Beneficial Effects of IncRNA-UC.360+ shRNA on Diabetic Cardiac Sympathetic Damage via NLRP3 Inflammasome-Induced Pyroptosis in Stellate Ganglion

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ABSTRACT: Hyperglycemia is one of the common symptoms of diabetes, and it produces excessive reactive oxygen species (ROS). This study investigated whether the long noncoding RNA (lncRNA) UC.360+ is involved in diabetic cardiac autonomic neuropathy (DCAN) mediated by NLRP3 inflammasome-induced pyroptosis in the stellate ganglion (SG). Using a rat type 2 diabetes model, we found that IncRNA UC.360+ short hairpin RNA (shRNA) ameliorated the dyslipidaemia of type 2 diabetic rats and reduced serum adrenaline and ROS production in SG under hyperglycemia. In addition, UC.360+ shRNA also reduced the expression of nuclear factor kappa-B (NF-κB), NLRP3, ASC, caspase-1, interleukin-1β (IL-1β), and IL-18 in the SG of diabetic rats and inhibited the phosphorylation of p38 mitogen-activated protein kinase (p38 MAPK). Therefore, IncRNA-UC.360+ shRNA may modulate the NLRP3 inflammasome/inflammatory pathway in the SG, which in turn alleviates diabetic heart sympathetic nerve damage.

INTRODUCTION

Diabetes is characterized by hyperglycemia and is caused by the implication of both genetic and environmental risk factors. The number and prevalence of diabetes are continuously rising. Diabetic patients in China account for more than 25% of the total number of adults with diabetes in the world, which is the country with the highest number and prevalence of diabetes. In terms of the type of diabetes, the diabetes population in China is mainly the type 2, accounting for more than 90%. As the progress of disease course, it will cause complications in various tissues and organs, and diabetic cardiac autonomic neuropathy (DCAN) is one of them. DCAN in diabetic patients involves only the vagus nerve injury at the early stage, but as the disease advances, both sympathetic and parasympathetic nerves as well as myelinated and unmyelinated fibers are damaged, causing various clinical symptoms. Studies have shown that the mortality rate of DCAN patients within 5 years is more than five times of that seen in non-DCAN patients. DCAN not only reduces the living quality of diabetic patients but also has seriously negative impact on the social economy. ROS at a proper level exhibit beneficial effects by regulating intracellular signal transduction and homeostasis, while ROS at a high concentration have a damaging effect on proteins, lipids, and DNA. The body’s antioxidant defense system maintains the dynamic balance of ROS, and disruption of this balance either by increased ROS formation or decreased antioxidant activity in biological systems may lead to oxidative stress. Increased ROS levels can promote the release of the proinflammatory transcription factor NF-κB, which mediates the occurrence of a series of diseases. ROS participate in pathological changes of diabetes implicated with cellular damage. Thus, targeting oxidative stress may be a strategy to prevent and treat diabetic complications with cellular damage.

In the human transcriptome, those RNAs having no apparent protein-coding role regions are collectively referred to as noncoding RNAs (ncRNAs), and the ncRNAs longer than 200 nucleotides are called long noncoding RNAs (lncRNAs). In recent years, with the progress of global transcriptome analysis, an increasing number of differential expression studies have demonstrated that lncRNAs can regulate many important biological processes, which may be implicated in the occurrence of human diseases. Among various organ systems, lncRNAs are preferentially expressed in...
the nervous system, and consequently, the aberrant expression of lncRNAs is involved in the most devastating neurological diseases. Therefore, lncRNAs may be key players in the normal development, physiological function, and pathological changes in the nervous system. To explore the possible role of lncRNAs in nerve damage, the SOLiD Whole Transcriptome Analysis Kit was used in the early stage of this project to complete Sprague Dawley rat multi-tissue transcriptome high-throughput sequencing. Further analysis showed that many noncoding TUs in peripheral ganglion tissue have poly (A) tail structures, similar to lncRNAs. After SOLiD high-throughput sequencing, this project also identified some lncRNAs through bioinformatics predictions and molecular biology verification. A few studies showed that abnormal expression of lncRNAs is related to neurological diseases, but the mechanism of action is still unclear.

