To investigate whether carbamylated erythropoietin can influence the coronary collateral circulation via extracellular signal regulated kinase 1/2 signaling pathway. A diabetic rat model was established. Vascular endothelial growth factor, angiotensin-I and phosphorylated-extracellular signal regulated kinase 1/2 expression were observed along with changes in hemodynamics and infarct area. The results showed that left ventricular diastolic pressure, left ventricular end diastolic pressure, maximum rate of left ventricular pressure rise, heart rate, (left ventricular systolic pressure-left ventricular diastolic pressure)* heart rate, microvessel density and vascular endothelial growth factor were significantly decreased and the percentage of myocardial infarct size, the levels of prostacyclin, endothelin 1, angiotensin II, von Willebrand factor, phosphorylated-fibroblast growth factor receptor 2, phosphorylated-phospholipase C gamma and phosphorylated-extracellular signal regulated kinase 1/2 were significantly increased in other groups compared with control group (p<0.05). Left ventricular diastolic pressure, left ventricular end diastolic pressure, maximum rate of left ventricular pressure rise, heart rate and (left ventricular systolic pressure-left ventricular diastolic pressure)* heart rate were significantly increased in carbamylated erythropoietin group and significantly decreased in carbamylated erythropoietin+PD98059 group compared with the sham group (p<0.05). However, the myocardial infarction size and the levels of prostacyclin, endothelin 1, angiotensin II and von Willebrand factor were significantly decreased and the microvessel density, vascular endothelial growth factor, angiotensin-1, phosphorylated-fibroblast growth factor receptor 2, phosphorylated-phospholipase C gamma and phosphorylated-extracellular signal regulated kinase 1/2 were significantly increased in carbamylated erythropoietin and carbamylated erythropoietin+PD98059 groups compared with the sham group (p<0.05). Amino glycosylated erythropoietin was able to improve the hemodynamics of diabetic infarct rats and there was also an extracellular signal regulated kinase 1/2 conduction pathway in the neovascularization of diabetic infarct rats. This provides a new target for future clinical use to reduce myocardial damage caused by myocardial infarction.

Key words: Erythropoietin, diabetes mellitus, heart attack, extracellular signal regulated kinase, streptozotocin
circulation and cause side effects such as hypertension and vascular embolism, so its clinical application has certain limitations[7].

Carbamylated Erythropoietin (CEPO) is a carbamoylated derivative of EPO. When the conformation and function of the protein are changed, CEPO with a molecular weight of about 40 kDa can be formed[8,9]. Although CEPO did not promote erythropoiesis, it retained a variety of cell protective functions of EPO[10,11]. CEPO can prevent the release of mitochondrial contents (such as reactive oxygen species and cytochrome c) by limiting the opening of mitochondrial permeability transition pore, thus reducing cell apoptosis[5].

In addition, CEPO can also affect cell signal transduction through Extracellular signal Regulated Kinase (ERK)-1/2 pathway[12]. However, it is not clear whether CEPO can promote coronary neovascularization in DHA rats. In this study, we investigated the effect of CEPO treatment on coronary neovascularization induced by Streptozotocin (STZ) fed on a high-fat/high carbohydrate diet in DHA rats and its underlying molecular mechanisms.

MATERIALS AND METHODS

Source of reagents:
RhePO, STZ, Diethyl Pyrocarbonate (DEPC), sodium borate and potassium cyanate were purchased from Sigma company (Sigma, CA, USA). The Terminal deoxynucleotidyl transferase mediated dUTP Nick End Labeling (TUNEL) kit was obtained from Shanghai Shanran Biotechnology Co., Ltd (Shanran Shanghai, China). Caspase-3, B-cell lymphoma extra-large (Bcl-xl), ERK-1 and ERK-2 antibodies were purchased from Cell Signaling Technology (CST, CA, USA). ERK messenger RNA (mRNA) was obtained from Shanghai Lianmai Bioengineering Co., Ltd (Lianmai Shanghai, China).

Synthesis of CEPO:
CEPO was synthesized as previously described by Khan, et al. In short, potassium cyanate is added to sodium borate containing rhEPO (1 mg/ml) and 500 μl (1 m) to produce a solution with a final concentration of 1 mol/l[13]. They were mixed and incubated at 37°C for 1 d to form the reaction solution with in situ hybridization kit. Excess potassium cyanate was removed from the reaction solution by dialysis and then further concentrated by ultrafiltration (cut-off membrane, 10 kDa). The protein concentration was measured by Coomassie brilliant blue staining and the absorbance was read at 335 nm.

Diabetic rat model:
A total of 100 healthy male Wistar rats from animal center of Zhongshan University were used to establish diabetic models. They were fed with high fat and high carbohydrate diet (66.6 % basic rat food, 20 % sucrose, 10 % lard, 3 % egg yolk and 0.4 % cholesterol) for 4 w and then intraperitoneally injected with STZ (50 mg/kg). 1 w later, the same dose of STZ was injected again. According to the standard of fasting blood glucose concentration >18 mmol/l and lasting for 1 w. Finally, 80 rats with diabetes were identified and used in the following experiments.

Animal grouping:
According to the principle of random distribution, the rats were randomly divided into four research groups: control group (only left coronary artery was threaded without ligation after thoracotomy); sham group, CEPO group and CEPO+PD98059 group; CEPO group, CEPO+PD98059 group; CEPO group, CEPO+PD98059 group, CEPO group, CEPO+PD98059 group, CEPO group. The drug was given once 30 min before ligation of left coronary artery and twice a week after operation (ear vein injection, 50 μg/kg each time); The administration method of CEPO in CEPO+PD98059 group was the same as that of CEPO group, but PD98059 (0.3 mg/kg by ear vein injection) was given 30 min before ligation of left coronary artery and twice a week (0.3 mg/kg each time) after operation. The control group was given the same dose of saline water by ear edge vein injection at the same time, 20 animals in each group.

