ZP2495 Protects against Myocardial Ischemia/Reperfusion Injury in Diabetic Mice through Improvement of Cardiac Metabolism and Mitochondrial Function: The Possible Involvement of AMPK-FoxO3a Signal Pathway

Shuang Li,1,2 Hao Wu,3 Dong Han,1 Mingming Zhang,5 Na Li,4 Weihua Yu,3 Dongdong Sun,4 Zhongchan Sun,4 Sai Ma,4 Erhe Gao,6 Congye Li,4 Min Shen,4 and Feng Cao1

1Department of Cardiology & National Clinical Research Center of Geriatrics Disease, Chinese PLA General Hospital, Beijing 100853, China
2Department of Cardiology, Chengdu Military General Hospital, Chengdu 610083, China
3Department of Toxicology, School of Public Health, Fourth Military Medical University, Xi’an, Shaanxi 710032, China
4Department of Cardiology, Xijing Hospital, Fourth Military Medical University, Xi’an, Shaanxi 710032, China
5Department of Cardiology, Tangdu Hospital, Fourth Military Medical University, Xi’an, Shaanxi 710032, China
6Center of Translational Medicine, Temple University School of Medicine, Philadelphia, PA 19107, USA

Correspondence should be addressed to Feng Cao; fengcao8828@163.com

Shuang Li, Hao Wu, and Dong Han contributed equally to this work.

Received 23 April 2017; Revised 23 August 2017; Accepted 14 September 2017; Published 17 January 2018

Academic Editor: Jacek Zielonka

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Coronary heart disease patients with type 2 diabetes were subject to higher vulnerability for cardiac ischemia-reperfusion (I/R) injury. This study was designed to evaluate the impact of ZP2495 (a glucagon-GLP-1 dual-agonist) on cardiac function and energy metabolism after myocardial I/R injury in db/db mice with a focus on mitochondrial function. C57BLKS/lepr+/lepr+ (BKS) and db/db mice received 4-week treatment of glucagon, ZP131 (GLP-1 receptor agonist), or ZP2495, followed by cardiac I/R injury. The results showed that cardiac function, cardiac glucose metabolism, cardiomyocyte apoptosis, cardiac mitochondrial morphology, and energetic transition were improved or ameliorated by ZP2495 to a greater extent than that of glucagon and ZP131.

In vitro study showed that ZP2495, rather than glucagon, alleviated mitochondrial depolarization, cytochrome C release, and mitochondria ROS generation in neonatal rat ventricular myocytes subjected to high-glucose and simulated I/R injury conditions, the effects of which were weaker in the ZP131 group. Furthermore, the expressions of Akt, FoxO3a, and AMPK phosphorylation were elevated by ZP2495 to a greater extent than that of ZP131. In conclusion, ZP2495 may contribute to the improvement of cardiac function and energy metabolism in db/db mice after myocardial I/R injury by improving mitochondrial function possibly through Akt/FoxO3a and AMPK/FoxO3a signal pathways.

1. Introduction

Type 2 diabetes is one of the strong independent risk factors for cardiovascular disease and death [1, 2]. Lots of studies including our previous studies have demonstrated that patients with type 2 diabetes have higher vulnerability of cardiac ischemia/reperfusion (I/R) injury as a result of the exposure to abnormal substrate and cytokines [3, 4]. However, effective strategies which can reduce cardiac I/R injury under diabetic conditions are not well developed in a clinic setting.
Glucagon-like peptide-1 (GLP-1) is derived from a proglucagon precursor and secreted by intestinal L-cells in response to oral nutrient ingestion, which acts through G protein-coupled receptor (GLP-1R) on pancreatic beta-cells to elicit glucoregulatory and insulinosotropic actions [5, 6]. GLP-1R agonists have been reported to have cardiac and vascular actions in rodents and humans, including effects on myocardial contractility, blood pressure, cardiac output, and cardioprotection [7–9]. Glucagon exerts biochemical and physiological effects on heart muscle partly by stimulating Ca\(^{2+}\) currents via CAMP production and inhibition of phosphodiesterases [9–11], leading to increased contractility. Although cardiac contractility can be enhanced by glucagon administration, it will increase cardiac mortality since glucagon increases cardiomyocyte consumption of energy after administration, it will increase cardiac mortality since glucose to fatty acid metabolism contributes to the severity of an ischemic injury and can impair functional recovery during and following ischemia. This is particularly evident during the reperfusion phase of the ischemic myocardium [15–17].

As a result, diabetes-induced changes in energy metabolism have the potential to significantly impact on the ability of the heart to withstand an I/R injury [18]. This study was designed to evaluate the impact of a glucagon–GLP-1 dual-agonist ZP2495 on cardiac function and energy metabolism after myocardial I/R injury in db/db mice and to investigate the underlying mechanisms involved.

