Berberine elevates mitochondrial membrane potential and decreases reactive oxygen species by inhibiting the Rho/ROCK pathway in rats with diabetic encephalopathy

Lin Tian¹, Hong Ri¹, Jiping Qi¹, and Peng Fu²

Abstract

Objectives: Diabetic encephalopathy (DE) is a serious complication of diabetes mainly occurring in the elderly patients. Berberine (BBR) is an isoquinoline alkaloids extracted from Coptis chinensis that is applied in the treatment of diabetes clinically. This study explored the possible mechanism of BBR in relieving DE.

Methods: Wistar rats were injected with streptozotocin and fed a high fat diet to establish the model of DE. The model rats were treated with BBR. The body weight, blood glucose and insulin of rats were measured, and Morris water maze test was conducted to evaluate the learning and memory abilities. The pathological conditions of cortical tissues were detected. The cortical mitochondria membrane potential (MMP) and reactive oxygen species (ROS) were monitored. The expressions of Rho/ROCK pathway-related genes of rat cortex were detected. The changes of MMP and ROS were detected after the treatment of Rho/ROCK pathway activator.

Results: The body weight of model rats changed little, and levels of blood glucose and insulin were increased. The spatial learning and memory abilities were impaired, with disordered cortical neurons, and obvious neurons apoptosis and glia proliferation. BBR alleviated cognitive dysfunction and pathological damage in rats with DE. BBR enhanced cortical MMP and suppressed ROS. BBR treatment inhibited the Rho/ROCK pathway. Activation of the Rho/ROCK pathway reversed the effects of BBR on MMP and ROS.

Conclusion: BBR elevated MMP and reduced ROS in rats with DE by inhibiting the Rho/ROCK pathway activator. This study may offer novel insights for the management of DE.

Keywords

Diabetic encephalopathy, berberine, mitochondria membrane potential, reactive oxygen species, Rho/ROCK pathway

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Introduction

Diabetes represents a group of heterogeneous metabolic disorders characterized by hyperglycemia in common.¹ Diabetes may bring about serious complications such as cardiovascular diseases, blindness and encephalopathy.² Diabetic encephalopathy (DE) is defined as the cognitive impairments and neurophysiological or structural alterations in the brain resulted from diabetes,³ and it is closely concerned with the degeneration and dysfunction of the central nervous system.⁴ Clinically, DE is mainly manifested as learning and memory impairments, which
may eventually lead to dementia. The pathogenesis of DE has not been clarified completely, but it is acknowledged that DE is related to cerebrovascular abnormalities, oxidative stress and insulin abnormalities. With the increasing incidence of diabetes and proportion of the elderly, elucidating the molecular mechanism of DE and developing potent agents for diabetes patients are urgent for improving the patients’ quality of life and reducing the burden of social medicare.

Mitochondrial dysfunction is demonstrated to be implicated in the pathological progression of diabetes and diabetic complications. Mitochondrial homeostasis disorder and mitochondrial membrane potential (MMP) polarization are presented in human mononuclear cells with type 2 diabetes. Moreover, mitochondria are the major source of reactive oxygen species (ROS), which are generally viewed as the toxic byproducts of aerobic metabolism. High level of ROS is induced in the diabetic environment, leading to the brain tissue damages associated with diabetic complications. Therefore, restoring mitochondrial homeostasis may also be a breakthrough to ameliorate DE.

Chinese herbal medicine bears a bright clinical application prospect in the management of diabetes with the advantages of less toxicity and side effects. Berberine (BBR) is an isoquinoline alkaloid derived from Rhizoma Coptidis, Cortex Phellodendri, and Cortex Berberidis, which possesses multiple pharmacological effects such as improving glycolipid metabolism and alleviating insulin resistance. Notably, BBR can exert benign effects as improving glycolipid metabolism and alleviating insulin abnormalities. which possesses multiple pharmacological effects such as correcting the toxic byproducts of aerobic metabolism. Hence, BBR may exert therapeutic effects on high glucose-induced vascular inflammation. Whether BBR affects cortical MMP and ROS in DE via the Rho/ROCK pathway remains unclear. This study herein investigated the effects of BBR and Rho/ROCK on MMP and ROS in rats with DE, which shall shed lights on the development of pharmacotherapy for DE.

