 20, or 40 μ M) for another 24 h.

Cell Survival Assay

Cell viability was determined using the Cell Counting Kit-8 (CCK-8) assay (Dojindo Laboratories, Kumamoto, Japan). Cells were seeded in a 96-well plate at a density of 5×10^3 cells/well, and then 10 μ L of CCK-8 reagent was added into each well and incubated for another 2 h at 37°C according to the manufacturer's instructions. The optical density (OD) value of each well at 450 nm was assessed by an AMR-100 automatic enzyme analyzer (Allsheng, Hangzhou, China).

Cell Apoptosis Assay

Annexin V-FITC/PI Apoptosis Detection Kit (Thermo Fisher Scientific) was used for the detection of apoptotic cells according to the manufacturer's protocol. Cells were washed twice with pre-cooled PBS, and then stained with propidium iodide (PI; 10 μ L) and Annexin V-FITC (10 μ L) for 15 min in the darkness. Subsequently, a FACSsort flow cytometer (BD Biosciences, Mountain View, CA, USA) was applied to measure the percentage of the apoptotic cells.

Intracellular Reactive Oxygen Species (ROS) Detection

SH-SY5Y cells were stained with 10 μ M of 2,7-dichlorofluorescein diacetate (DCFH-DA, Beyotime, Haimen, China) in the dark for 20 min. After that, cells were collected and suspended in 500 μ L of PBS. Subsequently, the fluorescence intensity was determined using a FACSsort flow cytometer (BD Biosciences).

ELISA Assay

The cell culture supernatant samples were collected from SH-SY5Y cells. Then, superoxide dismutase (SOD), malondialdehyde (MDA), glutathione peroxidase (GSH-Px), interleukin-1 β (IL-1 β), tumor necrosis factor- α (TNF- α), interleukin-10 (IL-10), interleukin-6 (IL-6), and macrophage chemoattractant protein 1 (MCP-1) assay kits (#A001-3, A003-1, A005-1, H002, H052, H009, H007,

H115; Jiancheng, Bioengineering Institute, Nanjing, China) were used to measure SOD, MDA, GSH-Px, IL-1 β , TNF- α , IL-10, IL-6, and MCP-1 contents according to the manufacturer's protocol.

Western Blot Assay

Total protein concentration was detected using the BCA Protein Assay kit (Beyotime). Protein samples of 40 μ g per lane were separated by 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE), and then transferred onto a PVDF membrane (Millipore, Billerica, MA, USA). After that, the membrane was blocked with 5% non-fat milk for 1 h at room temperature, and then incubated overnight at 4°C with the following primary antibodies against: toll-like receptor 4 (TLR4; 1:1000, Abcam), phospho-NF- κ B p65 (p-p65; 1:1000, Abcam), NF- κ B p65 (p65; 1:1000, Abcam), phospho-I κ B (p-I κ B; 1:1000, Abcam), I κ B (1:1000, Abcam), β -actin (1:1000, Abcam), and further incubated with HRP-labeled Goat Anti-Rabbit IgG (H+L) antibody (Abcam) at room temperature for 1 h. Finally, the protein bands were visualized by an enhanced chemiluminescent (ECL) reagent (Thermo Fisher Scientific).

Animal Study

SD rats (8 weeks old) were obtained from Shanghai SLAC Animal Center. Middle cerebral artery occlusion (MCAO) was performed in accordance with methods described by Lee et al.¹⁶ Briefly, rats were anaesthetized with ketamine (30 mg/kg) and xylazine cocktail (20 mg/kg). After that, a 7–0 surgical nylon monofilament with a rounded tip was inserted into the internal carotid artery (ICA), then occluding the middle cerebral artery (MCA). After 2 h, nylon monofilament was retracted, and reperfusion began. Then, the rats were allowed to recover for 7 days. These rats were randomly divided into the four groups (n = 18 per group): Sham group; I/R group; I/R + physcion_20 mg/kg group; I/R + physcion_40 mg/kg group. Rats were treated with physcion (20 or 40 mg/kg) via oral gavage after I/R every day. The animal used and experimental protocols were approved by the Animal care Committee of the Dongfang Hospital Beijing University of Chinese Medicine (No. BUCM-20,190,822). All animals were treated in accordance with the National Institutes of Health Guide for Care and Use of Laboratory Animals. Physcion (purity: >98.0%) was obtained from MCE (cat. no. HY-N0108; Med Chem Express, Monmouth Junction, NJ, USA).

Triphenyltetrazolium Chloride (TTC) Staining

TTC staining was performed as described previously.¹⁶ The brain tissues (n = 6 per group) were cut into six slices with 1 mm thickness, and the sections were stained with 2% TTC (Sigma-Aldrich, St. Louis, MO, USA) for 20 min at 37°C. The ipsilateral infarct area was analyzed by ImageJ software. Infarcted area was expressed as pale unstained.

Determination of Cerebral Water Content

The brain tissues (n = 6 per group) were dried at 105°C for 24 h to measure the dry weight. Then, brain water content (%) was calculated as ((wet weight – dry weight)/wet weight x 100%).

Neurological Severity Scorers (NSS)

The neurological function was assessed by NSS described by Chen et al.¹⁷ NSS was conducted by an investigator who was blinded to animal treatments.