2. Materials and Methods

2.1. Animal Modeling and Grouping. Male C57BLKS/J-lepr\(^{db}\)/lepr\(^{db}\) diabetic (db/db; stock number 000642) and nondiabetic C57BLKS/J-lepr\(^{+/}\)/lepr\(^{+}\) (BKS; stock number 000662) mice (8–10 weeks) were obtained from Jackson Laboratory (Bar Harbor, Maine, USA) at 6 weeks of age. Animals were housed in a 12:12 h light-dark cycle at 22°C with free access to food and water. db/db mice at the age of 12–16 weeks (50–60 g) when they had developed overt diabetes were used. All experiments were performed in accordance with the National Institutes of Health Guidelines on the Use of Laboratory Animals and were approved by the Fourth Military Medical University Committee on Animal Care (ID 2013052).

db/db mice were randomly allocated into the following groups with n = 20 each: (1) BKS + sham (sham); (2) db/db + sham (db/db); (3) db/db + I/R (I/R); (4) db/db + ZP2495 + I/R (ZP2495); (5) db/db + glucagon + I/R (glucagon); and (6) db/db + ZP131 + I/R (ZP131). The ZP2495 group was subcutaneously injected daily with ZP2495 at 70 nmol/kg of body weight for 4 weeks. The glucagon group received subcutaneous injections of glucagon at 70 nmol/kg of body weight daily for 4 weeks. The mice in the sham group were injected with saline at the same volume instead.

Mice were anesthetized with 2% isoflurane in 1.5 L/min O\(_2\). After exteriorizing the heart via a left thoracic incision, myocardial ischemia/reperfusion was induced by ligation of the left descending coronary artery for 30 minutes as previously described [19]. After 30 minutes of ischemia, hearts were harvested 3 hours (for TUNEL, caspase activity, and Western blot assay) or 24 hours (for echocardiographic, hemodynamic, and metabolic assessments and infarct size measurement) after the myocardium reperfusion.

2.2. Echocardiographic and Hemodynamic Measurements. Cardiac function was determined by echocardiography (VisualSonics, Canada) and invasive hemodynamic evaluation by a 1.4 French micromanometer (Millar Instruments, USA) at different time points after ischemia/reperfusion.

Echocardiography was performed under anesthesia using a 30 MHz transducer on a Vevo 2100 ultrasound system (VisualSonics, Canada) as previously described [20].

2.3. \(^{18}\)F-Fluorodeoxyglucose Positron Emission Tomography/Computed Tomography to Assess Cardiac Glucose Metabolism. Positron emission tomography/computed tomography (PET/CT) scanning was performed using a Nano PET/CT (Mediso, Hungary) as described previously. Before imaging study, mice were maintained under fasting condition for 12–14 h, and fasting blood glucose levels were maintained between 6.0 and 7.5 mmol/L. Every animal was injected with 3.7 MBq (100 μCi) of \(^{18}\)F-FDG via tail vein injection. To obtain the best signal-to-background ratio, the animals were anesthetized under 2% isoflurane in 1.5 L/min O\(_2\) and acquired for whole body static PET/CT imaging at 30 min postadministration with 10 min acquisition time, energy window at 400–600 keV, and coincidence relation with 1:3. CT scan was subsequently obtained with 45 kV and 179 mA for 10 min. The PET images were reconstructed using ordered subset expectation maximization reconstruction (OSEM) with the SSRB 2D LOR algorithm with decay correction, CT imaging-based attenuation correction, and random corrections from raw framed sinograms. The voxel size of the PET image was 0.796 × 0.861 × 0.861 mm, for a total of 128 × 128 × 159 voxels. After PET-CT registration, 3-dimensional regions of interest (3D-ROI) were drawn over the heart region on whole body axial PET/CT fusion images. Relative accumulation of the radioactivity in particular regions of interest was expressed as standardized uptake value (SUV). Nano PET/CT images were analyzed using InterView™ Fusion software (Mediso, Hungary).

2.4. Myocardial Infarct Size Measurement. Myocardial infarct size was evaluated by Evans Blue/TTC staining as previously described [21].

2.5. Detection of Cardiomyocyte Apoptosis in Hearts. TUNEL staining and caspase-3 activity assay were used to determine cardiomyocyte apoptosis as previously described [22].
2.6. Transmission Electron Microscopy (TEM). Ultrathin sections were observed under a TEM (JEM-2100, JEOL, Tokyo, Japan) at 60 kV. Mitochondria were imaged at ×10,000 (10 K) and ×40,000 (40 K) magnifications.

