Anti-Oxidant Effect of Lycium Barbarum Polysaccharides on the Retina of Streptozotocin-induced Diabetes Rats

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Introduction

Diabetic retinopathy is one of the most common complications of diabetes and its pathogenesis remains unclear. Studies have shown that mitochondria oxidative damage in retinal nerve cells and neuroglia cell may be the common pathway for the occurrence and development of diabetic retinopathy [1]. Lycium barbarum polysaccharides (LBP) is extracted from the traditional Chinese herbs medicine, medlar, which has a strong anti-oxidant effect [2]. In this study, LBP intervention was used in streptozotocin-induced diabetes rats to confirm the anti-oxidant effect through observations of the changes of retina damage.

Materials and Methods

Materials Twenty SPF male SD rats from Shanghai SLAC Laboratory Animal Co. Ltd. were used. No diseases were found in the outer eye and fundus oculi. The rats were raised in the Rodentia SPF Laboratory of the Experimental Animal Center of Fujian Medical University. Blood glucose was measured by the OneTouch Ultra2 Blood Glucose Meter (≤6.75mmol/L). Streptozotocin was dissolved in sterile citric acid–citrate buffer solution. It was prepared in 1% solution before use. LBP (1.05g) which was provided by prof. LY Huang was weighed for dissolution in 17.5ml physiological saline.
Method

Establishment of Streptozotocin-Induced Diabetic Rat Model

The [3] SD rats were randomly divided into the DM group (15 rats) and the control group (5 rats). They were raised adaptively for 1 week, and fasted for 10 hours before modeling. Streptozotocin solution was injected into rats in the DM group via the lower left abdominal cavity (65mg·kg⁻¹). Citric acid–citrate buffer solution of the same quantity was injected into 5 rats in the control group. 72 hours after, blood was taken from the caudal vein to measure blood glucose. When the rats showed a blood glucose level ≥ 16.7mmol·L⁻¹, the measurement of blood glucose level was repeated after 1 week. The rats which blood glucose level ≥ 16.7 mmol·L⁻¹ were picked out for the following experiment.

Experimental Grouping and Observation

Thirteen rats, which passed the modeling, were randomly divided into two groups: the diabetic model group (DM group) and the LBP treatment group (LBP group). Every morning, LBP was intragastrically administered to the rats of the LBP group, whereas physiological saline was intragastrically administered to the DM group and the control group. Body weight and blood glucose were measured every four weeks.

Observation of SOD and MDA Level of Retina

The rats were weighed and blood glucose were measured after drug administration for 24 weeks. The rats were intraperitoneally injected with napental(60mg·Kg⁻¹) and the eyeballs were removed. The retina was peeled under a ophthalmologic microscope. The retina tissue was added into the pre-cooled physiological saline at 4°C and then cleaned twice. The physiological saline at 4°C was added at a ratio of 1:9. The solution was then homogenized and centrifugated (2000rpm, 10min). The supernatant fluid was retrieved for measurements with a reagent kit, according to the manufacturer’s instructions.

Expression of VEGF mRNA in the Rat Retina Tested by RT-PCR

The eyeballs of the rats in each group were taken to peel the retina under a microscope for ophthalmologic operation. The TRIzol method was used to extract total RNA and prepare cDNA by reverse transcription. The cDNA was PCR-amplified, and then agarose gel electrophoresis was performed. A gel imaging system was used for gray scale scanning. The absorbance value was recorded and the VEGF/β-Actin specific value was determined.

Ultrastructure Observation by Transmission Electron Microscopy

Glutaraldehyde myocardial perfusion was performed and the eye ball was rapidly removed. Retina tissue was taken and cut into small rectangular pieces, which were then fixed in glutaraldehyde-potassium ferrocyanide solution overnight. After fixing in osmic acid-potassium ferrocyanide for cleaning, alcoholized uranyl acetate dye solution was used for En Bloc staining. Alcohol–acetic acid was used for dehydration in the gradient slope. Epoxy resin embedding medium was used for embedding. The specimens were sliced into ultrathin sections. Uranyl acetate and lead citrate were used for staining; they were observed under a Philips transmission electron microscope.

