

# SIRT3 is a downstream target of PPAR- $\alpha$ implicated in high glucose-induced cardiomyocyte injury in AC16 cells

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**Abstract.** Diabetic cardiomyopathy (DCM) is a worldwide public health concern that continues to display rapid growth trends. This study investigated the function of sirtuin 3 (SIRT3), a primary mitochondrial deacetylase with important roles in antioxidant defense and oxidative metabolism, during high glucose-induced cardiomyocyte (AC16 cell) injury. Peroxisome proliferator-activated receptor- $\alpha$  (PPAR- $\alpha$ ) is directly related to the occurrence of DCM. Hence, we further examined the relationship between SIRT3 and PPAR- $\alpha$ . AC16 cells were treated with various concentrations of glucose. Relative mRNA expression and protein levels were detected by RT-qPCR and western blot analysis, respectively. Cell proliferation and apoptosis were assessed using CCK8 and Annexin V-FITC apoptosis detection kits, respectively. DCFH-DA assay was used to measure reactive oxygen species (ROS) accumulation. The results indicated that high glucose treatment reduced the expression of mRNA and protein of SIRT3 and PPAR- $\alpha$  in AC16 cells. Moreover, high glucose inhibited cell proliferation, as well as induced apoptosis, intracellular hydrogen peroxide production, and JNK1/2 phosphorylation. These effects were antagonized by SIRT3 overexpression or treatment with the PPAR- $\alpha$  agonist, Wy14643. Conversely, inhibition of SIRT3 via 3-TYP led to similar phenomena as those induced by high glucose treatment in AC16 cells, which were blocked by Wy14643. Lastly, chromatin immunoprecipitation (ChIP) and luciferase assays demonstrated SIRT3 as a direct target of PPAR- $\alpha$ . Taken together, the results provide evidence for an important role

of SIRT3 in high glucose-induced cardiomyocyte injury and regulation of JNK1/2 signaling. Further, SIRT3 is a direct downstream target of PPAR- $\alpha$ .

## Introduction

Diabetes mellitus is a life-threatening and complex metabolic disorder affecting multiple systems in the body. The morbidity rate of diabetes in China is 9.7% and continues to grow (1). Diabetes is considered a major risk factor for coronary heart disease, with the mortality rate of cardiovascular disease in diabetic patients estimated at 65% (2). Diabetic cardiomyopathy (DCM) is a type of diabetic heart disease first discovered in 1972 (3) and appears as myocardial dysfunction in diabetic patients without valvular heart disease, hypertension, or coronary artery disease (CAD) (3,4). Epidemiologic studies indicated that DCM was closely associated with progressive increase of relative wall thickness, left atrium and left ventricular mass and impaired glucose tolerance (4). Research on the etiopathogenesis of DCM revealed that hyperglycemia played a decisive role in the development of DCM (5).

Diabetes mellitus is characterized by several important changes in micro-vascular architecture, including subendothelial matrix deposition, abnormal capillary permeability, fibrosis surrounding arterioles and micro-aneurysm formation (6). Hyperglycemia can activate protein kinase C and stimulate vascular endothelial cells to enhance the production of vasoconstrictor prostanoids, which can promote endothelial dysfunction, as well as ventricular and myocardial hypertrophy (7,8). On the other hand, hyperglycemia also can aggravate intracellular oxidative stress, which then induces myocardial injury (9). Moreover, oxidative stress is exacerbated by reactive oxygen species (ROS) production in the mitochondria of diabetic cardiac tissues (10). Aberrant ROS production also increases pro-inflammatory response and myocardial apoptosis (11).

Peroxisome proliferator-activated receptors (PPARs) are important nuclear hormone receptors (12) consisting of 3 types (PPAR- $\alpha$ , PPAR- $\gamma$  and PPAR- $\delta$ ), each of which is encoded by separate genes and distinguished by specific ligands, functions, and distributions (12). A previous study described that transgenic mice with PPAR- $\alpha$  overexpression develop cardiomyopathy similar to the diabetes mellitus condition (13).

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Acetylation is a critical mechanism for regulating the activity of widespread enzymes associated with mitochondrial metabolism, such as the fatty acid oxidation, citric acid cycle, antioxidant defense and the electron transport system (14–16). Sirtuin 3 (SIRT3) is the primary deacetylase in the mitochondria (17). Intriguingly, SIRT3 activity and expression were found to be decreased in an obesity animal model, possibly leading to obesity-related reduction of antioxidant defense and oxidative metabolism (18). However, very little is known about the function and mechanism of SIRT3 in DCM. In this study, the AC16 cell line was used as a cardiomyocyte model coupled with SIRT3 overexpression or inhibition (by 3-TYP), as well as treatment with PPAR- $\alpha$  agonist (Wyl4643) or antagonist (GW6471) under high glucose/euglycemia conditions. The interaction between SIRT3 and PPAR- $\alpha$  was investigated.