2.7. Measurement of Neonatal Rat Ventricular Myocyte (NRVM) Apoptosis Induced by Simulated Ischemia/Reperfusion (S/IR) Injury. Primary cultures of NRVMs were obtained from 1- or 2-day-old SD rats as described previously [23]. Media were replaced to different conditions after confluence: normal glucose medium (5.5 mmol/L), high-glucose medium (25 mmol/L), and high-glucose plus ZP2495 (10⁻⁷ mol/L), glucagon (10⁻⁷ mol/L), or ZP131 (10⁻⁷ mol/L). Simulated I/R (S/IR) was performed by transferring cardiomyocytes into an ischemic buffer adapted from Esumi et al. [24] containing (in mmol/L) NaCl 137, KCl 3.8, MgCl₂ 0.49, CaCl₂·2H₂O 0.9, and HEPES 4 supplemented with deoxyglucose 10, sodium dithionate 0.75, KCl 12 and lactate 20, and pH 6.5, for 3 h in a humidified cell culture incubator (21% O₂, 5% CO₂, and 94% N₂, 37°C). After simulated ischemia, cells were transferred to DMEM for 3 h in a humidified cell culture incubator (21% O₂, 5% CO₂, and 74% N₂, 37°C) to simulate reperfusion.

NRVM apoptosis was determined using an Annexin V-FITC/PI Kit (Merck, USA) as previously described [25].

2.8. Measurement of Mitochondrial Membrane Potential (ΔΨm) (MMP). The impact of ZP2495, glucagon, or ZP131 on mitochondrial ΔΨm in cardiomyocytes was determined by sensitive and relatively mitochondrion-specific lipophilic cationic probe fluorochrome JC-1 as previously described [26]. Cardiomyocytes were incubated with JC-1 Mitochondrial Potential Sensors (Invitrogen, USA), and images were visualized and photographed with confocal microscopy (Olympus, Japan). In order to measure the internalization of JC-1, cardiomyocytes were stained with the JC-1 Assay Kit for flow cytometry (Mitoprobe, Invitrogen, USA) according to the manufacturer’s protocol and analyzed by subsequent flow cytometry (BD Bioscience, USA). To exclude cell debris and doublets, cells were gated the same number in each group, 1 × 10⁵ gated events per sample were collected from three to four independent samples per treatment condition, and the experiments were repeated at least three times. The presented data are normalized to the control group for the treated group (100%).

2.9. Measurement of Respiration of Mouse Cardiomyocyte. Respiration was assayed in freshly isolated mitochondria with a high-throughput-automated 96-well extracellular flux analyzer (XF96; Seahorse Bioscience, USA) as previously described [26].

2.10. Assessment of Mitochondrial Cytochrome c Release and ATP Content. To evaluate the subcellular localization of cytochrome c, we used confocal imaging of cells double labeled with Mitotraker Red CMX Ros (Molecular Probes, USA) and cytochrome c antibody (Cell Signaling Technology, USA). The ATP content of the myocardium was measured using an ATP bioluminescent assay kit (Beyotime, China).

2.11. Detection of Reactive Oxygen Species (ROS) by Mitochondria. ROS production by the mitochondria was assessed by using Mito SOX Red (Invitrogen, USA) according to product protocol. Briefly, slides were labeled with Mito SOX Red (300 nM) and Mito Tracker green (100 nM) for 20 min at 37°C, then observed by confocal microscopy (FV1000, Olympus) with the excitation wavelength at 488 and 543 nm.

2.12. Western Blot Assay. Cardiac tissues from the area at risk (AAR) were lysed, sonicated, and centrifuged. Western blotting was performed following standard protocol. The following primary antibodies were used: anti-phospho-FoxO3a (Thr32) (rabbit polyclonal IgG, Abcam, 1:100); anti-phospho-FoxO3a (Ser413) (rabbit polyclonal IgG, Cell Signaling Technology, 1:1000); anti-FoxO3a (rabbit polyclonal IgG, Cell Signaling Technology, 1:1000); anti-phospho-Akt (Ser473) (rabbit polyclonal IgG, Cell Signaling Technology, 1:1000); anti-Akt (rabbit polyclonal IgG, Cell Signaling Technology, 1:1000); anti-AMPK (rabbit polyclonal IgG, Cell Signaling Technology, 1:1000); anti-phospho-AMPK (Thr172) (rabbit polyclonal IgG, Cell Signaling Technology, 1:1000); anti-caspase-3 (rabbit polyclonal IgG, Merck Millipore, 1:200); anti-cleaved caspase-3 (rabbit polyclonal IgG, Merck Millipore, 1:200); anti-Bim (rabbit polyclonal IgG, Abcam, 1:1000); anti-Bax (rabbit polyclonal IgG, Abcam, 1:1000); anti-Bcl-2 (rabbit monoclonal IgG, Abcam, 1:500); anti-Bad (rabbit polyclonal IgG, Cell Signaling Technology, 1:1000); anti-phospho-Bad (rabbit polyclonal IgG, Cell Signaling Technology, 1:1000); anti-MnSOD (rabbit polyclonal IgG, Abcam, 1:5000); anti-Catalase (rabbit polyclonal IgG, Abcam, 1:2000); anti-Sirt1 (rabbit polyclonal IgG, Abcam, 1:1000); anti-PGC-1α (rabbit polyclonal IgG, Abcam, 1:1000); anti-phospho-PGC-1α (Thr563) (rabbit polyclonal IgG, Abcam, 1:2000); anti-α-Syn (rabbit polyclonal IgG, Abcam, 1:1000); anti-Therm (rabbit polyclonal IgG, Abcam, 1:2000); and β-actin (rabbit polyclonal IgG, Santa Cruz Biotechnology, 1:1000). The details are described previously [25].

