Adiponectin improves diabetic nephropathy by inhibiting necrotic apoptosis

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Abstract

Introduction: This study aimed to investigate the effect of adiponectin (Apn) on necrotic apoptosis (Nec) in vitro and in vivo to clarify the possible role of Apn in the pathogenesis of diabetic nephropathy (DN).

Material and methods: Rat glomerular endothelial (RGE) cells were treated with high glucose (HG, 30 mmol/l) for 24 h and the effects of Apn on cell viability, RIP1 and RIP3 expression and p-p38MAPK activation were assayed by CCK-8, immunofluorescence and western blot. Then a streptozotocin (STZ)-induced DN rat model was established. The body weight, left kidney weight, left kidney weight/body weight (KW/BW), creatinine clearance rate (Ccr), 24 h urine protein and blood glucose were recorded. The expression of RIP1, RIP3 and p-p38MAPK in renal tissues was examined by immunohistochemistry and western blot.

Results: Treatment of RGE cells with HG induced significant cytotoxicity and increased expression levels of RIP1, RIP3 and p-p38MAPK, which were abrogated by Apn in a concentration-dependent manner. In vivo, compared with the control group, the Ccr, 24 h urine protein and blood glucose level of the rats in the model group were significantly increased, effects which were abrogated by Apn intervention. Moreover, the expression levels of RIP1, RIP3 and p-p38MAPK were also significantly increased in the model group, effects which were canceled by Apn intervention.

Conclusions: Apn can alleviate the inflammatory response and damage of DN by inhibiting Nec via p-p38MAPK signaling.

Key words: rat glomerular endothelial cells, diabetic nephropathy, necroptosis, p38MAPK.

Introduction

Diabetic nephropathy (DN) is one of the common and serious microvascular complications of diabetes mellitus (DM). It is clinically characterized by proteinuria, edema and hypertension, and it further develops into azotemia and renal failure. 30–40% of the patients with a 15-year course of diabetes have DN, which is the leading cause of death in diabetic patients. The occurrence and development of DN is related to a variety of factors, including genetic susceptibility, glomerular sclerosis and hemodynamic changes, urinary albumin excretion, glucose toxicity [1–3] and so on.
Adiponectin (Apn) was first isolated and cloned in the mouse 3T3-L1 adipocyte cell line in 1995. It is named as adipocyte complement-associate protein 30 and is a specific protein secreted by adipocytes with a molecular mass of about 30 kD [4]. In recent years, studies have found that Apn has a certain correlation with obesity, type 2 DM, insulin resistance, atherosclerosis and diabetic vascular disease. It has been confirmed by animal experiments that insulin resistance is improved and blood glucose levels are reduced after injection of recombinant Apn into obese mice, suggesting that Apn has an important protective effect on diabetic damage [7].

In 2005, Degterev et al. reported a new way of cell death: Necroptosis (Nec) [8] is programmed cell death, which has morphological features similar to necrosis, while the cell death pattern is non-caspase-dependent, that is, the combination of death receptor and ligand can trigger necrotic apoptosis under the condition of caspase inhibition. Tumor necrosis factor (TNF)-α combining with TNFR1 on the plasma membrane causes the TNF receptor-associated death domain (TRADD) to send receptor interacting protein 1 (RIP1) signals to recruit RIP3, thereby forming a necrosome [9].

Inflammation is the main cause of Nec and increases the incidence rate of cardiovascular events in diabetic patients and the death rate of patients with renal failure [10]. Signal transduction pathways of mitogen-activated protein kinases (MAPKs) are the most critical pathways in the inflammatory transduction network [11]. This study aimed to investigate the effect of Apn on necrotic apoptosis of rat glomerular endothelial cells under high glucose (HG) and renal tissue in diabetic animal models, to clarify the possible role of Apn in the pathogenesis of DN and its signaling, and to provide a new theoretical basis for the prevention and treatment of DN.

**Material and methods**

**Cell culture**

Rat glomerular endothelial (RGE) cells were purchased from ATCC, USA, incubated in DMEM medium containing 10% fetal bovine serum (FBS) and maintained in 5% CO₂ at 37°C. When the cell density reached 90%, the cells were digested and counted; 10⁵ cells were inoculated into new culture flasks, cultured to 80% fusion in serum-free medium for 24 h. Subsequently, they were divided into a control group (5 mmol/l glucose), HG group (30 mmol/l glucose), Apnl group (30 mmol/l glucose + 1 μg/ml Apn), ApnM group (30 mmol/l glucose + 5 μg/ml Apn), ApnH group (30 mmol/l glucose + 25 μg/ml Apn), and mannitol group (5 mmol/l glucose + 25 mmol/l mannitol), and continued to culture for 24 h.