Histological Examination

The same brain area of rat in each group (n = 3 per group) was used to perform the hematoxylin and eosin (H&E) staining and TUNEL assays. The brain tissues were fixed in 4% paraformaldehyde at 4°C, and then embedded in paraffin, and cut into 4 μ m-thick sections. The sections were stained with hematoxylin and eosin, and visualized via a Leica DM 5000B microscopy (Leica Microsystems, Wetzlar, Germany). The TUNEL assay was conducted by using a In Situ Cell Death Detection Kit (Roche; Indianapolis, IN, USA). The image of TUNEL-positive cells in brain tissues were captured by a Leica DM 5000B microscopy (Leica Microsystems).

High-Performance Liquid Chromatography-Mass Spectrometry (HPLC-MS)

Cerebrospinal fluid (CSF) samples were mixed with acetonitrile at room temperature, and then centrifuged at 10,000 x g for 5 min. After that, the supernatant of each sample was collected.

Samples were analyzed by LC-MS using an LC-20AD HPLC (Shimadzu, Japan) coupled to an API 400 triple quadrupole MS (AB Sciex, Beijing, China). HPLC separation of physcion was carried out on a Venusil C18 Plus column (50 \times

2.1 mm, 5 μ m particles diameter) manufactured by Agela (Torrance, CA, USA). Mobile phase A was ultrapure water, and mobile phase B was acetonitrile. The flow rate was 0.8 mL/min and the elution gradient were programmed as following: initial, 0% B; 0.01 min, 50% B; 0.5 min, 95% B; 1.3 min, 95% B; 1.4 min, 50% B; 2 min, stop. The MS detector was equipped with the electrospray ionization and operated in the negative ion mode and MRM scan type. Operational electrospray ionization parameters were: spray voltage -4200 V, turbo ion spray temperature 500°C .

Statistical Analysis

Each group was performed at least three independent experiments. Statistical analysis was analyzed by GraphPad Prism v7.0 (GraphPad Software, Inc., La Jolla, CA, USA) and all data were presented as the mean \pm standard deviation (SD). The statistical analysis of the results was analyzed with

a one-way analysis of variance (ANOVA) followed by Tukey's test. $P < 0.05$ was considered significant.

Results

Physcion Protects Cell Against OGD/R-Induced Neuronal Injury

CCK-8 assay was firstly used to assess the effect of physcion on the viability of SH-SY5Y cells. As shown in **Figure 1A**, 20 μM physcion significantly increased the viability of SH-SY5Y cells, while physcion (10, 40 or 80 μM) had a limited effect on the viability of SH-SY5Y cells. In addition, OGD/R remarkably reduced the viability, and increased the release of LDH in SH-SY5Y cells (**Figure 1B** and **C**). Conversely, physcion (10, 20 or 40 μM) reversed OGD/R-induced SH-SY5Y cell injury by increasing cell viability and reducing the release of LDH (**Figure 1B** and **C**). Moreover, OGD/R significantly induced the apoptosis of SH-SY5Y cells;