## Materials and methods

**Chemicals and reagents.** Glucose was from Sigma-Aldrich (Shanghai, China). Annexin V-FITC apoptosis detection kit, CCK8 assay kit and cellular reactive oxygen species detection kit were obtained from Beyotime Biotechnology. 3-TYP and Wyl4643 were from Selleck Chemicals, and GW6471 from R&D Systems China. TRIzol reagent and reverse transcription kits were obtained from Thermo Fisher Scientific, Inc. PPAR- $\alpha$  antibody was from Abcam. SIRT3, cleaved caspase-3, Bax, Bcl2, JNK1/2, phosphorylated JNK1/2 (p-JNK1/2) and glyceraldehyde 3-phosphate dehydrogenase (GAPDH) antibodies were from Cell Signaling Technology.

The study was approved by the Ethics Committee of Central Hospital of Minhang District (Shanghai, China).

**AC16 cells.** AC16 human cardiomyocyte cell line was obtained from the Chinese Academy of Sciences Cell Bank (<http://www.cellbank.org.cn/>, Shanghai, China). Cells were incubated with DMEM media (HyClone) supplemented with 100 U/ml penicillin and 10% fetal bovine serum (FBS; Gibco; Thermo Fisher Scientific, Inc.) at 37°C (5% CO<sub>2</sub>). AC16 cells at logarithmic phase were used for follow-up experiments.

### Experimental groups

**Effects of glucose treatment on SIRT3 and PPAR- $\alpha$  expression in AC16 cells.** AC16 cells were treated with different concentrations of glucose (5.5, 10, 30 and 50 mM, added into glucose-free DMEM media) for 48 h. Relative mRNA expression and protein levels of SIRT3 and PPAR- $\alpha$  were detected at 24 and 48 h after treatment.

**Effects of SIRT3 overexpression and PPAR- $\alpha$  activation/inhibition on AC16 cells under high glucose condition.** AC16 cells were divided into 5 groups: Control group (5.5 mM glucose); G 30 mM + EPC group (30 mM glucose + empty plasmid control); G 30 mM + SIRT3 OE group (30 mM glucose + SIRT3 overexpression); G 30 mM + SIRT3 OE + GW6471 group (30 mM glucose + SIRT3 overexpression + PPAR- $\alpha$  antagonist, GW6471); and G 30 mM + Wyl4643 group (30 mM glucose + PPAR- $\alpha$  agonist, Wyl4643). First, glucose (5.5 or 30 mM) was added into glucose-free DMEM media in respective groups for 24 h. Then, SIRT3 OE or EPC lentivirus (JRDun Biotech) (Table I), Wyl4643 (100  $\mu$ M, dissolved in DMSO) and GW6471 (10  $\mu$ M,

dissolved in DMSO) were added into AC16 cells, respectively for 72 h. Proliferation of AC16 cells was examined at 0, 24, 48 and 72 h after lentivirus treatment. Cell apoptosis and reactive oxygen species levels were measured at the end of the experiment, along with SIRT3 mRNA expression and SIRT3, cleaved caspase-3, Bax, Bcl2, JNK1/2 and p-JNK1/2 protein levels.

**Effects of SIRT3 inhibitor (3-TYP) and PPAR- $\alpha$  agonist (Wyl4643) on AC16 cells under euglycemia condition.** AC16 cells were divided into 3 groups, and received the following treatments: 5.5 mM glucose (control group); 5.5 mM glucose + 30 mM 3-TYP (G (5.5 mM) + 3-TYP group); and 5.5 mM glucose + 30 mM 3-TYP + 100  $\mu$ M Wyl4643 [G (5.5 mM) + 3-TYP + Wyl4643 group]. AC16 cells in each group were treated with glucose (added into glucose-free DMEM media) for 24 h. Subsequently, the indicated chemicals were added and cells were cultured for 72 h. Proliferation of AC16 cells was assessed at 0, 24, 48 and 72 h after treatment. Cell apoptosis, reactive oxygen species levels, and JNK1/2 and p-JNK1/2 protein levels were examined at the end of the experiment.

**Interaction between PPAR- $\alpha$  protein and SIRT3 promoter.** Cells were treated with Wyl4643 (100  $\mu$ M), GW6471 (10  $\mu$ M) or DMSO for 48 h. Chromatin immunoprecipitation (ChIP) and Dual-Lucy Assay kit (Promega, Beijing, China) were employed to test the interaction between PPAR- $\alpha$  and SIRT3.

### Experimental methods

**Cell proliferation and apoptosis assay.** Cultured AC16 cells were harvested at relative time-points. Proliferation of AC16 cells was examined using CCK8 assay kit following the manufacturer's instructions. Cell absorbance was measured at 450 nm using a microplate reader (Thermo Fisher Scientific, Inc.).

Cultured AC16 cells were harvested and incubated with FITC-labelled Annexin V and PI at 25°C for 20 min following the manufacturer's instructions. Subsequently, the intensity of Annexin V or PI fluorescence was analyzed by FACScan (Becton-Dickinson). For each sample, 10,000 cells were tested.