This project targeted an lncRNA (lncRNA-UC.360+ as searched online) that exists in the neck sympathetic ganglia of a rat model of type 2 diabetes for experiments. Pyroptosis can occur in vivo and in vitro. The typical features of cell pyroptosis involve these events: the plasma membrane loses its integrity, cell permeability increases, the cell membrane ruptures to form numerous small pores with a diameter of 1–2 nm, and intracellular substances are released outside the cell. Among the several types of inflammasomes that have been discovered, NLRP3 is linked to type 2 diabetes. The inflammasome adapter ASC self-oligomerizes to form a macromolecular complex, activates caspase-1, and then cleaves IL-1β and IL-18 precursors into active forms that can induce the synthesis of other pro-inflammatory factors and amplify local and systemic inflammatory responses.

**RESULTS**

Effect of UC.360+ shRNA on Fasting Blood Glucose (FPG) and Blood Lipids. Before UC.360+ shRNA injection (at the sixth week of the experiment), the FPG level in the diabetes mellitus (DM) group was higher than that in the control group (Figure 1A), suggesting that our model of diabetic rats was successfully established. At the eighth week, the FPG values in the DM group was still higher than that in the control group (p < 0.001, Figure 1B). After UC.360+ shRNA treatment for 2 weeks, the FPG values in the DM rats was significantly reduced (p < 0.001, Figure 1B). The DM group did not significantly differ from the DM + NC shRNA group (p > 0.05, Figure 1B). Thus, UC.360+ shRNA treatment could reduce the blood glucose level of type 2 diabetic rats.

Blood lipids mainly include T-CHO, LDL-C, TG, and HDL-C. The results showed that the levels of T-CHO, TG, and LDL-C were higher in the DM group than in the control group (p < 0.001, Figure 1C–F). After treatment with UC.360+ shRNA, the values were significantly reduced (p < 0.001, Figure 1C–F). The change in the value of HDL-C was contrary to the aforementioned three lipids under the same conditions (Figure 1F). The DM group did not significantly differ from the DM + NC shRNA group (p > 0.05, Figure 1C–F). Our observations suggested that UC.360+ shRNA treatment could improve dyslipidaemia in type 2 diabetic rats.

![Image](https://example.com/image.png)

**Figure 1.** Effect of UC.360+ shRNA on blood glucose and lipids in DM rats. ELISA was used to detect FPG (A, B) and lipids (C–F) in each group. ***p < 0.001 vs control group, ****p < 0.001 vs DM group; n = 3 per group.

| Table 1. Concentration of Serum Epinephrine (EPI, pg/mL) and ROS in SG Cells of Rats a |
|-----------------|-----------------|-----------------|-----------------|
|                 | control         | DM              | DM + uc.360+ shRNA | DM + NC shRNA   |
| EPI content     | 62.13 ± 1.29    | 168.77 ± 7.94*** | 116.17 ± 16.29****| 170.20 ± 12.64***|
| ROS content     | 62.5 ± 6.98     | 229.67 ± 10.65## | 82.83 ± 5.88***   | 231.83 ± 12.64**|

a###p < 0.001 vs control, ***p < 0.001 vs DM; n = 15 per group.
Effect of UC.360+ shRNA on Serum Epinephrine and ROS Content. ELISA was used to detect the serum content of epinephrine (Table 1). The epinephrine content in the DM group was higher than that in the control group ($p < 0.001$). The content in the DM+ UC.360+ shRNA group was lower than that in the DM group ($p < 0.001$). The DM group did not significantly differ from the DM + NC shRNA group ($p > 0.05$).