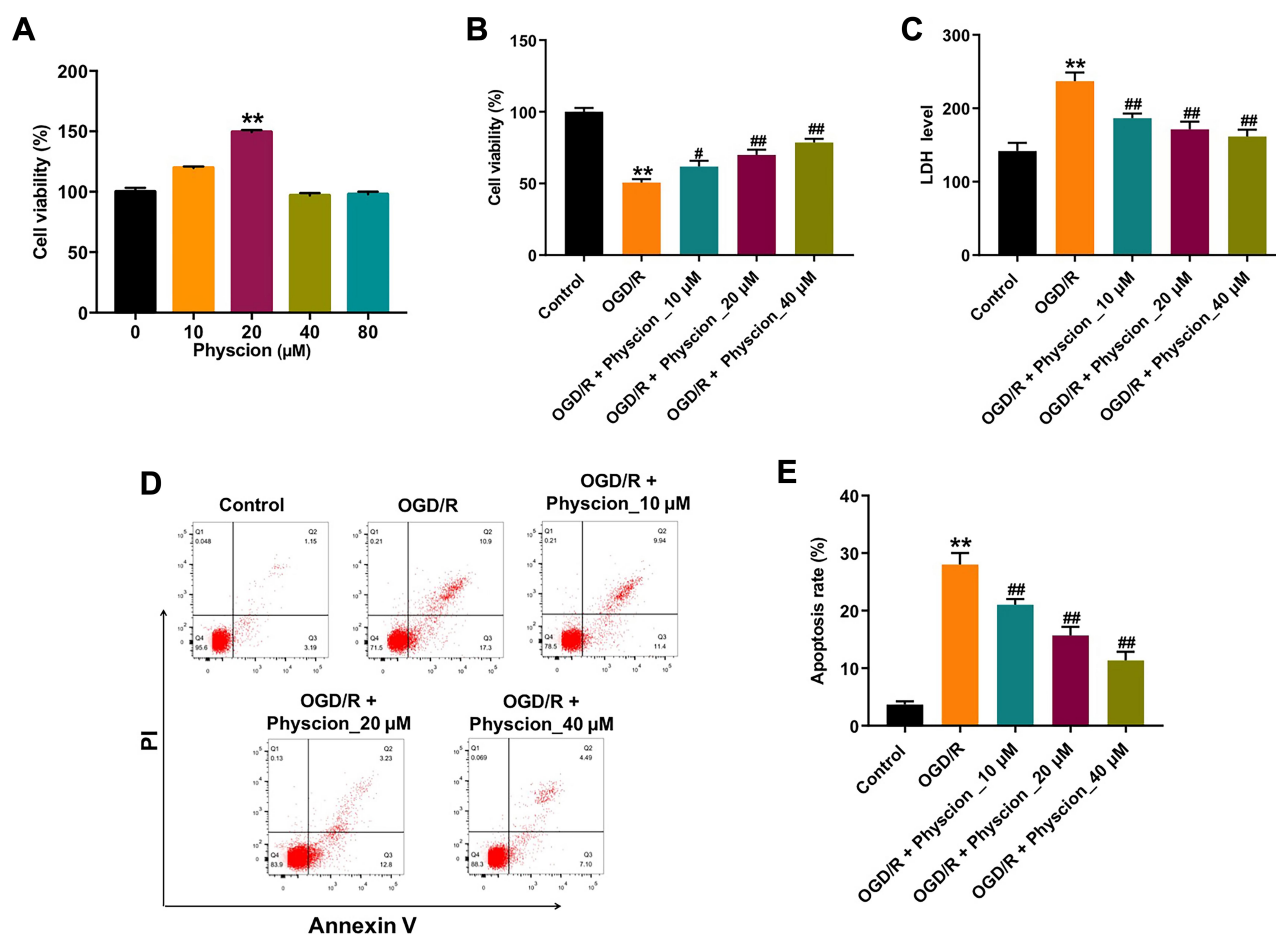


Figure 1 Physcion protects cell against OGD/R-induced neuronal injury. **(A)** SH-SY5Y cells were treated with 0, 10, 20, 40 or 80 μM physcion for 24 h. Cell viability was determined using CCK-8 assay. **(B)** SH-SY5Y cells were exposed to OGD for 4 h, and then reoxygenated in the presence of 10, 20, or 40 μM physcion for 24 h. Cell viability was determined using CCK-8 assay. **(C)** Cell cytotoxicity was determined using LDH release assay. **(D** and **E)** Apoptotic cells were measured by flow cytometry. ** $P < 0.01$ compared with control group; # $P < 0.05$, ## $P < 0.01$ compared with OGD/R group.

Abbreviations: OGD/R, oxygen-glucose deprivation/reperfusion; PI, propidium iodide.

however, OGD/R-induced apoptosis in SH-SY5Y cells was notably reversed by physcion (10, 20 or 40 μM) (Figure 1D and E). All these data demonstrated that physcion could protect cell OGD/R-induced neuronal injury.

Physcion Attenuates OGD/R-Induced Oxidative Stress in SH-SY5Y Cells

Next, we investigated the effects of physcion on OGD/R-induced oxidative stress in SH-SY5Y cells by detecting intracellular ROS generation. As shown in Figure 2A and B, physcion (40 μM) treatment significantly reduced OGD/R-induced ROS production in SH-SY5Y cells. In addition, OGD/R markedly increased MDA level, and reduced SOD and GSH levels in SH-SY5Y cells; however, these changes were markedly reversed by physcion treatment during the course of reperfusion (Figure 2C–E).

These results indicated that physcion could attenuate OGD/R-induced oxidative stress in SH-SY5Y cells.

Physcion Suppresses OGD/R-Induced Inflammatory Response in SH-SY5Y Cells

To investigate whether physcion could suppress the inflammatory response in SH-SY5Y cells in the condition of OGD/R insult, the levels of the pro-inflammatory factors were detected. As shown in Figure 3A–E, OGD/R notably increased the levels of IL-1 β , TNF- α , IL-6 and MCP-1, and reduced IL-10 level in the supernatants of SH-SY5Y cells; however, these OGD/R-induced changes were significantly reversed by physcion treatment during the course of reperfusion. These data suggested that physcion could suppress OGD/R-induced inflammatory response in SH-SY5Y cells.