**Dichlorodihydrofluorescein diacetate (DCFH-DA) flow cytometry.** ROS production was assessed using DCFH-DA assay according to a previous study (19). Briefly, AC16 cells (10<sup>6</sup>/ml) were cultured with 5  $\mu$ M of 2'-7'-dichlorodihydrofluorescein diacetate (DCFH-DA) for 20 min at 37°C (forward-reverse mixing once per 3 min). DCFH-DA (non-fluorescent) entered cells and hydrolyzed into cell-impermeable and non-fluorescent DCFH. DCFH was oxidized into highly fluorescent dichlorofluorescein (DCF) by intracellular ROS. The green fluorescence intensity was proportional to ROS level. DCF fluorescence was assayed at 525 nm after excitation of cells at 480 nm using flow cytometry analysis (Becton-Dickinson). Results were expressed as the ratio of fluorescence intensity of AC16 cells in the other groups to that in control group.

**ChIP and luciferase assay.** ChIP is a powerful tool to investigate interactions between intracellular DNA and specific proteins, and detect their genomic localization (20). In the present study, interaction between PPAR- $\alpha$  protein and SIRT3 promoter was tested by ChIP assay according to a previous study (21). Briefly, cell samples were subjected to standard cross-linking, chromatin shearing, immunoprecipitation,

Table I. SIRT3 (NM\_001017524)/SIRT3 promoter sequence and primers.

Gene name	Primer sequence (5'-3')
SIRT3 CDS	F: CGGAATTCATGGCGTTCTGGGGTTG R: CGGGATCCCTATTTGTCTGGTCCATCAAGC
Semi-qRT-PCR (ChIP)	F: CCCCTCCGTCTCCCTCTATC R: CAACCCCAAGAACGCCATG
SIRT3 promoter	F: CCCTCGAGACGGCGGAAGTGGTTG R: CCAAGCTTTCCTGCCCCAAG

F, forward; R, reverse; SIRT3, sirtuin 3; ChIP, chromatin immunoprecipitation.

reverse cross-linking, DNA precipitation and PCR analysis. The PCR products were determined using semi-quantitative RT-PCR (primers are shown in Table I).

Cells were co-transfected with a dual-luciferase reporter plasmid containing SIRT3 promoter-reporter plasmid (JRDun Biotech), in combination with Wy14643 or GW6471 treatment for 48 h in 24-well plates. Luciferase activity was measured using the Dual-Lucy Assay Kit (Promega) following the manufacturer's protocol (primers are shown in Table I).

**RT-qPCR.** Relative mRNA expression was determined by RT-qPCR. Total mRNA was isolated using TRIzol, and 2  $\mu$ g of total RNA from each sample was reverse transcribed into cDNA using First Strand cDNA Synthesis kit (Thermo Fisher Scientific, Inc.) on RT-qPCR machine (ABI-7300; Applied Biosystems).

Primers used for the RT-qPCR are shown in Table II. Relative mRNA expression was evaluated by  $2^{-\Delta\Delta Ct}$  relative quantitative analysis against GAPDH.

**Western blot analysis.** Total protein from each sample was extracted, and protein concentration was measured using BCA protein assay kit. Protein (30  $\mu$ g) was separated by 10% SDS-PAGE and transferred onto PVDF membranes, which were then blocked in 5% skimmed milk at 25°C for 1 h, followed by incubation with primary antibodies overnight at 4°C. After washing, membranes were incubated with secondary antibodies for 1 h. Finally, protein bands were visualized using ECL-detection kit (Beyotime Biotechnology). The following primary antibodies were used: PPAR- $\alpha$  (1:1,000), SIRT3 (1:1,000), cleaved caspase-3 (1:1,000), Bax (1:1,000), Bcl2 (1:1,000), JNK1/2 (1:1,000), p-JNK1/2 (1:1,000) and GAPDH (1:1,000; CST). GAPDH served as a loading control.

**Statistical analysis.** Data were expressed as mean  $\pm$  SD (n=3). Difference was evaluated using ANOVA. Duncan multiple range test was determined using SPSS 20.0 software (IBM Corp). P<0.05 was considered as statistically significant.

## Results

**Glucose treatment downregulates expression of SIRT3 and PPAR- $\alpha$  in AC16 cells.** In the current study, both mRNA

Table II. Primers used in real-time fluorogenic PCR assays.