Statistical Analysis

Data were represented as mean±SD. Statistical analysis was performed by SPSS13.0 statistical software. Statistical differences between groups were analyzed by one-way analysis of variance (ANOVA). P<0.05 was considered statistically significant.

Results

The difference of Body Weight During Observation

The body weight of the rats increased in control group whereas decreased in DM group and LBP group (P<0.01 VS control group). The change of body weight is the same level between DM group and LBP group (P=0.984) (Table 1, Figure 1).

Table 1: The difference of body weight during observation.

| Grouping      | The body weight before treatment(g) | The body weight after treatment(g) | The difference of body weight(g) | P   |
|---------------|-------------------------------------|-----------------------------------|----------------------------------|-----|
| control group | 281.0±11.81                         | 564.9±38.61                      | 283.9±33.31                     | <0.01*|
| DM group      | 283.7±10.14                         | 254.3±31.16                      | -29.4±29.60                     | <0.01*|
| LBP group     | 283.0±13.46                         | 253.3±41.32                      | -29.7±35.49                     | 0.984#|

*P<0.01 compare with the control group. #P=0.984 compare between the LBP group and DM group.
The Variation of Blood Glucose During Treatment

The blood glucose of the rats in DM group and LBP group is higher than the glucose in control group ($P<0.01$ VS control group). The blood glucose is the same level between DM group and LBP group ($P=0.503$) (Table 2, Figure 2, 3).

**Figure 1.**

**Figure 2.**
Table 2: The variation of blood glucose during treatment.

| Grouping    | Before Treatment | After Treatment |
|-------------|------------------|-----------------|
|             | Blood glucose (mmol/L) | P     | Blood glucose (mmol/L) | P     |
| control     | 4.97±0.75         | <0.01*         | 4.84±0.60         | <0.01*         |
| DM group    | 23.01±2.44        | <0.01*         | 27.49±2.79        | <0.01*         |
| LBP group   | 22.54±2.19        | <0.01*         | 26.66±3.64        | <0.01*         |

*P<0.01 compare with the control group. #P=0.503 and P=0.654 compare between the LBP group and DM group.

The SOD and MDA Level of Retina

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The SOD of retina in DM group decreased and the MDA increased (P<0.05, VS control group). The SOD of retina in LBP group increased and the MDA decreased (P<0.05, VS DM group) (Table 3, Figure 4, 5).

Table 3: The effect of LBP on the SOD and MDA of DM rats retina.

| Grouping  | SOD activity (U·mgprot-1) | SOD P* | SOD P# | MDA content (mol·mgprot-1) | MDA P* | MDA P# |
|-----------|--------------------------|--------|--------|-----------------------------|--------|--------|
| Control   | 171.51±4.79              |        |        | 4.12±0.43                   |        |        |
| DM        | 129.70±6.97              | <0.05  | <0.05  | 8.23±0.43                   | <0.05  | <0.05  |
| LBP       | 179.82±4.02              | <0.05  | <0.05  | 4.58±0.25                   | >0.05  | <0.05  |

*Compare with control group. #Compare between DM group and LBP group.

Expression of VEGF mRNA by RT-PCR

The expression of VEGF mRNA in the rat retina was highest in DM group and lowest in control group. The level of VEGF mRNA decreased in LBP group (P<0.05, VS DM group) (Table 4, Figure 6).
Table 4: The effect of LBP on the VEGFmRNA of DM rats retina.

| Grouping | Gray scale (target gene/β-actin) | P*   | P#   |
|----------|----------------------------------|------|------|
| Control group | 0.085±0.011                        |      |      |
| DM group  | 0.438±0.044                       | <0.05|      |
| LBP group | 0.159±0.056                       | <0.05| <0.05|

*Compare with control group. #Compare between DM group and LBP group.