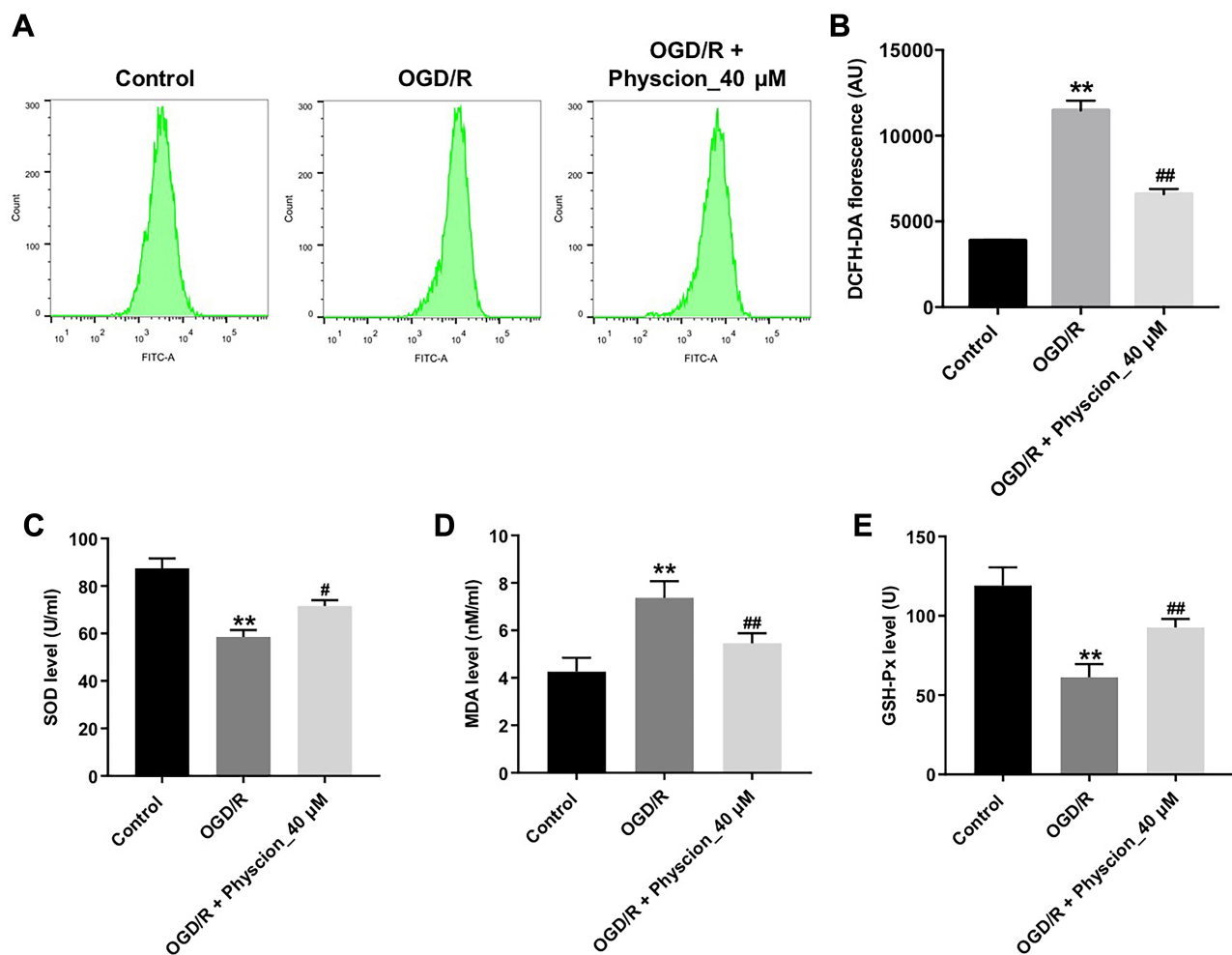


Figure 2 Physcion attenuates OGD/R-induced oxidative stress in SH-SY5Y cells. SH-SY5Y cells were exposed to OGD for 4 h, and then reoxygenated in the presence of 40 μM physcion for 24 h. (A and B) Intracellular ROS generation was assessed by flow cytometry. (C–E) Levels of SOD, MDA and GSH-Px in cells were detected with ELISA. ** $P < 0.01$ compared with control group; # $P < 0.05$, ## $P < 0.01$ compared with OGD/R group.

Abbreviations: DCFH-DA, 2,7-dichlorofluorescein diacetate; GSH-px, glutathione peroxidase; MDA, malondialdehyde; OGD/R, oxygen-glucose deprivation/reperfusion; SOD, superoxide dismutase.