Gene name	Primer sequence (5'-3')
SIRT3	F: CCTTGGCTTGGCATCCTC
SIRT3	R: GCACAAGGTCCCAGCATCTC
PPAR- $\alpha$	F: TCACGGACACGCTTTCACC
PPAR- $\alpha$	R: CCCCAGATTCTACATTCG
GAPDH	F: AATCCCATCACCATCTTC
GAPDH	R: AGGCTGTTGTCATACTTC

F, forward; R, reverse; SIRT3, sirtuin 3; PPAR- $\alpha$ , peroxisome proliferator-activated receptor- $\alpha$ ; GAPDH, glyceraldehyde 3-phosphate dehydrogenase.

expression and protein levels of SIRT3 and PPAR- $\alpha$  in AC16 cells showed no change with time under euglycemia condition (5.5 mM of glucose). However, with increasing dose ( $\geq 30$  mM), SIRT3 and PPAR- $\alpha$  mRNA expression (Fig. 1A and B) and protein levels (Fig. 1C-E) were reduced, and further diminished over time (Fig. 1).

**Effects of SIRT3 OE and PPAR- $\alpha$  activation/inhibition on AC16 cells under high glucose (30 mM) condition.** The regulatory function of SIRT and whether it was regulated by PPAR- $\alpha$  under high glucose (30 mM) condition were investigated. The present results indicated that the proliferation of AC16 cells was inhibited by high glucose (30 mM) treatment at 48 h (34.67 $\pm$ 3.21%) and 72 h (47.13 $\pm$ 5.42%), but was antagonized by SIRT3 OE, SIRT3 OE + GW6471 and Wy14643 treatment (Fig. 2A). Moreover, intracellular hydrogen peroxide production and apoptosis were induced by glucose (30 mM) treatment for 72 h, but reduced by SIRT3 OE, SIRT3 OE + GW6471 and Wy14643 treatment (Fig. 2B-D). In addition, SIRT3 mRNA expression and protein level were diminished by high glucose (30 mM) but increased by SIRT3 OE, SIRT3 OE + GW6471 and Wy14643 treatment (Fig. 3A-C). Moreover, the protein levels of cleaved caspase-3, Bax and p-JNK1/2 displayed trends consistent with apoptosis, whereas Bcl2 protein level showed a trend opposite to Bax. These results suggested that SIRT3 might be positively regulated by PPAR- $\alpha$  in AC16 cells (Fig. 3B and C).

**Effects of 3-TYP and Wy14643 treatments on AC16 cells under euglycemia condition.** Based on the above findings, it was hypothesized that the downregulation of SIRT3 might play an important role during the process of high glucose-induced myocardial injury. Therefore, the effects of 3-TYP and Wy14643 treatments on AC16 cells under euglycemia condition were analyzed. In the present study, the inhibition of SIRT3 (by treatment with 3-TYP) in AC16 cells led to similar results under euglycemia condition as high glucose treatment, including inhibition of cell proliferation (Fig. 4A), stimulation of intracellular hydrogen peroxide production and induction of apoptosis (Fig. 4B-D), and increased phosphorylation of JNK1/2. Furthermore, these phenomena could be relieved by activation of PPAR- $\alpha$  (Fig. 5A and B).