**Materials and methods**

**Ethics statement**

The study got the approval of the Ethical Committee of The First Affiliated Hospital of Harbin Medical University. All experimental procedures were implemented on the Ethical Guidelines for the study of experimental pain in conscious animals.

**Animal treatment**

Male Wistar rats (180–200 g, 6–8 weeks) were purchased from Beijing Vital River Laboratory Animal Technology Co., Ltd, (Beijing, China) [SYXK (Beijing) 2017-0033]. The rats were raised in a specific pathogen-free animal laboratory with 2 rats per cage. The rats were fed with a common chow for 2 weeks and then a high glucose and high fat diet (composed of 67.5% standard laboratory rat feed, 20% sugar, 10% lard, 2% cholesterol and 0.5% bile salt) for 4 weeks. In the 5th week, the rat model of DE was induced by injection of 25 mg/Kg streptozotocin (STZ) via tail vein. In the 6th week, oral glucose tolerance test (OGTT) was performed. The rats were given 2 g/kg glucose solution according to body weight, and then the blood glucose was detected using a glucose analyzer (ACCU-CHEK, Roche Diagnostics, Basel, Switzerland). The fasting blood glucose after fasting for 12h, and postprandial blood glucose at 1 h and 2 h after meal were measured. The rats with at 2 h postprandial postprandial blood glucose over 11.2 mmol/L were regarded as successful modeled rats. The model rats were assigned into model group, BBR low-dose (BBR-L) group and BBR high-dose (BBR-H) group. The rats in the experimental groups were intragastrically administered with 50 mg/Kg/d BBR and 200 mg/Kg/d BBR in the 6th week, respectively. The model rats were intragastrically administered with equal amount of 5% CMC-Na (Macklin, Shanghai, China). The lateral ventricle of rats were injected with 0.01 mol/L Rho/ROCK pathway activator lysophosphatidic acid (LPA) (5 µL) within 20 min with a microinjector (LPA group), and the rats in the control group were injected with 0.01 mol/L phosphate-buffered saline (PBS). The injector was retained for 10 min and slowly withdrawn, and then the rats were sterilized and sutured. The rats in the normal group were fed with rodent diet and water. In the 9th week, all the rats were evaluated for 14–16 h, and their body weight was measured in the next morning.

**Homeostasis model assessment of insulin resistance (HOMA-IR)**

The blood drop (3 µL) of rats was obtained through tail docking, and blood glucose level was detected using the One Touch Ultra Link Glucometer (LifeScan Inc., Milpitas, CA, USA). The plasma insulin was detected in line with the instructions of ELISA kit (EZRMI-13, Millipore, MA, USA). Insulin resistance was assessed by the formula: HOMA-IR = fasting insulin (mU/L) × fasting blood glucose (mmol/L)/22.5.
Morris water maze test

The circular pool for water maze was 160 cm in diameter and 60 cm in depth. The platform was 11 cm in diameter. The water volume corresponding to the platform height was added to the pool in advance, and then 250 mL of ink was added to make the water opaque. The platform was placed according to the relative fixed position shown on the computer. The water temperature was heated to 22–25°C before the experiment. In the first two days of training period, the height of the platforms was arranged to be 1 cm below the water level. In the 3rd, 4th, 5th day of training period and the 6th day of testing period, the height of the platforms was arranged to be 1 cm above the water level. This experiment was mainly divided into positioning navigation (fixed escape platform) test and space exploration test. For the positioning navigation test, the rats were oriented to the pool wall from four quadrants (A, B, C and D) in turn; the rats were put into the pool along the wall with the head raised up, adapted for 1–2 s above the water surface, and then released gently. The motion trace of rats in the next 1 min was recorded, and the test time lasted 60 s. If the rats found the platform and landed within 60 s, the computer automatically recorded the escape latency, and the rats were allowed to stay on the platform for 15 s and then taken away. If the rats did not find the platform within 60 s, they were guided to land the platform and stay for 15 s, and the software automatically recorded the escape latency time as 60 s. The rats entered water from four different quadrants in turn in a least interval of 20 min. The operation was repeated for 6 consecutive days. The system automatically recorded the escape latency (time from water entry to landing platform). For space exploration test, the platform was removed, and the staying time of rats in the target quadrant within 60 s was recorded according to the above operations, and the percentage of rats staying in the target quadrant was counted.