2.13. Statistics Analysis. All data were expressed as mean ± SD and were analyzed using ANOVA, followed by a Bonferroni correction for post hoc test (with exception of Western blot). Western blot densities were analyzed with the Kruskal-Wallis test, followed by Dunn’s post hoc test. A value of P < 0.05 was considered to be statistically significant. All statistical tests were performed using SPSS software package version 14.0 (SPSS, Chicago, IL, USA).

3. Results

3.1. Coagonist Treatment Improved Cardiac Function of db/db Mice after Myocardial I/R Injury. LVEF and LVFS were enhanced significantly in the glucagon or ZP131 group compared with the db/db + I/R group. ZP2495 enhanced LVEF and LVFS more significantly as compared with that of the glucagon or ZP131 group (Figures 1(a)–1(c)). Maximal
velocity of pressure development and decline ($\pm$LV $dP/dt$ max) also revealed a significant improvement in response to glucagon or ZP131 pretreatment in diabetic mice which underwent cardiac I/R injury. The $\pm$LV $dP/dt$ max was significantly higher in the ZP2495 group compared with the glucagon or ZP131 group (Figures 1(d) and 1(e)). Moreover, administration of glucagon or ZP131 reduced infarct size after cardiac I/R injury in db/db mice. The mice treated with ZP2495 exhibited much greater decline of infarct size as compared with the glucagon or ZP131 group. There was no significant difference in the area at risk (AAR) between groups (Figures 1(h)–1(j)).

3.2. Regulation of Cardiac Glucose Metabolism Prevented from Cardiac I/R Injury in db/db Mice. The mice treated with glucagon or ZP131 reduced infarct size as compared with the glucagon or ZP131 group. There was no significant difference in the area at risk (AAR) between groups (Figures 1(h)–1(j)).

3.3. Coagonist Exerted Antiapoptotic Effect on Cardiomyocyte in db/db Mice after Cardiac I/R Injury. Apoptotic cardiomyocytes labeled by TUNEL positivity were more frequently observed in the db/db + I/R group, glucagon group, and ZP131 group compared with the ZP2495 group (Figure 2(a)). Quantitative analyses demonstrated that the percentage of TUNEL-positive cardiomyocytes was significantly less in the ZP2495 group compared with the db/db + I/R group or the glucagon group. However, there was no significant difference between the percentages of TUNEL-positive cardiomyocytes in the ZP2495 group and ZP131 group (Figure 2(b)). Consistent with its antiapoptotic effect, ZP2495 attenuated the expression of the cleaved, activated forms of caspase-3 (Figures 2(c) and 2(d)).

3.4. Antiapoptotic Effect of ZP2495 on High-Glucose-Induced NRVMs under SI/R Injury. To analyze the antiapoptotic effect of ZP2495 on high-glucose-induced NRVMs after SI/R injury, we performed flow cytometry, TUNEL, and caspase-3 activity assays. Representative flow cytometry results (Figure 3(a)) indicated that high-glucose and SI/R injury significantly increased the percentage of apoptotic NRVMs. ZP2495 treatment decreased high-glucose and SI/R-induced apoptosis of NRVMs. As shown in Figures 3(b)–3(e), the antiapoptotic effect of ZP2495 on NRVMs was also evidenced by increased TUNEL (green) staining of cells (Figures 3(b)–3(d)) and upregulation of caspase-3 enzymatic activities (Figure 3(e)).

3.5. Antioxidation Effect of ZP2495 on Cardiomyocyte in db/db Mice under Cardiac I/R Injury via a Akt/FoxO3a-Dependent Mechanism. To further understand the mechanism of the protective effect of ZP2495, we examined the expressions of Akt, FoxO3a, and apoptosis-related proteins using Western blot (Figure 4). ZP2495 administration increased the levels of phospho-Akt in myocardial tissue of db/db mice after cardiac I/R injury (Figures 4(a) and 4(b)). Consistently, our data further revealed reduction of the Akt-engaged phosphorylation of FoxO3 at Thr32 following ZP2495 treatment. Diabetes with cardiac I/R injury decreased FoxO3a phosphorylation at Thr32 (Figures 4(a) and 4(d)). To further examine FoxO3a activation, we measured the expression levels of its apoptosis-related downstream targets, including proapoptotic proteins BimEL and Bax, as well as antiapoptotic protein Bcl-2. The representative Western blot results demonstrated that diabetes with cardiac I/R injury increased the expressions of BimEL (Figure 4(e)) and Bax and decreased the levels of Bcl-2. ZP2495 treatment balanced the expression between pro- and antiapoptotic proteins (Figures 4(a) and 4(g)). As demonstrated in Figures 4(a) and 4(f), diabetes and I/R injury could moderately induce myocardial mitochondrial antioxidative enzyme expressions including MnSOD and catalase. ZP2495 upregulated the expression of these enzymes.