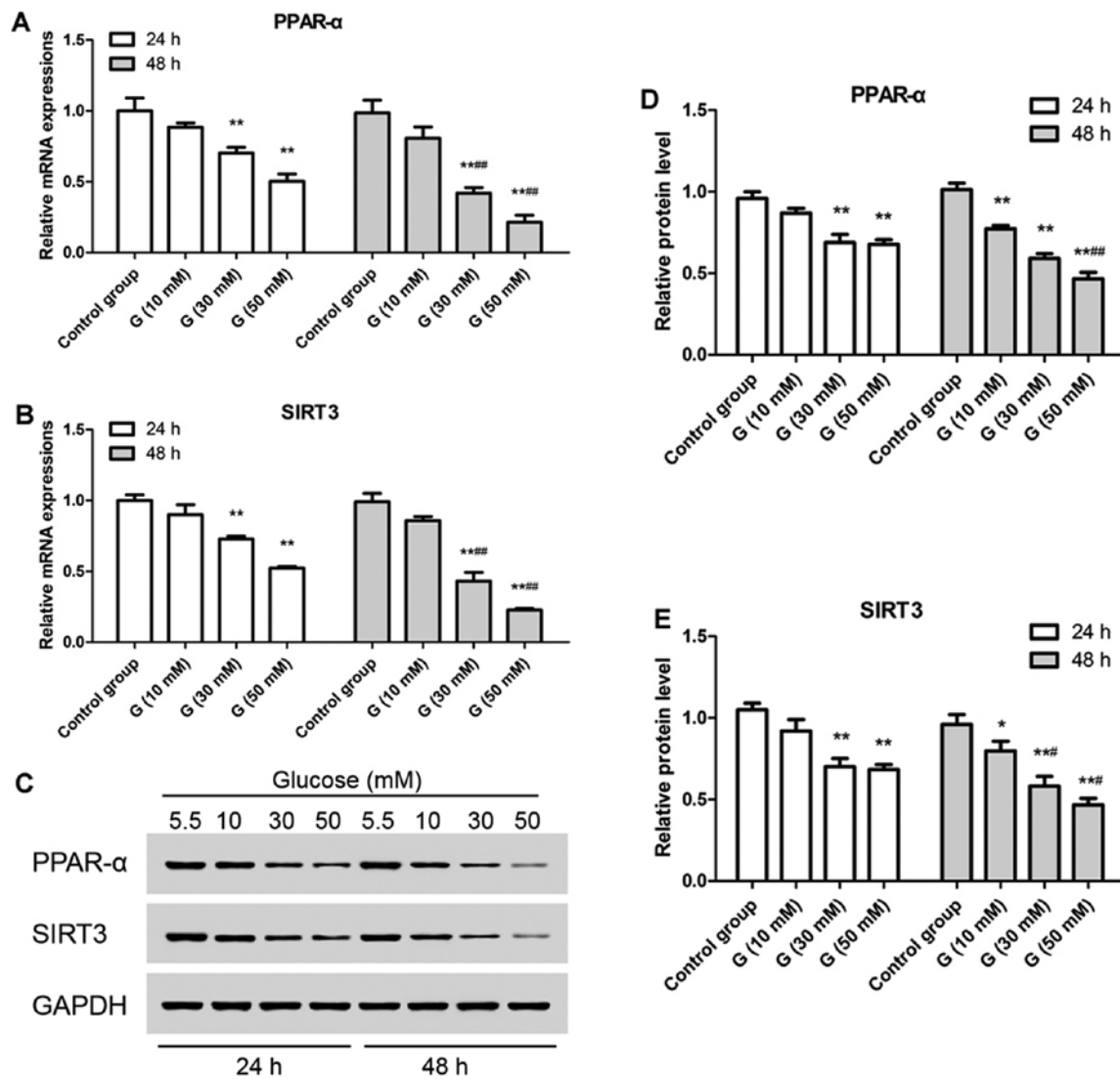


Figure 1. (A and B) Expression of mRNA and (C-E) protein levels of SIRT3 and PPAR- $\alpha$  were decreased by high glucose treatment (n=3). \*P<0.05 and \*\*P<0.01 compared with euglycemia condition (5.5 mM of glucose treatment) (n=3). #P<0.05 and ##P<0.01 compared with control group at different time points. PPAR- $\alpha$ , peroxisome proliferator-activated receptor- $\alpha$ ; SIRT3, sirtuin 3.

SIRT3 is a direct, positively regulated target of PPAR- $\alpha$ . The intracellular interaction between SIRT3 promoter and PPAR- $\alpha$  was tested using ChIP and luciferase assay. The present results suggested that intracellular PCR production of SIRT3 promoter was increased by PPAR- $\alpha$  agonist and decreased by PPAR- $\alpha$  antagonist (Fig. 6A and B). Furthermore, the ratio of Firefly and *Renilla* luciferase signals showed similar trends (Fig. 6C).

## Discussion

DCM has become a worldwide public health concern, yet the mechanisms and occurrence of DCM remain unknown (2,5). Herein, it was found that abnormally low expression of SIRT3 may play an important role in DCM. SIRT3 expression was weakened under high glucose condition in AC16 cells. Whereas, high glucose inhibited proliferation of AC16 cells and enhanced apoptosis and intracellular hydrogen peroxide production. These phenomena could be improved by SIRT3 overexpression. SIRT3 inhibition led to similar phenomena in

AC16 cells under euglycemia condition as under high glucose treatment.

The production of ROS is considered to be a contributing factor in the occurrence and progression of diabetic cardiomyopathy (22). The strategy of enhancing mitochondrial ROS scavenger system has been clinically proven to be effective in reducing cardiac dysfunction caused by diabetes (22). SIRT3 plays a key role in improving mitochondrial dysfunction and oxidative stress via the MAPK pathway (23,24). SIRT3 is able to maintain ROS homeostasis by regulating diverse mitochondrial enzymes, including superoxide dismutase 2 (SOD2), which can transform harmful superoxide free radicals into non-toxic hydrogen peroxide or oxygen (25). In the present study, ROS production was negatively correlated with SIRT3 expression, which may be related to the downregulation of JNK1/2 phosphorylation.

PPAR- $\alpha$  is a member of PPAR nuclear transcription factor family that is enriched in myocardium (26). PPAR- $\alpha$  maintains antioxidant defense and oxidant equilibrium, and exhibits anti-inflammatory action (26). PPAR- $\alpha$  also plays a crucial role