Hematoxylin and eosin (HE) staining

After the Morris water maze test, the rats were anesthetized by intraperitoneal injection of pentobarbital sodium (50 mg/kg) and then perfused with 0.9% normal saline and 4% paraformaldehyde into the hearts. The rat brain was removed, and the cortical tissues were isolated and fixed with 4% paraformaldehyde, then immersed in 30% sucrose at 4°C for 24 h. The paraffin-embedded cortical tissues were fixed in coronal section and sliced at 5 μm. The sections were placed onto the 3-aminopropyl triethoxysilane-treated slides and baked in an oven at 60–65°C for 30–60 min. After dewaxing and dehydration, the sections were stained with hematoxylin for 10–15 min, with the excess dye solution gently washed by running water. Then, the sections were differentiated with 5% hydrochloric acid ethanol for several seconds, and washed with running water for 3–5 min. Next, the sections were stained with eosin solution for 5 min. Afterwards, the sections were dehydrated with gradient alcohol, cleared with xylene and sealed with neutral gum. The pathological changes of tissues were observed under a light microscope.

Reverse transcription quantitative polymerase chain reaction (RT-qPCR)

Total RNA was extracted from 100 mg cortical tissues using TRIzol reagent (12183555, Thermo Fisher Scientific, Jiangsu, China), and then the concentration and purity of RNA were determined. The EP tube was added with 5 μL Mix reagent (AM8228G, Invitrogen), 5 μL total RNA and 10 μL RNase free H2O, and then centrifuged and mixed. The mixture was reacted in the qPCR instrument at 37°C for 15 min and 85°C for 5 s; the reaction was terminated at 4°C for qPCR or stored at –20°C. Primers (Table 1) were designed by Primer Premier 5 and synthesized by Sangon Biotech (Shanghai, China). The RT-qPCR was performed using ABI 7500 quantitative PCR instrument (Applied Biosystems, Inc., Carlsbad, CA, USA) on the provided instructions of SYBR® Premix Ex Taq™ II kit (Takara Bio Inc., Kyoto, Japan). PCR was performed on the following conditions: pre-denaturation at 95°C for 15 min; 40 cycles of denaturing at 94°C for 30 s, annealing at 55°C for 30 s and extension at 72°C for 30 s. After amplification, the dissolution curve analysis was conducted. The relative expression of genes was calculated by 2^(-ΔΔCt) method, with glyceraldehyde-3-phosphate dehydrogenase (GAPDH) as the internal reference.

Western blotting

Cortical tissue homogenate of rats was added with protein lysate (R0278, Sigma-Aldrich, Merck KGaA, Darmstadt, Germany), and then centrifuged at 14000 g and 4°C for 20 min to collect the supernatants. The total concentration of proteins in the homogenate was determined by intraperitoneal injection of pentobarbital sodium (50 mg/kg) and then perfused with 0.9% normal saline and 4% paraformaldehyde into the hearts. The rat brain was removed, and the cortical tissues were isolated and fixed with 4% paraformaldehyde, then immersed in 30% sucrose at 4°C for 24 h. The paraffin-embedded cortical tissues were fixed in coronal section and sliced at 5 μm. The sections were placed onto the 3-aminopropyl triethoxysilane-treated slides and baked in an oven at 60–65°C for 30–60 min. After dewaxing and dehydration, the sections were stained with hematoxylin for 10–15 min, with the excess dye solution gently washed by running water. Then, the sections were differentiated with 5% hydrochloric acid ethanol for several seconds, and washed with running water for 3–5 min. Next, the sections were stained with eosin solution for 5 min. Afterwards, the sections were dehydrated with gradient alcohol, cleared with xylene and sealed with neutral gum. The pathological changes of tissues were observed under a light microscope.