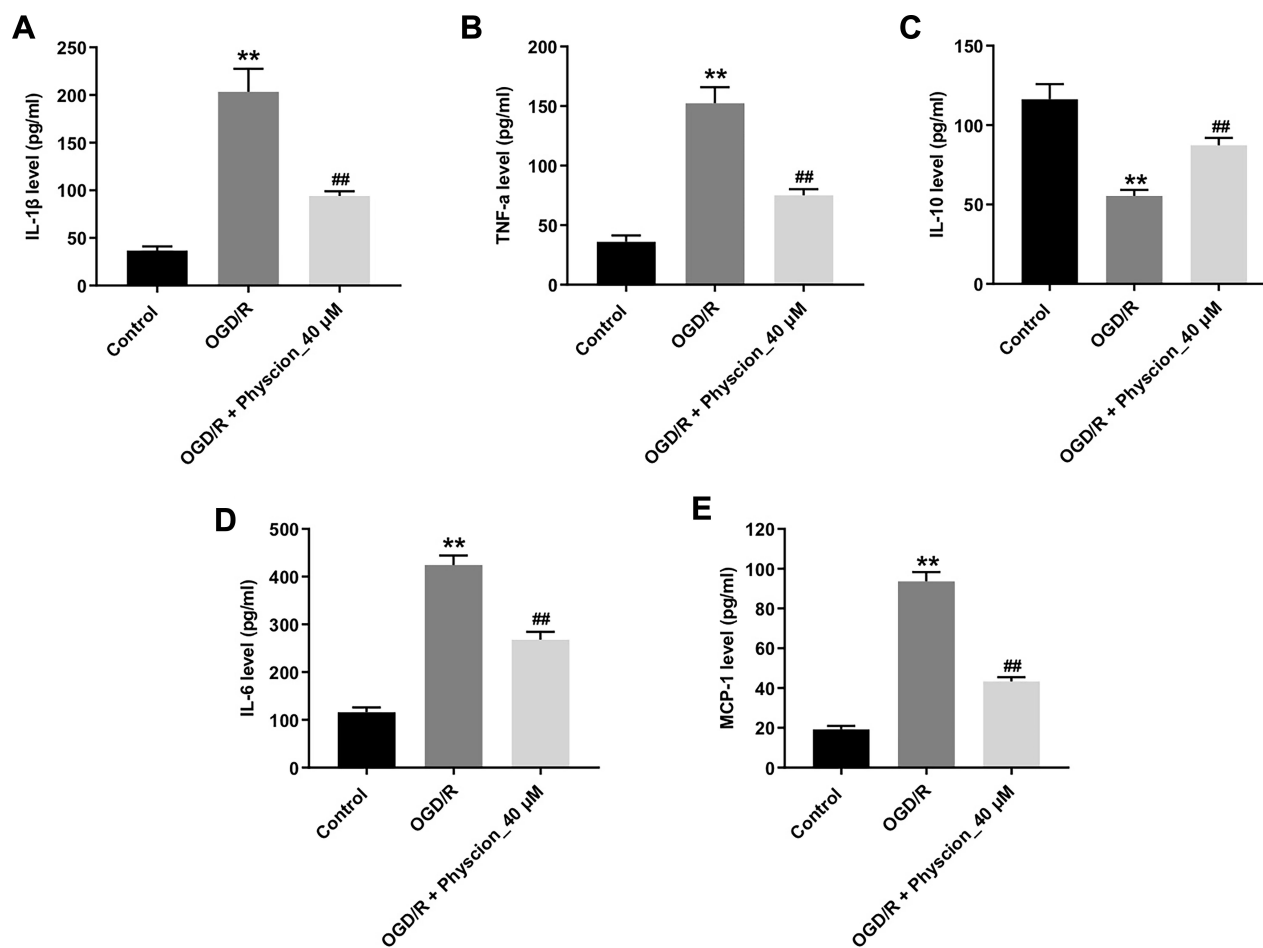


Figure 3 Physcion suppresses OGD/R-induced inflammatory response in SH-SY5Y cells. SH-SY5Y cells were exposed to OGD for 4 h, and then reoxygenated in the presence of 40 μ M physcion for 24 h. (A–E) Levels of IL-1 β , TNF- α , IL-10, IL-6 and MCP-1 in cells were detected with ELISA. ** $P < 0.01$ compared with control group; ## $P < 0.01$ compared with OGD/R group.

Abbreviations: IL-10, interleukin-10; IL-1 β , interleukin-1 β ; IL-6, interleukin-6; MCP-1, macrophage chemoattractant protein 1; OGD/R, oxygen-glucose deprivation/reperfusion; TNF- α , tumor necrosis factor- α .

Physcion Attenuates OGD/R-Induced Injury in SH-SY5Y Cells via Inhibiting the TLR4/NF- κ B Pathway

Evidence has shown that suppression of TLR4/NF- κ B signaling could improve cerebral I/R injury.¹⁸ Thus, to explore whether physcion protected OGD/R-induced neuronal cell injury via mediating the TLR4/NF- κ B pathway, Western blot assay was performed. As indicated in Figure 4A–D, the expressions of TLR4, p-p65 and p-I κ B were notably upregulated in OGD/R-treated SH-SY5Y cells; however, these OGD/R-induced changes were significantly reversed by physcion treatment in the course of reperfusion. These data indicated that physcion could attenuate OGD/R-induced injury in SH-SY5Y cells via inhibiting the TLR4/NF- κ B pathway.

Physcion Protects Rats Against Brain Damage After I/R via Inhibiting the TLR4/NF- κ B Pathway

A rat model of focal cerebral ischemia was established to explore the role of physcion in IS in vivo. To explore whether physcion is capable of crossing the blood-brain barrier (BBB) in rat brain, HPLC-MS/MS was performed to measure the level of physcion in CSF. The results indicated that the concentration of physcion in rat CSF was 196 ng/mL in I/R + physcion_40 mg/kg group, confirming that physcion was able to penetrate the BBB (Supplementary Figure 1A and B). In addition, TTC staining showed that the rats in I/R group exhibited marked areas of infarction (Figure 5A and B, Supplementary Table 1). However, physcion treatment obviously decreased areas of infarction compared to I/R group (Figure 5A and B, Supplementary Table 1).