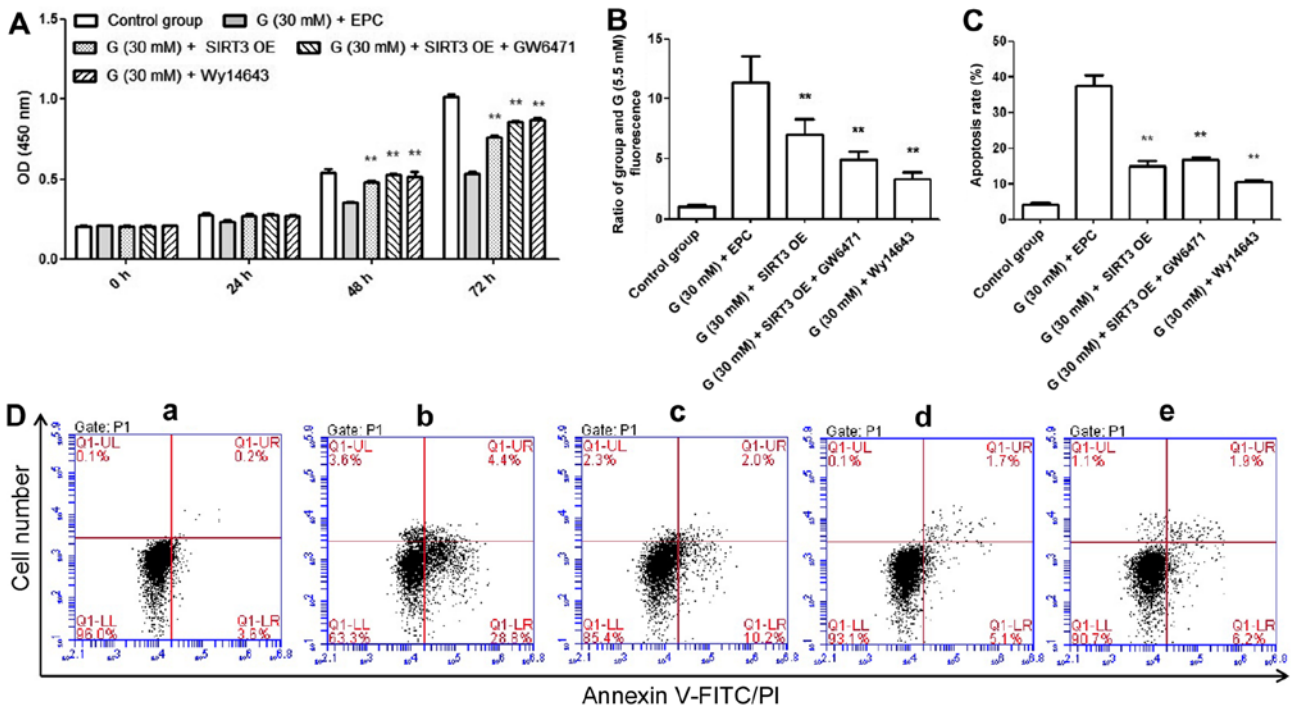


Figure 2. Effects of SIRT3 overexpression and PPAR- $\alpha$  activation/inhibition in AC16 cells under high glucose (30 mM) condition (n=3). (A) Effects of different treatments on the proliferation of AC16 cells. (B) Effects of different treatments on intracellular hydrogen peroxide production. Results are expressed as the ratio of fluorescence intensity of other groups to that of glucose (5.5 mM) group. (C) Effects of different treatments on cell apoptosis. (D) Flow cytometry image of apoptotic cells. \*\*P<0.01 compared with glucose (30 mM) + EPC group. a, control group; b, G (30 mM) + EPC; c, G (30 mM) + SIRT3 OE; d, G (30 mM) + SIRT3 OE + GW6471; e, G (30 mM) + Wy14643. PPAR- $\alpha$ , peroxisome proliferator-activated receptor- $\alpha$ ; SIRT3, sirtuin 3.