Table 1. Primer sequence for RT-qPCR.

| Gene    | Sequence                           |
|---------|------------------------------------|
| Rho C   | F: 5'-AAATGGACCCCAAGGTTAGG-3'      |
|         | R: 5'-TTAATTCCCCACCAGATAGT-3'      |
| ROCK1   | F: 5'-AAAAATGGAACACCTCGTGC-3'      |
|         | R: 5'-GGAGAGAAAATCCAAACAT-3'       |
| ROCK2   | F: 5'-CGTGTACGGAGACCGCT-3'         |
|         | R: 5'-TTTCTTTCCTCAAGCAAGGA-3'      |
| GAPDH   | F: 5'-GTTGAAGGGTGAGTTACCGG-3'      |
|         | R: 5'-CCTGGAAGATGGTATGGATT-3'      |
determined. Then, 10% sodium dodecyl sulfate-polyacrylamide (SDS) separation gel and concentrated gel were prepared. The samples were mixed with the loading buffer, boiled at 100°C for 5 min, cooled in an ice-bath and centrifuged. Equal amount of protein on each lane was separated by SDS gel electrophoresis and transferred onto nitrocellulose membranes. The membranes were blocked with 5% skim milk at 4°C overnight. Afterwards, the membranes were cultured with the rabbit anti-mouse primary antibodies for 1 h: RhoC (1/2000, ab187026, Abcam Inc., Cambridge, MA, USA), ROCK1 (1/1000, ab134181, Abcam), ROCK2 (1/1000, ab125025, Abcam) and GAPDH (1/2000, ab181602, Abcam), following 3 PBS washes (5 min/time). Next, the membranes were cultured with secondary antibody goat anti-rabbit immunoglobulin G (IgG) (1/1000, ab205718, Abcam) for 1 h, following PBS washes (5 min/time). The membranes were immersed in the enhanced chemiluminescence reagent (Pierce, Waltham, MA, USA) for 1 min. The cortical tissues were cut into pieces in mitochondrial extraction buffer (225 mol/L sucrose, 75 mol/L sorbitol, 1 mol/L ethylene glycol tetraacetic acid, 5 mol/L Hepes, 0.1% fatty acid-free bovine serum albumin and 10 mmol/L Tris-HCl), homogenized and centrifuged at 200 g and 4°C for 5 min to collect the supernatants and remove the precipitate. The supernatants were centrifuged at 17000 g and 4°C for 30 min, with the precipitate kept and the supernatants removed. The precipitates were resuspended in a loading buffer containing 0.32 mol/L sorbitol and 5 mmol/L potassium phosphate, (pH = 7.5), and isolated and purified by sucrose gradient centrifugation. A small amount of acquired mitochondria were fixed with glutaraldehyde, embedded in epoxy resin, sectioned and stained with uranyl acetate. The mitochondria were identified under an electron microscope, and the mitochondria with purity above 90% were used in the subsequent experiments.

**Isolation and purification of mitochondria**

The cortical tissues were cut into pieces in mitochondrial extraction buffer (225 mol/L sucrose, 75 mol/L sorbitol, 1 mol/L ethylene glycol tetraacetic acid, 5 mol/L Hepes, 0.1% fatty acid-free bovine serum albumin and 10 mmol/L Tris-HCl), homogenized and centrifuged at 200 g and 4°C for 5 min to collect the supernatants and remove the precipitate. The supernatants were centrifuged at 17000 g and 4°C for 30 min, with the precipitate kept and the supernatants removed. The precipitates were resuspended in a loading buffer containing 0.32 mol/L sorbitol and 5 mmol/L potassium phosphate, (pH = 7.5), and isolated and purified by sucrose gradient centrifugation. A small amount of acquired mitochondria were fixed with glutaraldehyde, embedded in epoxy resin, sectioned and stained with uranyl acetate. The mitochondria were identified under an electron microscopy, and the mitochondria with purity above 90% were used in the subsequent experiments.

**Detection of MMP**

JC-1 is an ideal fluorescent probe widely used to detect MMP. MMP was detected using the JC-1 detection kit. The cells were incubated on the slides at 37°C for 24 h. Then the changes of MMP were observed under an inverted fluorescence microscope (IX51, Olympus, Tokyo, Japan). The average value of red fluorescence intensity of 5 visual fields was calculated using the Color Histogram of Image Pro Plus 6.0 software.

The average fluorescence intensity was used for statistical analysis to quantify the level of MMP.