ELISA was also used to detect the content of ROS in SG (Table 1). Our results showed that the ROS content in the DM group was higher than that in the control group ($p < 0.001$). The content was lower in the DM+ UC.360+ shRNA group than in the DM group ($p < 0.001$). The DM group did not significantly differ from the DM + NC shRNA group ($p > 0.05$).

Effect of UC.360+ shRNA on Activation of the p38 MAPK Pathway in SG. The p38 MAPK-mediated pathway is mainly involved in stress responses. Western blotting detection found that the p38 MAPK level in the control group did not significantly differ from the DM group after normalizing by the $\beta$-actin value (Figure 2A). In contrast, the value of phospho-p38 MAPK in the DM group was higher than that in the control group ($p < 0.05$) after normalizing to the P38 value. After UC.360+ shRNA injection, the value of phospho-p38 was reduced ($p < 0.05$). The DM group did not significantly differ from the DM + NC shRNA group ($p > 0.05$).

Phosphorylation of p38 MAPK may lead to the activation of the proinflammatory transcription factor NF-κB. Western blotting detection was performed (Figure 2B), and the value of NF-κB P65 protein was normalized by $\beta$-actin. The value of NF-κB P65 protein in the DM group was higher than that in the control group ($p < 0.01$), whereas the value was lower in the DM+ UC.360+ shRNA group than in the DM group ($p < 0.001$). The DM group did not significantly differ from the DM + NC shRNA group ($p > 0.05$).

Effect of UC.360+ shRNA on Expression of NLRP3, ASC, and Caspase-1 in SG. The NLRP3 inflammatory body

Figure 2. UC.360+ shRNA reduced the phosphorylation of p38 MAPK and content of NF-κB in the SG of DM rats. Western blotting was used to determine the protein levels of p38 MARK, p-p38 MARK (A), and NF-κB (B) in the SG. *$p < 0.05$, **$p < 0.001$ vs control group, *$p < 0.05$, ***$p < 0.001$ vs DM group; $n = 3$ per group.

Figure 3. UC.360+ shRNA reduced the expression of NLRP3, ASC, and caspase-1 in the SG of DM rats. The expression of NLRP3, ASC, and caspase-1 mRNA (A–C) and protein (D–F) was detected by real-time quantitative PCR and Western blotting, respectively. ***$p < 0.001$ vs control group, **$p < 0.01$, ***$p < 0.001$ vs DM group; $n = 3$ per group.
is a key factor inducing pyroptosis. The mRNA and protein values were standardized by β-actin. The mRNA (Figure 3A–C) and protein (Figure 3D–F) levels of NLRP3, ASC, and caspase-1 in the DM group were higher than those in the control group \( p < 0.001 \). The values were lower in the DM+UC.360+ shRNA group than in the DM group \( (p < 0.01 \text{ or } p < 0.001) \). The DM group did not significantly differ from the DM + NC shRNA group \( (p > 0.05) \).

Immunohistochemistry was performed to detect caspase-1 immune reactivity in rat SG (Figure 4). The extent of caspase-1 immune reactivity in the DM group was higher than that in the control group \( (p < 0.001) \). This could be significantly reduced after UC.360+ shRNA treatment \( (p < 0.001) \). The DM group did not significantly differ from the DM + NC shRNA group \( (p > 0.05) \).