Cell viability [12] was determined by CCK-8

RGE cells were inoculated in a 96-well culture plate. When the cells were grown to about 80% fusion, the above grouping was performed. They were incubated for different periods, then the supernatant was discarded, and washed 3 times with PBS; 90 μl of DMEM and 10 μl of CCK-8 were added to each well, incubated for 2.5 h in a 37°C incubator, and absorbance density (OD 490 nm) of the wells was recorded with a microplate reader. This experiment was repeated 5 times.

**Animal model establishment**

This study was approved by the Ethics Committee of Nanchang University and conducted in accordance with the Declaration of Helsinki and the Guide for the Care and Use of Laboratory Animals. Thirty male, standard, clean, healthy Sprague-Dawley (SD) rats were provided by the Experimental Animal Center.

Diabetes model preparation: STZ was dissolved in citrate buffer (pH = 4.5) in the weight ratio of 60 mg/kg; under fasting conditions, intraperitoneal injection was performed; the normal control group was only injected with an equal amount of citrate buffer. The blood glucose level was measured at 72 h, and ≥ 16.7 mmol/l indicated that the diabetes animal model was successfully established. Among the 30 rats, in 28 the blood sugar reached the standard, with a success rate of 93.3%.

DN model preparation: 24 h urine protein was measured at 4 weeks, and ≥ 30 mg/day indicated that the DN models were successfully established. Among the 28 rats, 20 reached the standard, with a success rate of 71.4%. The successful DN animal model was randomly divided into a model group (n = 10) and a DN/Apn group (n = 10), and 2 mg/kg/day Apn was injected into the tail vein of the rats in the DN/Apn group for 6 weeks. Eight SD rats with matched age and body weight were finally selected as the normal control group. At the end of the study, the animals were injected with isoflurane overdose and subjected to cervical dislocation to sacrifice the animals.

**Collection and testing of specimens**

In 6 weeks after the Apn administration, the rats were placed in a metabolic cage, and urine was collected in 24 h. After filtration, the rats’ urine was stored at -70°C for urine protein quantification. Serum creatinine and urine creatinine were measured using an automatic biochemical analyzer, and the creatinine clearance rate (Ccr)
was calculated using the following equation: Ccr = 
urine creatinine × urine volume/serum creatinine × 1440/body weight. Then, the rats were weighed, anesthetized by intraperitoneal injection with sodium pentobarbital at a weight ratio of 45 mg/kg, and the left kidneys of the rats were quickly weighed. After the right kidney was lavaged with normal saline, some right kidney tissues were taken and fixed with 10% neutral formaldehyde, and embedded in paraffin for immunohistochemistry and hematoxylin-eosin (HE) staining. Some kidney tissues were taken separately for western blot analysis.

Renal pathologic examination

Renal tissue was fixed with 10% neutral formaldehyde, embedded in paraffin, sliced in sections 4 μm thick, and stained with HE in order to observe the degree of mesangial matrix hyperplasia as well as glomerular, renal tubule and renal interstitial lesions. Glomerular hypertrophy was confirmed by measuring the mean glomerular diameter of 20 glomeruli. Renal tissue sections were routinely dewaxed and dehydrated, incubated in 3% hydrogen peroxide for 10 min at room temperature to eliminate endogenous peroxidase, and blocked with serum for 10 min, then rabbit anti-rat RIP1 and PIP3 antibody were added (1 : 200, overnight at 4°C), and washed with phosphate buffer saline (PBS) twice, 3 min each time. On the next day, biotin-labeled secondary antibody was added to each section, incubated at 37°C for 30 min, and then washed twice with PBS, 3 min each time. Next, HRP-labeled streptavidin was added to each section, incubated at 37°C for 30 min, and then washed twice with PBS, 3 min each time. After they were developed with diamobenzidine (DAB) for 5–20 min at room temperature, they were slightly counterstained with hematoxylin-eosin.

Western blotting

Cells or kidney tissues were lysed on ice for 30 min with cell lysis solution (containing protease inhibitor cocktail). They were centrifuged at 12 000 × g for 15 min at 4°C, and then the supernatant was taken. Protein quantification was performed by the bicinchoninic acid (BCA) method. A 50 μg sample was loaded in each group, and subjected to sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) using 5% concentrated gel and 8% or 10% separating gel to separate the proteins (RIP1, PIP3, p-p38MAPK and t-p38MAPK). These proteins were transferred to a polyvinylidene fluoride (PVDF) membrane at 220 mA electric current for 1 h; then they were blocked for 1 h at room temperature; after that they were washed 3 times with tris buffered saline Tween (TBST), and the corresponding primary antibodies were incubated overnight at 4°C. The next day, they were incubated with corresponding secondary antibody for 1 h at room temperature; then, they were incubated with an enhanced chemiluminescence solution for 2 min, and quantified using Alpha-Ease software version 2200 [13].