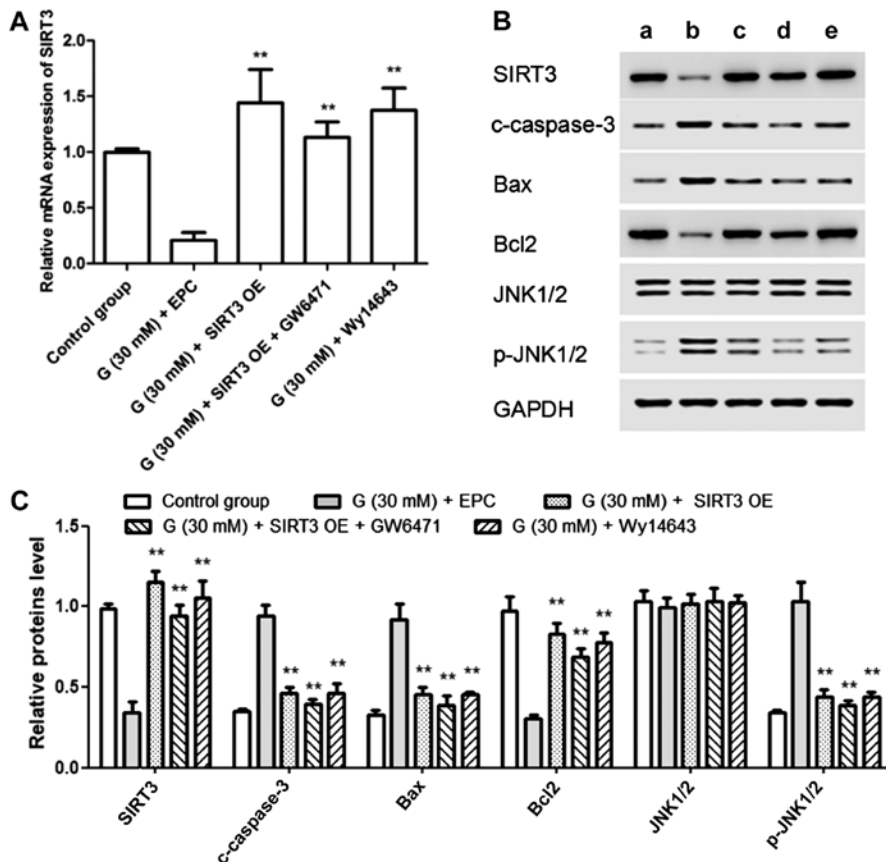


Figure 3. Effects of different treatments on SIRT3 mRNA and protein expression. (A) Effects of different treatments on SIRT3 mRNA expression. (B and C) Effects of different treatments on protein expression as assessed by western blotting. a, glucose (5.5 mM) group treatment group (control group); b, glucose (30 mM) + EPC group; c, glucose (30 mM) + SIRT3 OE group; d, glucose (30 mM) + SIRT3 OE + GW6471 group; e, glucose (30 mM) + Wy14643 group. \*\*P<0.01 compared with glucose (30 mM) + EPC group; SIRT3, sirtuin 3.

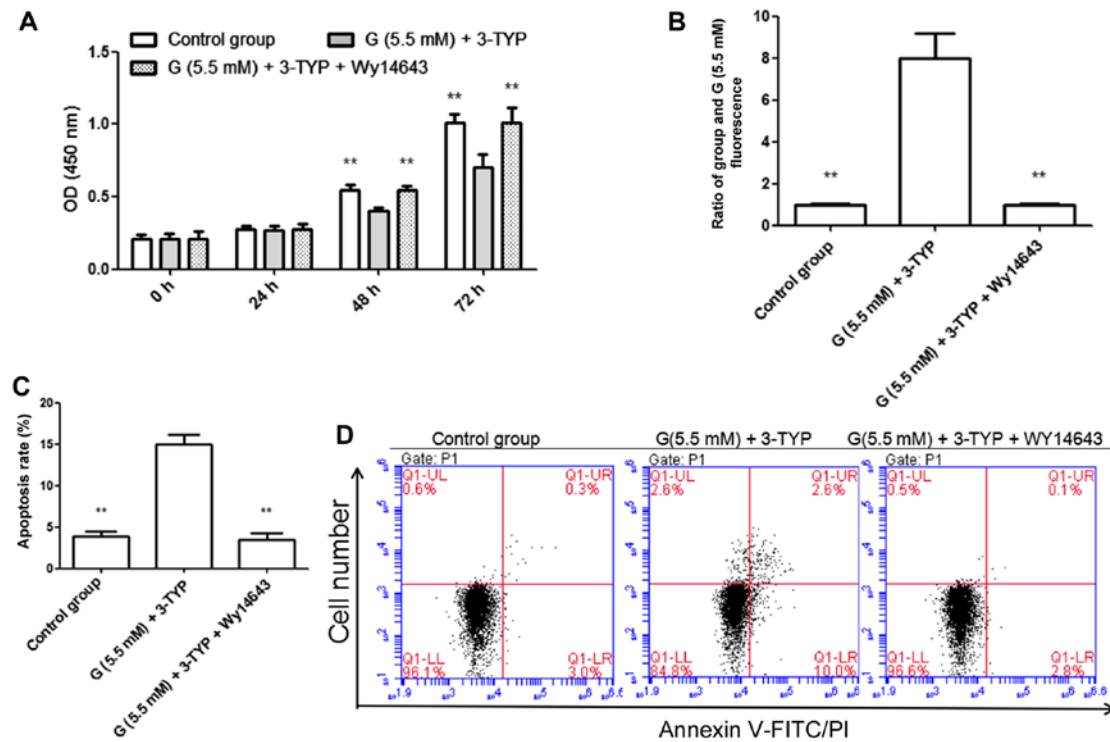


Figure 4. Effects of 3-TYP and Wy14643 treatments on AC16 cells under euglycemia condition (n=3). (A) Effects of 3-TYP and Wy14643 on proliferation of AC16 cells under euglycemia condition. (B) Effects of 3-TYP and Wy14643 on intracellular hydrogen peroxide production under euglycemia condition. (C) Effects of 3-TYP and Wy14643 on cell apoptosis under euglycemia condition. (D) Flow cytometry image of apoptotic cells. \*\*P<0.01 compared with glucose (5.5 mM) + 3-TYP treatment group.