**DISCUSSION**

In this study, the detection of fasting blood glucose was used to confirm whether the rat diabetic model was successfully established. The detection of blood lipids was to validate whether IncRNA-UC.360+ shRNA could improve dyslipidemia induced by type 2 diabetes. In diabetes, the state of high glucose can prompt cells to produce excess ROS, which in turn regulates the MAPK signaling pathway and the activation of various cytokines and transcription factors. The detection of ROS in the study is not only a further confirmation of the success of the model establishment but also serves as a guiding flag in the subsequent activation of the MAPK signaling pathway. The activation of p38MAPK regulates the expression...
of NF-κB, and NF-κB further regulates the activities of a series of components of the inflammasome NLRP3. Therefore, the experiments assessing p38 MAPK, NF-κB, NLRP3, ASC, caspase-1, IL-1β, and IL-18 by real-time PCR, Western blotting, and immunohistochemistry were followed. These experiments would reveal how lnRNA-UC.360+ shRNA treatment exhibits its potential beneficial action on counteracting the over-production of intracellular ROS and activation of the ROS-NLRP3-IL-1β signaling pathway in diabetes, thereby protecting cervical sympathetic ganglion nerve cells from pyroptosis damage.

Hyperglycemia is a typical feature of diabetes, and T2DM initially manifests as insulin resistance, which gradually progresses to β-cell damage and eventually leads to hyperglycemia. Epidemiological studies have shown that approximately 50% of type 2 diabetic patients suffer from dyslipidemia. There are studies showing that dyslipidemia also promotes the risk of developing T2DM, thus, dyslipidemia and T2DM are viewed to interact and cause each other. Therefore, in the study of type 2 diabetes, attention should be given to the phenomenon of dyslipidemia. Research has shown that LDL-C is involved in the activation of the NF-κB signaling pathway and elicits the production of ROS, and T2DM are viewed to interact and cause each other. Dyslipidemia. There are studies showing that dyslipidemia approximately 50% of type 2 diabetic patients suffer from glycemia. Epidemiological studies have shown that approximately 50% of type 2 diabetic patients suffer from glycemia. Epidemiological studies have shown that approximately 50% of type 2 diabetic patients suffer from glycemia. Epidemiological studies have shown that approximately 50% of type 2 diabetic patients suffer from glycemia. Epidemiological studies have shown that approximately 50% of type 2 diabetic patients suffer from glycemia.

The p38MAPK pathway primarily mediates the stress response and can be activated by many stimuli including ROS.22 Activation of p38 MAPK can regulate the expression of a variety of transcription factors, including NF-κB.23 Studies have shown that MAPK is a signal transduction factor in high glucose caused various complications of diabetes. p38 MAPK is involved in type 2 diabetic autonomic neuropathy.4,22 This study found that the content of p38 MAPK in SG was not significantly altered, but the content of its active phosphor-p38 MAPK in SG of DM rats was upregulated, suggesting the activation of redox sensitive signaling pathways. In addition, lnRNA-UC.360+ shRNA treatment decreased the content of phospho-p38 MAPK in the SG of DM rats, suggesting that lnRNA-UC.360+ shRNA could effectively inhibit the activation of the p38 MAPK signaling pathway. Moreover, the level of NF-κB protein in the SG of DM rats was increased, and it was relieved by lnRNA-UC.360+ shRNA treatment, indicating that the activation of p38 MAPK may regulate NF-κB.

Pyroptosis is accompanied by the activation of caspase-1/4/5/11,24 and the caspase-1-induced pyroptosis is called the classical cascade.25 The NLRP3 inflammasome is related to type 2 diabetes.26 Under physiological conditions, the level of NLRP3 in the body is extremely low, NLRP3 mRNA needs to be transcribed first. At this time, NF-κB activation is required, and then translation and posttranslational modifications ensue.20 Studies have shown that increasing the content of ROS in the cytoplasm can activate NLRP3 inflammasomes.27 The high glucose in diabetes promotes the production of a large amount of ROS in the cell, which in turn can activate the NLRP3 inflammasome. NLRP3 inflammasome is a key factor in inducing pyroptosis. Cell pyroptosis is involved in diabetic nephropathy16,32,33 and diabetic cardiomyopathy,34,35 but the process of diabetic autonomic neuropathy induced by type 2 diabetes remains to be studied in depth. Activation of the inflammasome leads to the processing and activation of caspase-1, turning pro-caspase-1 into active caspase-1. Afterward, pro-IL-1β and pro-IL-18 are cleaved into their mature forms by caspase-1. In this study, Western blotting detected upregulated protein expression of NLRP3, ASC, caspase-1, IL-1β, and IL-18 in the SG of type 2 diabetic rats. These upregulated changes indicate inflammasome activation. LnRNA-UC.360+ shRNA treatment effectively inhibited the increase in these signaling molecules. These findings suggest that ROS may regulate the expression of NF-κB by activating the p38 MAPK signaling pathway and in turn activate the NLRP3 inflammasome to elicit pyroptosis, whereas lnRNA-UC.360+ shRNA reduced inflammasome activation and alleviated diabetic injury-mediated pyroptosis.