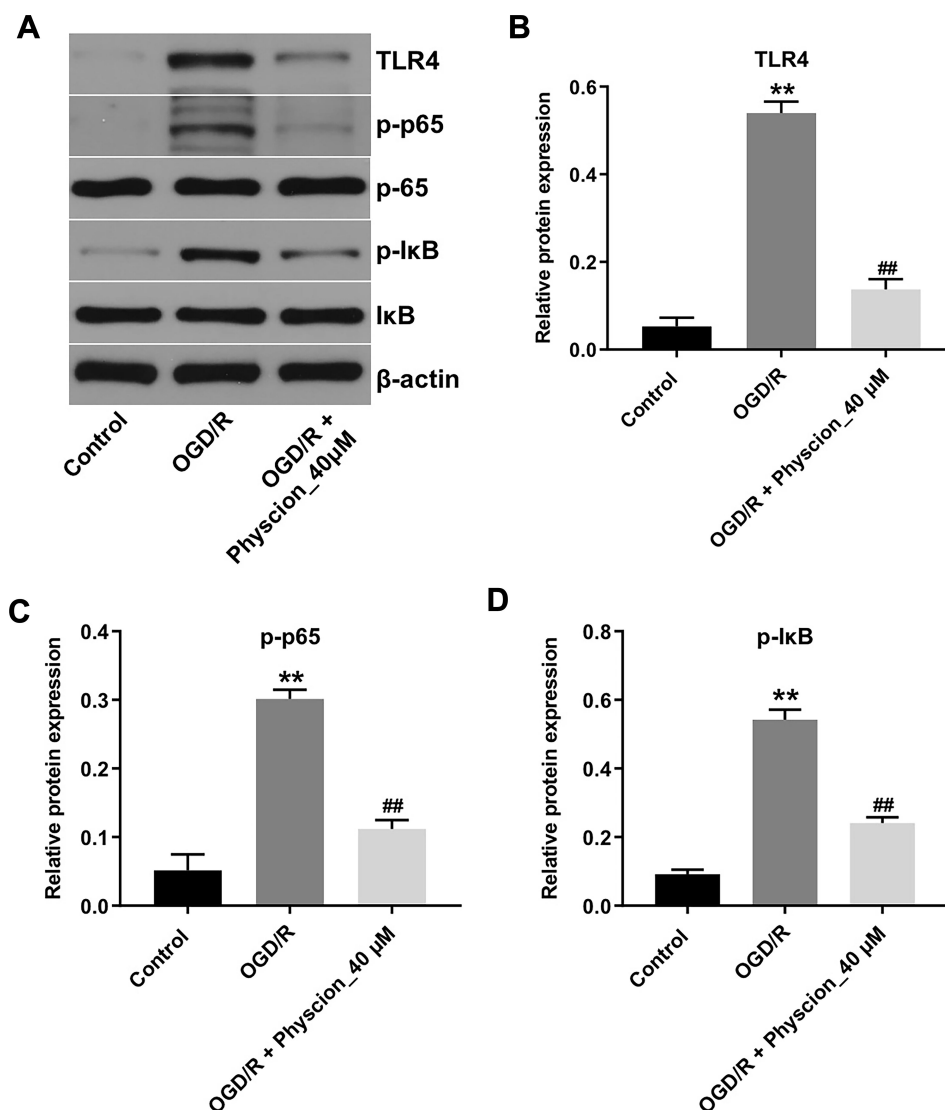


Figure 4 Physcion attenuates OGD/R-induced injury in SH-SY5Y cells via inhibiting the TLR4/p65 pathway. SH-SY5Y cells were exposed to OGD for 4 h, and then reoxygenated in the presence of 40 μ M Oroxylin A for 24 h. (A) Expression levels of TLR4, p-p65, p-I κ B in SH-SY5Y cells were detected with Western blotting. (B–D) The relative expressions of TLR4, p-p65, p-I κ B in cells were quantified via normalization to β -actin, p65 and I κ B. ** $P < 0.01$ compared with control group; ## $P < 0.01$ compared with OGD/R group.

Abbreviations: OGD/R, oxygen-glucose deprivation/reperfusion; p-I κ B, phospho-I κ B; p65, NF- κ B p65; p-p65, phospho-NF- κ B p65; TLR4, toll-like receptor 4.

Meanwhile, physcion significantly reduced the brain water content and the NSS in I/R rats, compared to I/R group (Figure 5C and D, Supplementary Tables 2 and 3). Moreover, HE and TUNEL assays showed that physcion treatment had protective effects against tissue injury in the brain slice of rats in the condition of cerebral I/R injury (Figure 5E and F, Supplementary Table 4).

Furthermore, the expressions of TLR4, p-p65 and p-I κ B were markedly upregulated in whole brain tissue of I/R rats; however, these changes were reversed by physcion treatment in the course of reperfusion (Figure 6A–D). All these data illustrated that physcion could protect rats against brain

injury after cerebral I/R via inhibiting the TLR4/NF- κ B pathway.

Discussion

It has been shown that physcion plays a therapeutic role in central nervous system (CNS) diseases, including cerebral IS.¹⁹ In this study, we performed a study to investigate the regulatory effect of physcion in cerebral I/R injury. Our data indicated that OGD/R significantly reduced the viability, increased the release of LDH and cell apoptosis in SH-SY5Y cells; however, these effects were reversed by the treatment with physcion (10, 20 or 40 μ M).