3.6. Coagonist Treatment Improved Mitochondrial Ultrastructural Morphology in Myocardium of db/db Mice Subjected to Cardiac I/R Injury. Alterations in mitochondrial ultrastructure were observed by TEM. As shown in Figure 5(a), in the db/db + I/R group, most of the mitochondria from the peri-infarct zone presented significant disorders, including abnormal cristae or areas of the matrix. In some mitochondria, the cristae and matrix were cleared out, resulting in vacuoles; some mitochondria were swelling and presented with crista disorientation and breakage. The derangement in the ultrastructural morphology of mitochondria was improved in the ZP2495 treatment group, as evidenced by normalized cristae density and architecture. Most of the mitochondria in this group presented sharply defined cristae. The number of swelling mitochondria was less, and no obvious vacuoles could be found in the ZP2495 treatment group.

3.7. Coagonist Treatment Improved Mitochondrial Energetic Transition in Myocardium of db/db Mice Subjected to Cardiac I/R Injury. We measured mitochondrial respiration, coupling of oxidative phosphorylation (OxPhos), and the relative ATP level in isolated mitochondria from hearts of all groups. Mitochondrial respiration was first quantified in the presence of substrates of complex I, II, and IV and the coupling of OxPhos under each condition (Figures 5(b)–5(d)). The basic bioenergetic behavior of mitochondria from hearts of the ZP2495 treatment group was noticeably improved. The RCR (state 3/4), a measure of the degree of coupling of OxPhos, was markedly decreased at the level of the three main respiratory complexes in mitochondria of the db/db and db/db + I/R groups (Figure 5(e)). In these two groups of mitochondria, a trend to higher values of state 4 respiration was evident in complex I, II, and IV (Figures 5(b)–5(d)). Likewise, the relative ATP levels in the db/db and db/db + I/R groups, the mycardia were also significantly decreased, and ZP2495 treatment preserved the reduction (Figure 5(f)).
Figure 1: Effects of ZP2495, glucagon, and ZP131 on cardiac function and hemodynamic and cardiac glucose metabolism in db/db mice after I/R injury. (a) Representative M-mode echocardiography images. (b) Measurement of left ventricular ejection fraction (LVEF). (c) Measurement of left ventricular fractional shortening (LVFS). (d, e) Measurement of maximal velocity of pressure development and decline ($\pm$LVdp/dt). (f) Representative PET/CT scan images from each group. Higher glucose uptake level is evidenced by an increase in the intensity of red color. (g) Quantification of accumulated $^{18}$F-fluorodeoxyglucose ($^{18}$F-FDG) in the heart. Relative accumulation of the radioactivity in particular regions of interest was expressed as standardized uptake value (SUV). (h) Myocardial infarct size was assessed by Evans blue/2,3,5-triphenyl-2H-tetrazolium chloride (TTC) double staining. Evans blue stained areas (black) indicated nonischemic/reperfused area. TTC-stained areas (red staining) indicated ischemic but viable tissue. Evans blue/TTC staining negative areas indicated infarcted myocardium. (i) Summary of infarct area (INF) per area at risk (AAR). (j) Summary of AAR per left ventricle (LV). Sham: BKS + sham group; db/db: db/db + sham group; I/R: db/db + I/R group; ZP2495: db/db + I/R + ZP2495 group; glucagon: db/db + I/R + glucagon group; and ZP131: db/db + I/R + ZP131 group. Presented values are mean ± SEM. *P < 0.05 versus the sham group; $^\#$P < 0.05 versus the db/db group; $^\&$P < 0.05 versus the db/db + I/R group; and £P < 0.05 versus the db/db + I/R + ZP2495 group.
3.8. Coagonist Treatment Prevented Mitochondrial Depolarization and Reduced Cytochrome c Release and Mitochondria ROS Generation in High-Glucose-Induced NRVMs after SI/R Injury. As is shown in Figure 6(a), there were stronger red fluorescent signals and weaker green fluorescent signals in the control group, indicating intact high mitochondria membrane potential. However, red fluorescent signals had almost disappeared and green fluorescent signals were greatly enhanced in the HG (high-glucose) and HG + SI/R group, suggesting typical mitochondrial membrane potential collapse. While in the ZP2495 treated group, there were more red fluorescent signals remaining, along with fewer green fluorescent signals, which indicated that ZP2495 treatment prevented mitochondrial depolarization (Figure 6(b)). In the control group, cytochrome c immunoreactivity was colocalized with mitotracker red fluorescence, indicating that cytochrome c was confined to the mitochondria. After high-glucose and SI/R injury, cytochrome c immunoreactivity was diffusely distributed throughout the cytoplasm and an increase in mitochondria-associated cytochrome c, indicating that cytochrome c was released from mitochondria to the cytoplasm. On the contrary, fewer cells exhibited such cytochrome c release pattern in the ZP2495 treatment group (Figure 6(c)). High glucose with SI/R injury-induced mitochondrial ROS generation was assessed by MitoSox fluorescence (Figure 6(d)). Mitochondria were visualized with Mitotracker. Image merging of MitoSox with Mitotracker showed that the diminished ROS was generated from mitochondria in the ZP2495 treatment group.