In conclusion, hyperglycemia in diabetes may promote the over-production of ROS in cells, and such oxidative stress can activate the p38 MAPK-mediated signaling pathway, which stimulates NF-κB and further regulates the activities of a series of components of the NLRP3 inflammasome. LnRNA-UC.360+ shRNA treatment effectively downregulated the production of intracellular ROS and inhibited the activation of the ROS-NLRP3-IL-1β signaling pathway, thereby reducing the pyroptosis of cervical sympathetic ganglion nerve cells and improving diabetic cardiac sympathetic nerve damage.

**METHODS**

**Animals.** Male Sprague–Dawley rats weighing 180–220 g were purchased from Changsha Tianqin Experimental Animal
Center. The animal operation protocol was reviewed and approved by the Ethics Committee of Medical School of Nanchang University. The rats were placed in clean cages, and the room was kept at 22 ± 2 °C and 50% humidity. After adaptation, the rats were randomly divided into a control group and a DM group. After 4 weeks of feeding with a high-sugar and high-fat diet, rats for type 2 diabetic model were injected intraperitoneally (i.p.) with streptozotocin (STZ) (30 mg/kg). The establishment of a rat diabetic model and drug treatment were performed as previously described. In short, in this experiment, rats were fed a high-sugar and high-fat diet to induce insulin resistance. Later, a small dose of STZ was injected i.p. to impair the islet function and reduce insulin secretion. Without sufficient insulin levels, the body cannot maintain normal blood sugar levels, thus establishing a rat model of type 2 diabetes.

**Blood Glucose.** A Roche blood glucose meter was used to measure blood glucose after 12 h of fasting (fasting at 21:00 the night before, blood glucose measurement at 9:00 the next morning, free drinking). After the tail of the rat was sterilized and approximately 3 mm of tail tip was cut off, the blood was dropped onto the blood glucose test paper that was inserted into the meter to read the value of blood glucose. After waiting for 2 s, the blood glucose value was read on the blood glucose meter and recorded.

**Blood Lipids.** Fresh blood was collected and kept at 25 °C for 30 min. The samples were centrifuged at 2000 × g for 15 min at 4 °C, and the upper layer of pale-yellow clear liquid was serum. The serum was collected in disposable endotoxin-free test tubes and kept on ice for testing. If serum could not be tested on the same day, it was stored at −80 °C until use. The contents of T-CHO, TG, LDL-C, and HDL-C in the serum of rats were measured by assay kits (Elabscience Biotechnology, China).

**Epinephrine (EPI).** Blood collection was conducted as described above. The serum adrenaline concentration of rats was measured according to the kit instructions (Elabscience Biotechnology, China). This kit uses a competitive ELISA method. The EPI antigen is coated on the ELISA plate. During the measurement, the EPI in the sample or standard competes with the coated EPI to bind to the site on the biotin-labeled anti-EPI monoclonal antibody. Horseradish peroxidase-labeled avidin was added, and after the reaction, the EPI concentration was measured.