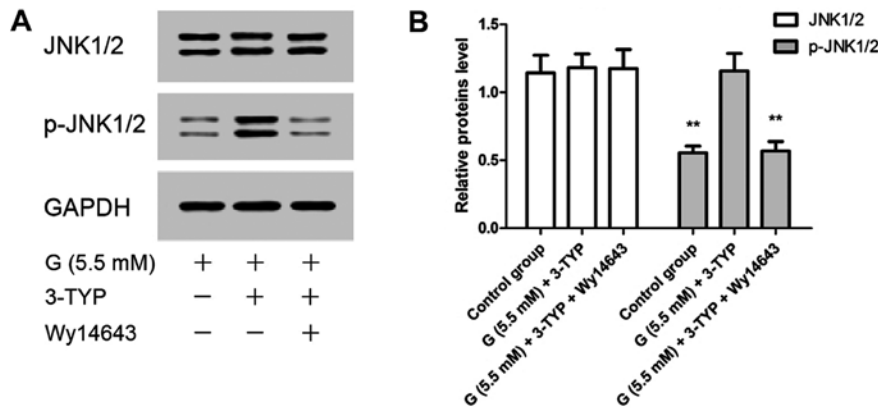


Figure 5. Effects of 3-TYP and Wy14643 on protein levels of JNK1/2 and p-JNK1/2 under euglycemia condition. (A) Protein bands of western blotting and (B) quantification. \*\*P<0.01 compared with glucose (5.5 mM) + 3-TYP treatment group.

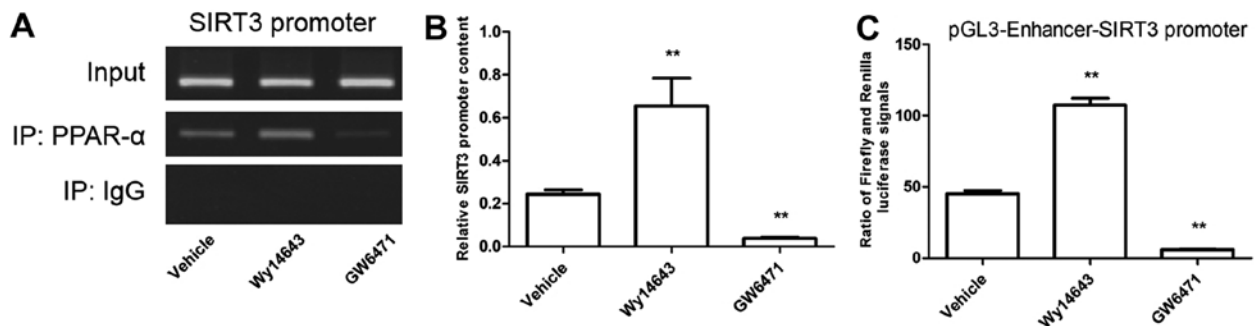


Figure 6. SIRT3 is a direct, positively regulated target of PPAR- $\alpha$ . (A) Electrophoretic bands of PCR-amplified products in ChIP assay. (B) Statistical analysis of ChIP assay. (C) Results of double fluorescent reporter gene system. \*\*P<0.01 compared with vehicle group (n=3). PPAR- $\alpha$ , peroxisome proliferator-activated receptor- $\alpha$ ; SIRT3, sirtuin 3; ChIP, chromatin immunoprecipitation.

in the regulation of lipoprotein transport and assembly, as well as mitochondrial fatty acid oxidation (27). Decrease in PPAR- $\alpha$  expression has been suggested to be a self-adaptation process that transforms the metabolic substrate of cardiac energy from fatty acid to glucose (28). It was found in this study that PPAR- $\alpha$  expression was downregulated in AC16 cells under high glucose condition. Furthermore, PPAR- $\alpha$  agonists improved myocardial cell injury induced by high glucose. Results from ChIP and luciferase assay attested that SIRT3 was a direct, positively regulated target of PPAR- $\alpha$ .

In conclusion, the present results indicated that diminished expression of SIRT3 played an important role during high glucose-induced injury in AC16 cells. The function of SIRT3 and its interaction with PPAR- $\alpha$  were further evaluated in AC16 cells, and the results demonstrated that SIRT3 was a downstream target of PPAR- $\alpha$ .

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### Availability of data and materials

The datasets used and/or analyzed during the present study are available from the corresponding author on reasonable request.

### Authors' contributions

XZ contributed to the conception and design of the study, supervised the progress of individual study experiments and wrote the manuscript. KC and GY performed cell proliferation and apoptosis assay and flow cytometry. ZW, QS and DY were responsible for RT-qPCR and western blot analysis. PL, WH and YC contributed to observation indexes analysis. The final version was read and adopted by all the authors. All authors read and approved the final manuscript.

### Ethics approval and consent to participate

The study was approved by the Ethics Committee of Central Hospital of Minhang District (Shanghai, China).

### Patient consent for publication

Not applicable.

### Competing interests

The authors declare that they have no competing interests.

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