**Detection of ROS in mitochondria**

ROS in mitochondria can oxidize dichlorofluorescein diacetate (DCFH-DA) to dichlorofluorescein (DCF), and the formation rate of DCF is proportional to the content of ROS in mitochondria. Therefore, the fluorescence intensity of DCF can be used to reflect the production of ROS in mitochondria. The cells were incubated with 10 μmol/L DCFH-DA at 37°C for 60 min and then the fluorescence intensity was detected under an inverted fluorescence microscope at an excitation wavelength of 488 nm and an emission wavelength of 527 nm.

**Detection of glutathione (GSH) and malondialdehyde (MDA) in mitochondria**

The concentration of GSH and MDA in mitochondria was detected using the GSH kit (S0053, Beyotime, Shanghai, China) and MDA kit (MAK085, Sigma-Aldrich).

**Statistical analysis**

Data analysis was introduced using the Prism 5 (GraphPad, San Diego, CA, USA). Data are expressed as mean ± standard deviation. The t-test was used for comparison between two groups. One-way or two-way analysis of variance (ANOVA) was adopted for the comparisons among multiple groups, followed by Tukey’s multiple comparisons test. The p value was obtained from a two-tailed test, and p < 0.05 meant a statistically significance.

**Results**

**The rat model of DE was induced by STZ**

The body weight, fasting blood glucose and plasma insulin levels of rats were measured periodically. One week after STZ injection (Figure 1(a)), a total of 70 rats with fasting blood glucose ≥11.2 mM were identified after 12 h of fasting, and the remaining 8 rats that did not meet the standard or died were excluded from the subsequent research. The body weight of rats with type 2 diabetes (T2D) encephalopathy was no longer changed notably after injection of STZ, while the weight of control rats continued to increase and eventually exceeded from the two-tailed test, and p < 0.01; Figure 1(b)). STZ injection increased the postprandial blood glucose level of rats with T2D encephalopathy from 6 mM to 28 mM within one week (Figure 1(c)). Plasma insulin and HOMA-IR were also notably promoted after injection of STZ.
To confirm the rat model of T2D encephalopathy, we conducted Morris water maze test to evaluate the learning ability of rats through hidden platform training (Figure 1(f) to (g)). We found that the swimming speed of rats remained unchanged (Figure 1(h)). However, the reference learning and procedural memory of rats with T2D encephalopathy were reduced; compared with the control rats, the rats with T2D encephalopathy needed longer latency time to reach the platform (all $p < 0.01$; Figure 1(i)). In addition, we also adopted space exploration test to assess the ability of the rats to maintain the spatial memory. Compared with the control rats, the rats with T2D encephalopathy had shorter retention time and less retention percentage in the target quadrant (all $p < 0.01$; Figure 1(j)). These results indicated that the spatial learning and memory ability of DE rats were impaired.

### BBR alleviated cognitive dysfunction and pathological damage in rats with DE

To determine whether BBR could improve memory impairment, we examined the effects of BBR on hyperglycemia and insulin resistance in rats with T2D encephalopathy. The results exhibited that 4 weeks of BBR treatment increased body weight and decreased blood glucose of rats with T2D encephalopathy (all $p < 0.01$; Figure 2(a) and (b)). The learning ability of rats was measured by hidden platform training (Figure 2(c) to (d)). The swimming speed of all the rats remained unchanged (Figure 2(e)). However, reference learning and procedural memory were reduced in rats with T2D encephalopathy. After training, the time to reach the platform in the T2D + BBR-H group was significantly less than that in T2D group (Figure 2(e)). BBR treatment also significantly reduced the escape latency of rats (all $p < 0.01$; Figure 2(f)) and increased the retention time in target quadrant (all $p < 0.01$; Figure 2(g)). These results suggested that BBR improved the impaired spatial learning and memory ability of rats with T2D encephalopathy. HE staining revealed that the rats with T2D encephalopathy showed obvious pathological changes of cortical tissues due to long-term hyperglycemia, disordered cortical neurons large neuron apoptosis and glial proliferation. However, the control rats had normal number of cortical neurons, without obvious neuronal apoptosis and glial proliferation, while neuronal apoptosis and glial...
proliferation were significantly reduced after BBR treatment (Figure 2(h)).