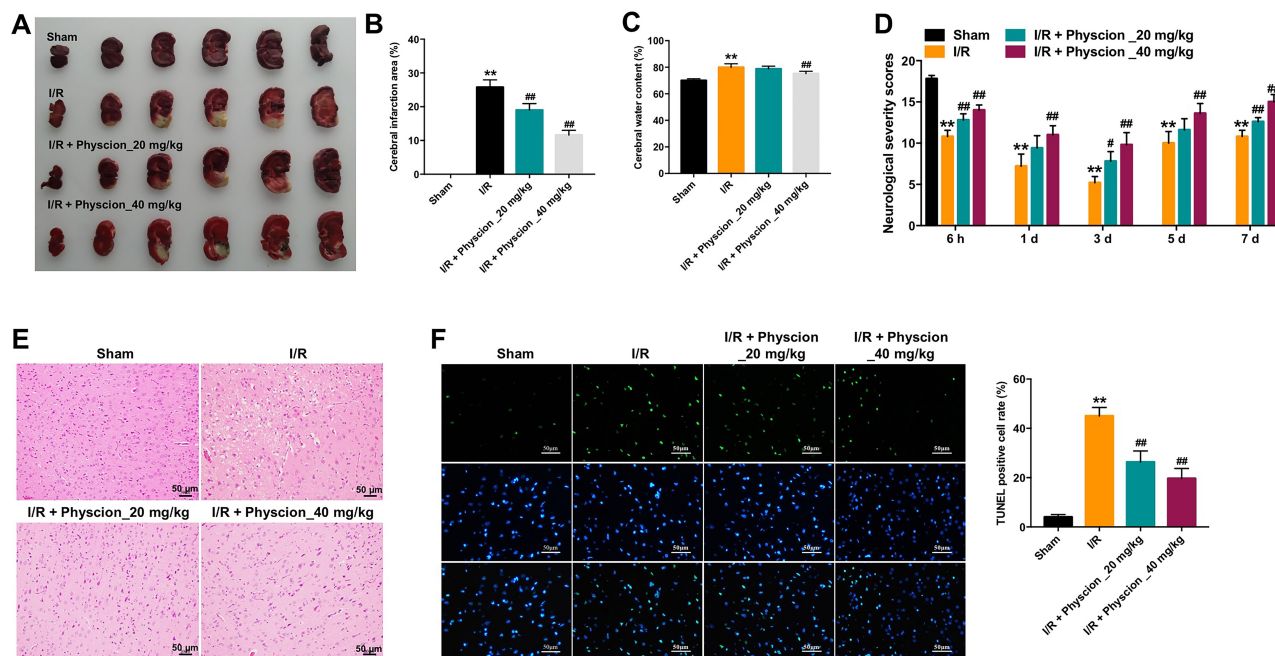


Figure 5 Physcion protects rats against brain damage after I/R. (A and B) TTC staining was applied to assess infarct area (magenta: healthy tissue; white: damaged tissue). (C) Cerebral water content was calculated. (D) Neurobehavioral tests were performed at 6 h, 1, 3, 5 or 7 days after MCAO. Neurological severity scores (NSS). (E) HE staining assay was performed to detect brain tissue injury in I/R rats (magnification x 200). (F) TUNEL assay was performed to detect cell apoptosis in I/R rats challenged with MCAO (magnification x 400). **P<0.01 compared with sham group; #P<0.05, ###P<0.01 compared with I/R group.

Abbreviations: I/R, ischemia-reperfusion; TUNEL, terminal deoxynucleotidyl transferase dUTP nick end labeling.

Importantly, 40 μ M physcion exhibited the most pronounced effect in preventing OGD/R-induced SH-SY5Y cell injury. In addition, our results indicated that physcion could attenuate I/R-mediated neuron injury via attenuating oxidative stress, inflammatory response, and inactivating the TLR4/NF- κ B pathway. Consistently, Li et al found that 26.4 mg/kg physcion exhibited anti-ischemic effect in rats with cerebral ischemia.¹³ In this study, we found that

physcion (20 or 40 mg/kg) protected against cerebral I/R injury in rats via inactivating the TLR4/NF- κ B pathway. These data indicated that physcion could protect against cerebral I/R injury in vitro and in vivo via inhibition of TLR4/NF- κ B signaling.

ROS generation is implicated in the I/R injury, increased ROS could lead to neuronal cell death.^{16,20} In addition, oxidative damage can consume intracellular