Ultrastructure Observation by Transmission Electron Microscopy

The retinal ultrastructure of the rats in the control group was normal. In the retina of the DM group, the damage of retinal nerve tissue was present in all layer of retina. A large number of huge mitochondria was found in the inner nuclear layer. Crista was tumid, cracked and reduced. Part of the mitochondria was round and vacuole-like. The nuclear membrane was non-continuous, with an increased number of unevenly distributed heterochromatin. Chromatin was concentrated below the nuclear membrane at different block mass sizes. The perinuclear space was enlarged. The cell nucleus of the partial bipolar cells was shrunk and cracked. Mitochondria and synaptic vesicle were visibly reduced in the neuron axon. The Müller cell volume increased with numerous loose endochylema and vacuoles. Numerous abnormal mitochondria were also found. The perinuclear space was enlarged. The membranous disc of the outer segment was disorderly with an unclear layered structure. The mitochondria in the inner segment ellipsoid was reduced and crista was vacuole-like. The microvilli of Müller cells decreased and exhibited disorderly arrangement. A small number of microvascular pericytes showed heterochromatin condensation and margination, as well as a non-continuous nuclear membrane. No clear change was found in the vascular endothelium cell and basilar membrane. In the LBP treatment group, no obvious abnormality was found in the retinal ganglionic cells and receptor cells. In the bipolar and Müller cells, the partial mitochondria crista became shorter and fewer. No abnormal change was found in the cell nucleus. The inner nuclear layer cells were regularly and tightly arranged. (Figure 7).

Figure 7.

Discussion

Diabetic retinopathy is one of the most common eye complications, considered a long-term pathological change in blood capillary level. Recent clinical and animal studies have shown that the pathological changes in retinal nerve tissues occur in the early stage of diabetic. Clinically, abnormal visual functions, including visual electrophysiology, color vision, contrast sensitivity and visual fields, show up before retinal microangioopathy in early stages. Research showed that pathological changes in retinal nerve tissue, such as degeneration and apoptosis of ganglionic...
cells, photoreceptor cells and neuroglial cells, occurred in the first week in diabetic animal models [4]. Barber believed that diabetic retinopathy is a degenerative disease of the nervous system manifesting in the eyes [5]. Our experimental results had shown that pathological changes in different degrees occur in almost all retinal nerve tissues of streptozotocin-induced diabetic rats in the 24th week, as represented by the change in ganglionic cells, photoreceptor cells, bipolar cells, and neurogial cells in terms of the number and morphology of mitochondria, increased nucleolus heterochromatin and reduced synaptic vesicle in the axon. Our study revealed obvious and extensive pathological changes in retinal nerve tissues, whereas no clear pathological changes were found in vascular endothelial cells and basilar membrane. Moreover, diabetic retinopathy possibly originated from the pathological changes in nerve tissues.