**BBR increased cortical MMP and decreased ROS in rats with DE**

Insulin resistance is the major factor leading to cognitive dysfunction of DE, and is also related to mitochondrial dysfunction and adenosine triphosphate decline. Therefore, we measured the MMP and levels of ROS, GSH and MDA in rat cortex. The MMP of rats with T2D encephalopathy was lower than that of control rats, and the MMP of BBR-treated rats was promoted with the increase of BBR concentrations (all $p < 0.01$; Figure 3(a)). ROS level in the cortical mitochondria of rats with T2D encephalopathy was notably increased, and BBR treatment lowered the ROS level with the increase of BBR concentrations (all $p < 0.01$; Figure 3(b)). Moreover, rats with T2D encephalopathy showed lower GSH level and higher MDA level compared with the control rats, and BBR effectively promoted the production of GSH and inhibit the production of MDA (all $p < 0.01$; Figure 3(c) and (d)).

**BBR inhibited the Rho/ROCK pathway in rats with DE**

BBR improves the behaviors of rats with DE via the SIRT/ER pathway, and BBR can alleviate the diabetes-related memory impairments. The renal protective effect of BBR on diabetic rats partly depends on RhoA/ROCK inhibition. Rho/ROCK pathway has been demonstrated to play a vital part in DE and participate in the production of ROS. RhoA/ROCK also participates in the pathogenesis of long-term complications of diabetes mellitus. Whether BBR improves DE via the Rho/ROCK pathway needs further exploration. The expressions of the Rho/ROCK pathway-related genes (RhoC, Rock1 and Rock2) were detected. The results revealed that the expressions of RhoC, Rock1 and Rock2 of rats with T2D encephalopathy were higher than those of the control rats; compared
with those of rats in the T2D group, the expressions of RhoC, Rock1 and Rock2 of rats in the T2D + BBR-L and T2D+BBR-H groups were significantly reduced (all \( p < 0.01 \); Figure 4(a) and (b)). Briefly, BBR effectively inhibited the Rho/ROCK pathway.

**Figure 3.** BBR increased cortical mitochondrial membrane potential (MMP) and decreased ROS level. (a) Fluorescence intensity of MMP in JC-1 stained cells was observed under an inverted fluorescence microscope; (b) fluorescence intensity of ROS in DCFH-DA stained cells (200 ×); (c/d) levels of GSH and MDA in rat cortical mitochondria were detected using GSH detection kit and MDA detection kit. \( N = 6 \). The experiment was repeated three times. Data are expressed as mean ± standard deviation. One-way ANOVA was applied to assess the data, followed by Tukey’s multiple comparisons test. *** \( p < 0.001 \) vs. the control group; * \( p < 0.05 \), ** \( p < 0.01 \), *** \( p < 0.001 \) vs. the T2D group.

**Figure 4.** BBR inhibited the Rho/ROCK pathway in rats with DE. (a/b) Levels of RhoC, ROCK1 and ROCK2 were detected using RT-qPCR and Western blotting. \( N = 6 \). The experiment was repeated three times. Data are expressed as mean ± standard deviation. One-way ANOVA was applied to assess the data, followed by Tukey’s multiple comparisons test. *** \( p < 0.001 \) vs. the control group; # \( p < 0.01 \), ## \( p < 0.001 \) vs. the T2D group.

**BBR increased MMP and decreased ROS in rats with DE by inhibiting the Rho/ROCK pathway**

To further confirm that BBR inhibited ROS production by regulating the Rho/ROCK pathway, we injected
LPA, an activator of the Rho/ROCK pathway, into BBR-treated rats. The protein levels of RhoC, Rock1 and Rock2 were notably promoted after LPA treatment (all \( p < 0.01 \); Figure 5(a)). Compared with that in the T2D + BBR-H group, the rats in the T2D + BBR-H + LPA group showed reduced MMP and elevated ROS level (all \( p < 0.01 \); Figure 5(b) and (c)), as well as decreased GSH level and increased MDA level (all \( p < 0.01 \); Figure 5(d) and (e)). These results suggested that BBR increased MMP and decreased production of ROS by inhibiting the Rho/ROCK pathway in rats with DE.

**Discussion**

BBR regulates glucose metabolism through a variety of mechanisms and signaling pathways, such as enhancing insulin sensitivity and increasing glucose transporters, thereby playing a crucial role in the treatment of diabetic complications including DE. The present study demonstrated that BBR exerted protective effects on rats with DE by increasing MMP and reducing ROS.