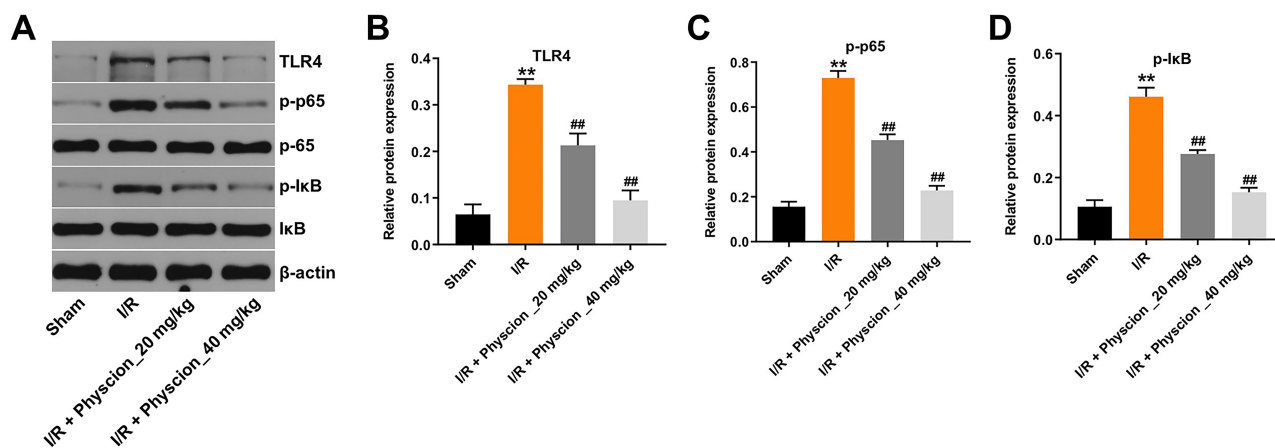


Figure 6 Physcion protects rats against brain damage after I/R via inhibiting the TLR4/p65 pathway. (A) Expression levels of TLR4, p-p65, p-IκB in the whole brain tissues of I/R rats were detected with Western blotting. (B–D) The relative expressions of TLR4, p-p65, p-IκB in the whole brain tissues of I/R rats were quantified via normalization to β -actin, p65 and IκB. **P<0.01 compared with sham group; ###P<0.01 compared with I/R group.

Abbreviations: I/R, ischemia-reperfusion; p-IκB, phospho-IκB; p65, NF- κ B p65; p-p65, phospho-NF- κ B p65; TLR4, toll-like receptor 4.

antioxidants (GSH and SOD), and lead to the production of MDA.^{21,22} Evidences have shown that cerebral I/R injury could increase ROS generation and induce oxidative stress by suppressing the activities of antioxidant biomarkers,^{23,24} which were again confirmed in the current study. Sun et al found that cannabidiol can protect hippocampal neurons against OGD/R-induced injury via attenuating oxidative stress.²⁵ Wu et al indicated that curcumin could attenuate OGD/R-induced injury in SH-SY5Y cells via increasing the levels of SOD and GSH.²⁶ In this study, we found that physcion could reduce OGD/R-induced oxidative stress in SH-SY5Y cells, as shown by the decreased generations of ROS and MDA, and the increased activities of GSH and SOD. These data suggested that physcion could protect SH-SY5Y cells against OGD/R-induced oxidative stress. In addition, consistent with previous studies,^{27,28} we found that OGD/R insult significantly reduced cell viability via inducing apoptosis, which was reversed by the administration of physcion.

Inflammatory mediators are key contributing factors in cerebral ischemic injury.^{29,30} It has been shown that TLR4 contributes to I/R-induced inflammatory response.³¹ Our data indicated that the expression of TLR4 was increased in SH-SY5Y cells exposed to OGD/R and in brain tissues of rats subjected to I/R injury. In addition, NF- κ B is an essential transcription factor that plays an important role in cell growth and inflammation.^{9,32} NF- κ B exists in cytoplasm as an inactive dimeric form binding to I κ B.³¹ TLR4/MyD88 signaling can phosphorylate I κ B, and then release the active form of NF- κ B (NF- κ B translocation from the cytoplasm to the nucleus).³³ NF- κ B participates in the pathological process of IS, which can be triggered by ROS, and several inflammatory cytokines.³² Moreover, it also has the ability to regulate the production and release of pro-inflammatory factors, including IL-1 β , IL-6, and TNF- α in neurons.³⁴ Ye et al found that meisoindigo protected against cerebral I/R injury by inhibiting TLR4/NF- κ B signaling pathway.¹⁸ Zhu et al indicated that berberine attenuated I/R injury via inhibiting NF- κ B nuclear translocation.⁷ Meanwhile, physcion was reported to possess an anti-inflammatory effect in LPS-stimulated HepG2 cells.³⁵ In this study, we found that TLR4, p-p65, and p-I κ B proteins, and inflammatory factors, including IL-1 β , TNF- α , IL-6 and MCP-1, were all upregulated in SH-SY5Y cells during OGD/R injury. However, these OGD/R-induced changes were significantly reversed by physcion treatment in the course of reperfusion, indicating that physcion-inhibiting inflammatory responses were likely to contribute to attenuating cerebral I/R injury.

Collectively, physcion attenuated OGD/R-induced inflammatory response in SH-SY5Y cells via inhibition of TLR4/NF- κ B pathway. Furthermore, wang et al found that curcumin could reduce the apoptosis and inflammation of PC12 cells after OGD/R insult via inactivating the TLR4/NF- κ B pathway.³⁶ In addition, Gao et al indicated that pterostilbene could inhibit oxidative stress in a mouse model of acute renal I/R injury via inhibition of TLR4/NF- κ B pathway.³⁷ These data indicated that physcion could attenuate the neuronal apoptosis, oxidative stress and inflammatory response induced by I/R injury via inactivating the TLR4/NF- κ B signaling.

Conclusion

In summary, the present study demonstrated that physcion could protect against rats' cerebral I/R injury via inhibition of the TLR4/NF- κ B signaling. Our findings might provide a potential therapeutic strategy for IS.

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Disclosure

The authors declare no competing financial interests.

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