The pathogenesis of diabetic retinopathy currently remains unclear. Several pathogenic mechanisms have been posited, including the polyol pathway, generation of advanced glycation endproducts (AGES), protein línase C (PKC) and aminoheoxose pathway. Recent studies have shown that the generation of a large number of reactive oxygen species (ROS) induced by high glucose may be the common pathogenic factor of many kinds of chronic diabetic complications, including diabetic retinopathy [6]. A uniform pathogenic mechanism of diabetic complications was proposed in 2005 [7], the authors argued that the pathogenic mechanism of all diabetic complications is a unique pathway, the electron transfer chain of mitochondria induced by high glucose produces excessive superoxide. The pathogenic pathway determined by previous studies, including the polyol pathway, PKC activation, generation of AGE precursors in cells and activation of amidohexose pathway, resulted from the increase in mitochondrial ROS. ROS consists of a series of functional groups, such as O$_2^-$, H$_2$O$_2$, and NO. Many pathways can generate ROS while diabetic occurs. For example, the glycosylation and electron transfer chain of mitochondria can be activated. AGES, insulin and angiotensin II can induce ROS by activating NADPH oxidase combined with epicyte [8,9]. The study of diabetic renal disease indicated that the inhibition of NADPH oxidase can prevent AGE-induced renal damage in diabetic patients [10]. Under normal conditions, the antioxidant system in organisms can eliminate the oxidative substances produced by metabolism, and maintains the dynamic balance of the internal environment. For example, superoxide dismutase (SOD) can eliminate the oxidative substances produced by metabolism, and maintains the dynamic balance of the internal environment. In severe cases, the entire mitochondria exhibited vacuolar changes, indicating the importance of mitochondrial change in the pathological changes of mitochondrial ROS. ROS consists of a series of functional groups, such as O$_2^-$, H$_2$O$_2$, and NO. Many pathways can generate ROS while diabetic occurs. For example, the glycosylation and electron transfer chain of mitochondria can be activated. AGES, insulin and angiotensin II can induce ROS by activating NADPH oxidase combined with epicyte [8,9]. The study of diabetic renal disease indicated that the inhibition of NADPH oxidase can prevent AGE-induced renal damage in diabetic patients [10]. Under normal conditions, the antioxidant system in organisms can eliminate the oxidative substances produced by metabolism, and maintains the dynamic balance of the internal environment. For example, superoxide dismutase (SOD) can eliminate all kinds of oxygen radicals. When an organism produces more oxygen radicals beyond its antioxidant ability, ROS damages its tissues and organs. These attack the mitochondrial membrane, resulting in apoptosis in the mitochondrial pathways [11]. Excessive ROS also activates the action pathways of all known diabetic complications, including the polyol pathway, PKC, AGES precursors and amidohexose pathway, thereby inducing diabetic complications in multiple organs [12]. Malondialdehyde (MDA) is a lipid peroxidation product. It affects the electron transfer chain of mitochondria and key enzyme activity, aggravating the oxidative damage of cells. Therefore, MDA is also one of the key indicators of oxidative reaction in an organism. In our experiment, the SOD activity of diabetic rats induced by streptozocin decreased, whereas MDA increased. This finding confirms that oxidative stress has an important effect on the pathological changes in the retinal nerve tissues of diabetic rats. The ultrastructure of diabetic rats retina showed that the most prominent changes in retinal neurons and neurogial cells are the changes of mitochondria in the number, size and morphology. Several huge mitochondria can be found in the inner nuclear layer and inner molecular layer. The number of mitochondria in axons, photoreceptors and other positions decreased, with short, cracked and decreased cristae. Part of the cristae was beaker-like. In severe cases, the entire mitochondria exhibited vacuolar changes, indicating the importance of mitochondrial change in the pathological mechanism of diabetic retinopathy. Lycium barbarum polysaccharides (LBP) is extracted from the Chinese traditional herbs medicine, Ningxia medlar. Modern pharmacological studies found that LBP can remove redundant free radicals in the body [13], improve the activity of antioxidant enzymes [14,15] and increase the survival rate and promote the growth of rat retinal ganglion cells [16]. In our study, no significant difference in blood glucose level was found between the LBP group and the DM group before treatment, showing that the two groups were comparable. For blood glucose and body weight after treatment, no significant difference was also found between the two groups. These findings show that LBP could not reduce the blood glucose of animals. This result differs from that derived by Liu Ping [17]. The method they adopted was low dosage combined with high amounts of fat and sugar. The blood glucose of the modeled animal was 13.43 ± 1.36 mmol-L$^{-1}$. Our study adopted the high-dosage streptozocin simple modeling method. The blood glucose of the diabetic model rats were more than 20 mmol-L$^{-1}$. Therefore, our research can rule out the possibility of LBP indirectly curing diabetic retinopathy by reducing blood glucose. Significant pathological changes were not found in the gangliocyte and photoreceptor cell of the LBP group. A small amount of mild mitochondrial change could be observed in the endochylema of bipolar and Müller cells. This change manifested as the shortening and reduction in the number of cristae. SOD activity of the retina visibly increased and MDA decreased, indicating that LBP could alleviate the diabetic-induced pathological changes in retinal nerve tissues by reducing the oxidative damage caused by mitochondria pathway. The down-regulated VEGF expression showed that antioxidant therapy early can control the progression of retinal microangiopathy. However, the relation between the retinal nerve tissue pathological change and microangiopathy is unclear.

Thus, the early retinopathy of diabetic mainly occurs in the nerve cell and neuroglial cell by oxidative damage of mitochondria pathway. LBP can reduce the oxidative damage, alleviate the pathological changes in mitochondria, prevent the apoptosis of...
nerve cells and block the progression of diseases to the vascular tissue. LBP can be used in the early prevention and cure of diabetic retinal nerve pathological changes.

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