**ROS.** Fresh SG tissue was placed in precooled PBS to remove other impurities around the tissue. After cutting the SG into 1 mm³ pieces, a certain amount of pancreatin was added for digestion (37 °C, 30 min). The digestion was stopped with precooled PBS, and the samples were passed through a 300-mesh nylon mesh to remove tissue blocks and then centrifuged (1500 × g, 10 min). The supernatant was removed, and a single-cell suspension was obtained. The total number of cells used for fluorescence detection was ensured to be greater than 10⁶. The wavelength settings were as follows: excitation light of 500 nm and emission wavelength of 525 nm, the value obtained was the fluorescence value needed.

**Real-Time PCR.** After the diabetic model was established, the SG tissue was separated and washed in PBS to remove residual blood. The extraction kit was used for RNA extraction (Beijing TransGen Biotech Co. Beijing, China). After the sample was lysed by TRIzol and layered with chloroform, isopropanol and absolute ethanol were added successively for precipitation to obtain high-purity total RNA. Then, the instructions of the cDNA synthesis kit were followed to reverse transcribe the required cDNA (Thermo Fisher Scientific, USA). The total of 20 μL cDNA reverse transcription reaction system was set up according to the steps as follows: 4 μL of 5 × buffer, 2 μL of dNTPs, 1 μL of OligoDT, 1 μL of MMLV, 1 μL of RNasin, and 11 μL of RNA sample, they were mixed well in a 37 °C water bath for 1 h.

PCR system was configured according to the Promega instructions. Each primer and each cDNA template were added to three replicate wells, and each well was 20 μL, which included the following reaction components: 7 μL of riboyme-free water, 0.5 μL of both upstream and downstream primers, 10 μL of 2 × Master Mix, and 2 μL of cDNA. The primer sequences are shown in Table 2.

**Western Blotting.** The previously separated SG tissue was removed from −80 °C and ground with a homogenizer after 300 μL of lysate was added (the homogenizer was soaked in acid overnight, autoclaved, and used after drying, the whole procedure was performed on ice). The milled homogenate was transferred to a precooled EP tube and centrifuged. The supernatant was collected. Loading buffer was added according to the proportion, mixed and sealed, and boiled in boiling water until denatured. After cooling, the samples were transferred to −20 °C for storage. SDS–PAGE of the corresponding specifications was performed as required. After BCA quantification of protein content, 15 μL of the protein sample was added to each well. Gel electrophoresis was performed at 20 mA/gel for approximately 2 h. Then, the membrane was transferred to a constant current of 300 mA in an ice bath, and the transfer time was approximately 90 min. The blots on the membrane were incubated with primary and secondary antibodies and placed in a gel imager for development. The following primary antibodies (dilution ratio of 1:1000) were used: rabbit anti-p-p38 MAPK and rabbit anti-p38 MAPK (Cell Signaling Technology, Beverly, MA), rabbit anti-NF-κBp65 and rabbit anti-ASC (Affinity, Cincinnati, OH, USA), rabbit anti-Pro-Caspase-1 (Novus Biologicals, USA), rabbit anti-NLRP3, rabbit anti-Pro-ASC (Novus Biologicals, USA), rabbit anti-ASC, rabbit anti-Pro-IL-1β and rabbit anti-Pro-IL-18 (Abcam, UK), and mouse anti-β-actin (ZSGB-Bio, Beijing, China). Anti-mouse IgG and goat anti-rabbit IgG (ZSGB-Bio, Beijing, China) were used as the secondary antibodies with a dilution ratio of 1:2000. The optical density (OD) values were analyzed by ImageJ.

**Immunohistochemistry.** The isolated rat SG was fixed in 4% PFA, and then 10 μm frozen slices were prepared. The slices were deparaffinized and rehydrated after incubation with 3% H₂O₂ (10 min). The slices were blocked with 3% BSA for 90 min and incubated with rabbit anti-caspase 1 antibody (1200 overnight). Afterward, the slices were incubated with a secondary antibody for 2 h. The slides were mounted with neutral gum, and images were captured under a microscope.