3.9. Coagonist Increased Mitochondrial Biogenesis in Hearts of db/db Mice Subjected to I/R Injury through AMPK/FoxO3a and AMPK-SIRT1-PGC-1α Pathway. To investigate
the potential signaling pathways involved in which ZP2495 improved cardiomyocyte energy metabolism after diabetes with myocardial I/R injury, we examined the level of AMPK. Our results indicated that I/R injury markedly decreased phosphorylation of AMPK in db/db mice, which was abrogated by ZP2495 (Figures 7(a) and 7(b)). Furthermore, we found reduction of AMPK-activated Foxo3 phosphorylation at the Ser413 site following ZP2495 treatment (Figures 7(a) and 7(c)). As shown in Figures 7(a)–7(e), AMPK-α phosphorylation and SIRT1 expression level was dramatically lower and PGC-1α acetylation was increased in the db/db and the db/db + I/R groups. However, AMPK-α and PGC-1α protein levels of cardiac tissue did not display a measurable difference between groups. ZP2495 treatment stimulated AMPK-α phosphorylation and PGC-1α deacetylation without altering PGC-1α protein expression (Figures 7(a) and 7(e)). In the meanwhile, ZP2495 treatment significantly increased SIRT1 expression (Figures 7(a) and 7(d)). We next determined whether administration of ZP2495 could increase mitochondrial biogenesis in hearts of db/db mice subjected to I/R. ZP2495 treatment also significantly restored the protein levels of Nrf-1 and Tfam (Figures 7(a) and 7(f)). The effects of ZP2495 on mitochondrial biogenesis in db/db mice subjected to cardiac I/R injury were related to the AMPK-SIRT1-PGC-1α pathway (Figure 8).

4. Discussion

It is increasingly recognized that the occurrence of coronary artery disease contributes to an increased risk for the development of heart failure in diabetic patients. Although the pathophysiological mechanisms are certainly multifactorial,
Figure 4: ZP2495 regulated Akt, FoxO3a, and apoptosis-related protein expression. (a) Representative blots of p-Akt, Akt, p-Bad, Bad, p-FoxO3a, FoxO3a, BimEL, MnSOD, catalase, Bax, and Bcl-2 in cardiomyocyte of db/db mice subjected to I/R. (b–g) Semiquantitative analysis of the Western blots. Sham: BKS + sham group; db/db: db/db + sham group; I/R: db/db + I/R group; ZP2495: db/db + I/R + ZP2495 group; glucagon: db/db + I/R + glucagon group; and ZP131: db/db + I/R + ZP131 group. Presented values are mean ± SEM. *P < 0.05 versus the sham group; #P < 0.05 versus the db/db group; $P < 0.05 versus the db/db + I/R group; and £P < 0.05 versus the db/db + I/R + ZP2495 group.
emerging evidence suggests that derangements in cardiac energy metabolism play a fundamental role in the pathogenesis of diabetic cardiomyopathy. Recent studies focusing on the mitochondria show an important role of abnormal mitochondrial function [27]. Previous studies revealed that glucagon-GLP-1 dual-agonist ZP2495 was superior compared with glucagon alone in the improvement of cardiac function in subjects with insulin resistance (IR) [28]. The data by Axelsen et al. have presented that perfusion buffer containing ZP131 failed to change cardiac function in hearts from either control or IR rats compared with vehicle-perfused hearts. In contrast, glucagon and ZP2495 perfusion improved cardiac function in control hearts. By comparison, in hearts from IR rats, however, only ZP2495 significantly increased cardiac power. Although their findings did not elucidate the protective effect of ZP2495 in the presence of diabetes, these evidences might indicate the protective role of ZP2495 in the diabetic heart because insulin resistance was usually conceived as the earlier stage of diabetes. Furthermore, ZP2495 increased glucose oxidation and glycolytic rates in IR hearts as glucagon did avoid the concomitant accumulation of AMP or ADP [28]. In line with these findings, in our current study, dual-agonist ZP2495 induced superior cardioprotection (decreased infarct size, enhanced cardiac function, and ameliorated cardiac apoptosis) in diabetic hearts subjected to myocardial I/R compared with that of ZP131 or glucagon, which might be ascribed to its stronger capacity to upregulate cardiac glucose metabolism and mitochondrial function as compared to ZP131 or glucagon. Taken together, both our studies depicted the superiority of dual-agonist ZP2495 compared with ZP131 or glucagon in the treatment of cardiometabolic diseases with cardiac glucose metabolism defects.
The present study demonstrates that the metabolic effects observed in different regions of the ischemia-injured heart may in large part contribute to the cardioprotection observed with GLP-1 treatment [29]. However, our current study failed to analyze metabolic changes triggered by treatments in terms of ischemic versus remote regions. These changes could provide more information about a metabolic shift in viable regions of the hearts subjected to I/R injury, and we will explore this operation in our future experiments.