BBR is endowed with multiple pharmacological activities, especially the glucose- and cholesterol-lowering properties. This study exhibited that 4 weeks of BBR

![Figure 5](image-url)
treatment decreased blood glucose of rats with DE, which is consistent with the results revealed by a previous study. BBR also participates in insulin signaling pathway and improves insulin resistance to ameliorate cognitive impairments. We conducted Morris water maze test to examine the effect of BBR on cognitive function in rats with DE. The results demonstrated that BBR improved the impaired spatial learning and memory ability of rats with DE. Consistently, Wang et al. have demonstrated that BBR improves cognitive dysfunction in rats with DE by alleviating tau hyperphosphorylation and axonal injury. Increased neuronal apoptosis is responsible for the impaired learning and memory ability of diabetic mice. Additionally, the extensive proliferation of glial cells around the brain injury sites results in glial scarring, which may be the main obstacle to neuronal regeneration. We showed that neuronal apoptosis and glial proliferation in rats with DE were significantly reduced after BBR treatment. Sedaghat et al. have also revealed that BBR inhibits hippocampal CA3 neuronal loss and exerts neuroprotective effects on epileptic rats. In brief, BBR alleviated cognitive dysfunction and pathological damage in rats with DE.

Insulin resistance contributes to the major risk factor for diabetic cognitive impairments, and mitochondrial dysfunction is demonstrated to promote insulin resistance. Generally, mitochondrial dysfunction is related to the decrease of MMP and the increase of intracellular and mitochondrial ROS. In this study, we showed that the MMP of BBR-treated rats was significantly promoted with the increase of BBR concentrations. BBR combined with glucose stimulation can prevent mitochondrial depolarization and restore negative membrane potential in diabetic rats. The pathogenesis of diabetic complications involves excessive oxidative stress caused by overproduction of ROS and deficiency of insulin transduction pathway. Enhanced ROS directly affects the synaptic activity and neurotransmission of neurons, which resulting in cognitive dysfunction. ROS level in the cortical mitochondria of rats with DE was notably increased, and BBR treatment could reduce the ROS level. BBR can attenuate ROS production and reduce glucose neurotoxicity in high glucose-treated cells. These results indicated that BBR increased cortical MMP and decreased ROS in rats with DE.

Thereafter, we shifted to exploring the signaling pathway of BBR improving cognitive impairments in rats with DE. RhoA/ROCK is reported to be implicated in the pathogenesis of diabetic complications and ROCK inhibitor is viewed as a promising target for the management of diabetic complications. Importantly, the Rho/ROCK pathway is involved in the regulation of diabetes-induced cognitive impairments. We detected the expressions of Rho/ROCK pathway-related genes in BBR-treated rats, and found that BBR could effectively inhibit the Rho/ROCK pathway. Decreased ROCK2 expression can improve cognitive ability in animal models of diabetic dementia. Moreover, we injected LPA into BBR-treated rats to further confirm that BBR protected against DE via the Rho/ROCK pathway. The results revealed that activating the Rho/ROCK pathway reversed the effects of BBR on MMP and ROS. Inhibition of the Rho/ROCK pathway can decrease the production of ROS. Briefly, BBR increased MMP and decreased production of ROS by inhibiting the Rho/ROCK pathway in rats with DE.

Conclusion

To sum up, BBR elevated MMP and reduced ROS in rats with DE by inhibiting the Rho/ROCK pathway. Our data are expected to contribute to clarifying the potential protective mechanism of BBR in DE. In the future, we shall carry out more prospective trials to refine the clinical application of BBR in DE.

Data Availability

All the data generated or analyzed during this study are included in this published article.

Authors’ Contributions

Lin Tian is the guarantor of integrity of the entire study; Lin Tian contributed to the study concepts, study design, definition of intellectual content, literature research, manuscript preparation and manuscript editing and review; Hong Ri contributed to the clinical studies; Lin Tian and Jiping Qi contributed to the experimental studies and data acquisition; Peng Fu contributed to the data analysis and statistical analysis. All authors read and approved the final manuscript.

Declaration of Conflicting Interests

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