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**Table 2. Primer Sequences**

| Gene         | Primer Sequences                  |
|--------------|-----------------------------------|
| β-actin      | forward: 5′- TAAAGACCTCTATGCAACACAGT -3′ |
|              | reverse: 5′- CACGTAGGAGGCGGCGACTCATC -3′ |
| NLRP3        | forward: 5′- CTGCATGGCTATGCTGTTG -3′ |
|              | reverse: 5′- CCGGCTTTAGCAAAATCCAG -3′ |
| ASC          | forward: 5′- CTGTGATGCGAATGTCGAC-3′ |
|              | reverse: 5′- GAAACAGGTCCTTGGTCCAG -3′ |
| caspase-1    | forward: 5′- GGACGTTCAGTGCCAGTCATC -3′ |
|              | reverse: 5′- GCCGACCTTCTTGGTCCAG -3′ |

**Western Blotting.** The previously separated SG tissue was removed from −80 °C and ground with a homogenizer after 300 μL of lysate was added (the homogenizer was soaked in acid overnight, autoclaved, and used after drying, the whole procedure was performed on ice). The milled homogenate was transferred to a precooled EP tube and centrifuged. The supernatant was collected. Loading buffer was added according to the proportion, mixed and sealed, and boiled in boiling water until denatured. After cooling, the samples were transferred to −20 °C for storage. SDS–PAGE of the corresponding specifications was performed as required. After BCA quantification of protein content, 15 μL of the protein sample was added to each well. Gel electrophoresis was performed at 20 mA/gel for approximately 2 h. Then, the membrane was transferred to a constant current of 300 mA in an ice bath, and the transfer time was approximately 90 min. The blots on the membrane were incubated with primary and secondary antibodies and placed in a gel imager for development. The following primary antibodies (dilution ratio of 1:1000) were used: rabbit anti-p-p38 MAPK and rabbit anti-p38 MAPK (Cell Signaling Technology, Beverly, MA), rabbit anti-NF-κBp65 and rabbit anti-ASC (Affinity, Cincinnati, OH, USA), rabbit anti-Pro-Caspase-1 (Novus Biologicals, USA), rabbit anti-NLRP3, rabbit anti-Pro-ASC (Novus Biologicals, USA), rabbit anti-ASC, rabbit anti-Pro-IL-1β and rabbit anti-Pro-IL-18 (Abcam, UK), and mouse anti-β-actin (ZSGB-Bio, Beijing, China). Anti-mouse IgG and goat anti-rabbit IgG (ZSGB-Bio, Beijing, China) were used as the secondary antibodies with a dilution ratio of 1:2000. The optical density (OD) values were analyzed by ImageJ.

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Statistical Analysis. SPSS 21 and Origin 9.0 were used for data analysis. Prism 5.0 was used for data graphing. One-way ANOVA was used for comparisons between two or more data points, and the LSD method was used for comparisons between groups. Values are presented as the mean ± SD. P < 0.05 was defined as significant.

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Author Contributions
All authors approved the final version of manuscript. S.D.L., L.R.S., and G.D.L. conceived and designed the research, and drafted the manuscript. L.R.S. performed the statistical analysis and interpreted the data. L.R.S., Q.X.H., L.L., R.N.Y., X.M.X., J.P.D., L.F.Z., G.L.L. and S.M.L. performed the experiments.

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Notes
The authors declare no competing financial interest.

ABBREVIATIONS
ROS reactive oxygen species; LncRNA long noncoding RNA; DCAN diabetic cardiac autonomic neuropathy; SG stellate ganglion; NLRP3 NOD-like receptor thermal protein domain associated protein 3; ASC apoptosis-associated Speck-like protein containing a CARD; NF-κB nuclear factor kappa-B; IL-1β interleukin-1β; IL-18 interleukin-18; p38 MAPK p38 mitogen-activated protein kinase; DM diabetes mellitus; ELISA enzyme-linked immunosorbent assay

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