Myocardial I/R injury can induce additional injury to the myocardium, due to excessive oxygen free radicals, calcium overload, neutrophil infiltration, depletion of energy stores, and changes in subcellular activities including the opening of the mitochondrial permeability transition pore (MPTP) [30]. All of these changes are detrimental to ischemic cardiomyocyte and may reduce the beneficial effects of reperfusion [31]. Notably, the mechanisms activating during ischemia may lead to necrotic cell death in cardiomyocytes, while hallmarks of apoptosis mainly occur after reperfusion [32]. Accordingly, both necrotic and apoptotic cardiomyocyte cell deaths contribute to the final myocardial infarction (MI) size. Enormous preclinical and clinical studies have demonstrated that diabetes will increase the vulnerability of diabetic hearts subjected to ischemic injury and result in higher mortality [33]. Our results from the present study exhibited that ZP2495 significantly reduced infarct size after cardiac I/R injury in db/db mice, mainly by exerting an antiapoptotic effect on cardiomyocyte both in vivo and in vitro.

Cardiac mitochondria are vital organelles that supply energy to support the high ATP consumption of a beating heart [34]. The irreversible mitochondrial dysfunction is a crucial event in cardiomyocyte death during myocardial I/R.
Figure 7: ZP2495 increased mitochondrial biogenesis in hearts of db/db mice subjected to I/R. (a) Representative blots of p-AMPK, AMPK, p-FoxO3a, FoxO3a, Sirt1, Ace-lysine-PGC-1α, PGC-1α, Nrf-1, and Tfam in cardiomyocyte of db/db mice subjected to I/R. (b–e) Semiquantitative analysis of the Western blots. Sham: BKS + sham group; db/db: db/db + sham group; I/R: db/db + I/R group; ZP2495: db/db + I/R + ZP2495 group; glucagon: db/db + I/R + glucagon group; ZP131: db/db + I/R + ZP131 group. Presented values are mean ± SEM. *P < 0.05 versus the sham group; †P < 0.05 versus the db/db group; ‡P < 0.05 versus the db/db + I/R group; and §P < 0.05 versus the db/db + I/R + ZP2495 group. Data shown are representative of six independent experiments.
injury. After reperfusion, intracellular Ca\(^{2+}\) elevation triggers mPTP opening, leading to the depletion of ATP and loss of mitochondrial integrity [35, 36]. Therefore, alleviating mitochondrial dysfunction may decrease cardiac I/R injury.

Meanwhile, modulating mitochondrial function may improve insulin resistance and reduce subsequent cardiac mortality. In addition, many studies have documented signs of mitochondrial dysfunction, including impaired respiration, damaged oxidative phosphorylation, altered substrate utilization, decreased ATP production, and impaired ATP transfer [37]. In the present study, we observed both structural and functional mitochondrial impairment in the myocardium of db/db mice subjected to cardiac I/R injury. ZP2495 treatment improved mitochondrial ultrastructural morphology. These findings were in line with the alleviated functional damage observed in the mitochondria in ZP2495-treated mice.

A large accumulation of myocardial mitochondrial ROS was observed in both permanently occluded and reperfused myocardia, especially in diabetic animals, suggesting that oxidative stress in mitochondrial may contribute to cardiac dysfunction [4]. It would result in electron transport chain deterioration, mitochondrial membrane depolarization, apoptotic pathway activation, and cardiomyocyte death [4, 38]. In this study, administration of ZP2495 reduced high-glucose and SI/R injury-induced mitochondrial ROS generation, which may be responsible for preventing mitochondrial depolarization and attenuating the release of cytochrome c. These observations provide strong evidence for the notion that reduction in cardiac mitochondrial injury may represent a critical event in the cardioprotective effect of ZP2495.