Statistical analysis

Statistical analysis was performed using SPSS 19 statistical software. The measurement data were expressed by mean ± SD, and the comparison between groups was made by analysis of variance (ANOVA). P < 0.05 indicated that the difference was statistically significant.

Results

Apn protected HG-induced Nec

Figure 1 A shows that treatment of RGE cells with HG (30 mmol/l) for 24 h can induce significant cytotoxicity and thus reduce cell viability; compared with the normal control group, the difference is statistically significant. The cell viability was increased by co-treatment of the cells with 1, 5 or 25 μg/ml Apn and HG for 24 h, which was statistically significant (p < 0.05); the inhibitory effect of 25 μg/ml Apn on cytotoxicity was most pronounced.

We tested the expression of RIP1 and RIP3 in cells by further application of immunofluorescence and western blot. The expression levels of RIP1 and RIP3 in cells treated with HG for 24 h were significantly increased. Co-treatment of cells with 1, 5 or 25 μg/ml Apn and HG for 24 h could reduce the expression of RIP1 and RIP3. The differences were statistically significant (Figures 1 B, C).

The expression of p-p38MAPK was also significantly increased by treating the cells with HG for 24 h; compared with the normal control group, the difference was statistically significant (p < 0.01). The expression of p-p38MAPK was significantly decreased by co-treatment of cells with 1, 5 or 25 μg/ml Apn and HG for 24 h (p < 0.01, Figure 1 C).

Apn improved DN

After 10 weeks, STZ rats clearly showed symptoms of polydipsia, polyuria and polyphagia, and their body weights were significantly reduced. In addition to the death of 3 rats in the model group, the animals in other groups were in a good condition. The body weight, left kidney weight, left kidney weight/body weight (KW/BW, g/kg), Ccr, urine protein and blood glucose for each group of rats are shown in Figure 2. Compared with the rats in the control group, the body weights of the
Figure 1. Apn protects HG-induced Nec. A – Cell viability was assayed by CCK-8 assay. Treatment of RGE cells with HG (30 mmol/l) for 24 h can cause significant cytotoxicity. Cell viability was promoted by co-treatment of the cells with 1, 5 or 25 μg/ml Apn. B – The expression of RIP1 and RIP3 in cells were examined by immunofluorescence. C – The expression of RIP1, RIP3 and p-p38MAPK were assayed by western blot. The expression levels of RIP1, RIP3 and p-p38MAPK in HG-treated cells were significantly increased. Co-treatment of cells with 1, 5 or 25 μg/ml Apn could reduce the expression of RIP1, RIP3 and p-p38MAPK.

*p < 0.05 and **p < 0.01 vs. Control; *p < 0.05 and **p < 0.01 vs. HG.
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**Figure 2.** Apn improves DN animal models (n = 8). Compared with the rats in the control group, the body weights of the rats in the model group were significantly reduced from the 4th week, and were increased from the 6th week after Apn intervention. The left kidney weight and KW/BW of the rats in the model group were significantly increased, and KW/BW significantly decreased after Apn intervention. Ccr, 24 h urine protein and blood glucose from the rats in the model group increased significantly from the 4th week, and significantly decreased from the 4th week after Apn intervention.

*p < 0.05 and **p < 0.01 vs. Control; *p < 0.05 and **p < 0.01 vs. Model.
Figure 3. Apn reduced the expression of RIP1, RIP3 and p-p38MAPK (n = 8). A – HE staining. The glomerular diameter was significantly increased in the model group, and was decreased after Apn intervention. B – Immunohistochemistry of RIP1 and PIP3 in renal tissues. C – Western blot assay. The expression levels of RIP1, PIP3 and p-p38MAPK for the rats in the model group were significantly increased, while they were significantly decreased after Apn intervention.

**p < 0.01 vs. Control; *p < 0.05 and ##p < 0.01 vs. Model.
In its pathogenesis [14]. It has been confirmed that proteinuria in rats increased significantly at 4 weeks, and increases progressively with time. In animal models, we found that Apn significantly improved DN including increasing the body weight and KW/BW and decreasing Ccr, 24 h urine protein and blood glucose. Also, Apn reduced the expression of RIP1, RIP3 and p-p38MAPK and decreased the expression of RIP1 and RIP3.

In conclusion, Apn can alleviate the inflammatory response and damage of DN by inhibiting the p38MAPK pathway.

Conflict of interest
The authors declare no conflict of interest.

References