Several studies have found that PGC-1α is important for controlling metabolic pathways in the heart during development as well as in response to physiological stressors and pathological stimuli [39, 40]. PGC-1α is a positive regulator of mitochondrial biogenesis and respiration and plays roles in neurodegenerative disorders and heart failure and pathological conditions that are strongly associated with mitochondrial defects [39]. Recent studies have shown that PGC-1α acetylation was a kind of protein modification with inactivation effects [41]. PGC-1α activity has also been reported to be modulated by several modifications, such as phosphorylation by AMPK or deacetylation by AMPK and sirtuin 1 (SIRT1). In the current study, we observed that ZP2495 upregulated PGC-1α and its downstream Nrf-1 and Tfam expressions, which was associated with an increase in mitochondrial DNA copies in db/db mouse hearts. These results indicated that ZP2495 may exhibit its effects of cardioprotection through improving mitochondrial biogenesis by activating the AMPK/PGC-1α pathway.

Another important observation of our study is that FoxO3a may play an important role in protecting the myocardium from injury. The FoxO transcription factors, including FoxO1, FoxO3a, FoxO4, and FoxO6, belong to the

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**Figure 8**: Proposed scheme for the protective effect of ZP2495 on cardiomyocyte. A summary of our current hypothesis about the mechanisms that the glucagon and GLP-1 dual-agonist (ZP2495) protects against myocardial I/R injury in db/db diabetic mice. The protective effects of ZP2495 may be associated with the activation of Akt/FoxO3 and AMPK/FoxO3a signaling pathways.
forkhead family of transcriptional regulators. FoxO transcription factors have been implicated in regulating diverse cellular functions such as differentiation, metabolism, proliferation, and apoptosis [42, 43], especially for Akt-mediated antiapoptotic properties [23]. Furthermore, FoxO3a has been considered a converging point for Akt and AMPK signaling pathways [44, 45]. AMPK phosphorylation of FoxO3a at the site of Ser413 enhances the stress resistance and longevity [46–50]. On the other hand, Akt phosphorylation of FoxO3a at Thr32 site inhibits FoxO3a transcriptional activity, promoting antiapoptosis and insulin sensitivity [51]. Therefore, Akt and AMPK signaling may elicit transcriptional regulation via FoxO3a to allow organismal adaptation for physiological or pathophysiological changes. The convergence of the two pathways at the level of FoxO3a may play a critical role in the crosstalk between Akt and AMPK. Our data revealed that the protective effect of ZP2495 was related to FoxO3a activation mediated by the PI3K/Akt and AMPK pathways. We found that ZP2495 phosphorylated FoxO3a to inhibit the transcription of proapoptotic gene Bim, which resulted in decreased expression of proapoptotic proteins such as Bim and Bax. Consequently, cytochrome c was released into the cytosol and the caspase cascade was activated. In parallel, ZP2495 increased levels of the antiapoptotic proteins Bad and Bcl-2, which alleviated the apoptosis induced by diabetes and cardiac I/R injury. Collectively, we demonstrated that ZP2495 exerted an antiapoptotic effect by inhibiting ROS production, cytochrome c release, and caspase-3 activation through Akt/FoxO3a and AMPK/FoxO3a pathways. In the present study, myocardial I/R injury induced loss of phosphorylation of Akt and AMPK in db/db mice. These data suggested a potential role of Akt-FoxO3a and AMPK-FoxO3a in cardiac I/R injury and diabetes-induced mitochondrial dysfunction and apoptosis. Our data revealed a novel mechanism whereby ZP2495 exerted its protective effect by activating FoxO3a through the Akt and AMPK signaling pathways.

4.1. Limitations. Other members of the RISK pathway may also participate in the link between ZP2495 and its cardiac protection effects. In order to further demonstrate the efficacy of ZP2495 administration on cardiomyocyte mitochondrial biogenesis and function, AMPK-related pathways should be investigated in future studies. More studies should be performed to figure out whether ZP2495 activates Akt/FoxO3a and AMPK/FoxO3a pathways by different mechanisms in diabetic mice or nondiabetic mice which underwent cardiac I/R injury.

5. Conclusion

In conclusion, we provided evidences that glucagon and GLP-1 dual-agonist ZP2495 protected against myocardial I/R injury in db/db mice, which may be attributed to the improvement of cardiac mitochondrial function and energy metabolism. These protective effects of ZP2495 might be associated with Akt/FoxO3a and AMPK/FoxO3a signaling pathways.

Disclosure

This work was presented at The 25th Great Wall International Congress of Cardiology 2014. The corresponding meeting abstract (GW25-e4495) was also presented in Supplement of Journal of the American College of Cardiology (JACC, 21; 64(16):C44. DOI:10.1016/j.jacc.2014.06.213).

Conflicts of Interest

The authors have declared that no competing interest exists.

Acknowledgments

This work was supported by the National Key Research Project (2016YFA0100903), China National Funds for Distinguished Young Scientists (81325009), National Nature Science Foundation of China (nos. 81270168, 81300149, 81530058, and 81570272), Innovation Team Grant of Shanxi Province (no. 2014KCT-20) to Feng Cao (BWS12037), and Beijing Natural Science Foundation (7152131).

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