Monitoring of hemodynamic parameters in rats:
After anesthetized rats by intraperitoneal injection of chloral hydrate, the right skin of the rat neck was cut, and the fascia, muscle and carotid sheath were separated layer by layer and the common carotid artery was separated by 1-1.2 cm. The common carotid artery was cut by ophthalmic scissors and the catheter was inserted about 1-1.5 cm. After 15 min, Blood Pressure (BP) and mean arterial pressure (map) were recorded. Then, Med lab-u/8C biological signal acquisition and processing system was applied to monitor the Left Ventricular Diastolic Pressure (LVDP), maximum rate of left ventricular pressure rise (dP/dt max), Left Ventricular End Diastolic Pressure (LVEDP).

Measurement of infarct size:
The infarct size was measured by
2,3,5-Triphenyltetrazolium chloride (TTC) staining. The left ventricle was rinsed with normal saline and wrapped with tin foil and stored in a refrigerator at 20°C. 2-3 mm serial sections were made from the apex to the bottom of the heart in the direction perpendicular to the long axis of the heart to the level of the ligation line. The slices were placed in TTC dye solution and dyed in 37°C incubator for 20 min and then put into 4% formaldehyde solution for 15 min. White infarcted myocardial tissue and brick red viable myocardial tissue could be observed after taking out the tissue. Finally, the infarct area and non-infarct area were separated with scissors and the mass of infarct area was calculated by electronic balance.

The infarct area=left ventricular infarct mass/left ventricular ligation line myocardial mass×100%.

Enzyme-Linked Immunosorbent Assay (ELISA):

The antigens (prostacyclin (PGI2), Endothelin 1 (ET-1), angiotensin II (Ang II), von Willebrand Factor (vWF)) were diluted to 10 μg/ml with coated buffer and overnight at 4°C. The next day, add 0.1 ml serum of rats in each group into the coated reaction pore and incubate at 37°C for 1 h; 0.1 ml fresh diluted enzyme labeled second antibody was added and incubated at 37°C for 30 min; then 0.1 ml tetramethylbenzidine (TMB) substrate solution was added and incubated at 37°C for 10 min. Finally, add 0.05 ml 2 mol/l sulfuric acid to determine the Optical Density (OD value) at 450 nm wave length of ELISA detector. Meanwhile, the OD values of PGI2, ET-1, Ang II and vWF were determined and calculated in each group.

The Microvessel Density (MVD) was measured by immunohistochemistry:

In this study, vascular endothelium was labeled with CD34 and MVD was counted by immunohistochemistry. The infarct area was determined according to the existing studies[14]. 5 μm thick myocardial tissue was reserved at the junction of infarct area (gray white, wall thinning) and non-infarct area (pink). The frozen section (2 mm below the ligation line) was made in the sham operation group. The myocardium of each group was fixed, transparent, waxed, embedded, dewaxed and hydrated. After incubation overnight at 4°C, each slice was incubated with 50 μl of ready to use maxvision reagent at room temperature for 15 min and then 100 μl of freshly prepared Diaminobenzidine (DAB) solution was added. Under the microscope, the number of endothelial cells or clusters of endothelial cells stained brown was counted as a blood vessel and the vessels in the lumen with more than 8 red blood cells and thicker muscular layer were not counted. Under the microscope of 400x visual field, 5 visual fields were randomly selected from each section to count the number of microvessels (MVC) in the infarct border area and the average value was obtained. MVC in each field was MVD.

The expression of Vascular Endothelial Growth Factor (VEGF) and Ang-1 was detected by immunohistochemistry:

Paraffin sections of myocardial tissue were routinely dewaxed, hydrated and high-pressure antigen repair, then diluted primary antibody (rabbit anti-VEGF 1:50, rabbit anti-rat Ang-1 antibody 1:100) was dropped at 37°C for 2 h and washed with Phosphate Buffered Saline (PBS) three times. After that, the second antibody was incubated at room temperature for 15 min, washed with PBS for 3 times then dyed with DAB developer and hematoxylin. Taking the presence of brown particles in the cytoplasm or extracellular matrix of cardiac myocytes as the positive criterion, six visual fields were randomly selected under high power microscope (400 times) and their Integral Absorbance (IA) was counted respectively by automatic image analysis system and the numerical mean value was analyzed.

Western blot analysis:

After adding 30 mg fresh myocardial tissue into 100 μl tissue lysate for 1 h, the γ supernatant was centrifuged and the protein was quantified by Bicinchoninic acid (BCA) protein concentration quantitative kit. After Sodium Dodecyl-Sulfate Polyacrylamide Gel Electrophoresis (SDS-PAGE) electrophoresis, 50 μg protein/well was transferred to Polyvinylidene Fluoride (PVDF) membrane. 5% bovine serum albumin (BSA) was sealed at room temperature for 2 h. The working solutions of Fibroblast Growth Factor Receptors (FGFR), phosphorylated-fibroblast growth factor receptors (p-FGFR2), phosphorylated-phospholipase C gamma (p-PLCγ), phosphorylated ERK1/2 (p-ERK1/2), FGFR2, PLC, ERK1/2 and β-actin were prepared and sealed overnight at 4°C and that was washed three times at room temperature. Then incubated with Horseradish Peroxidase (HRP) labeled second antibody working solution for 2 h. The Electrochemiluminescence (ECL) working fluid was exposed to the high performance luminescent gel imaging system. The gray scale of the strips was calculated by image J software and the relative expression intensity of phosphoprotein was detected by the ratio of phosphorylated protein to total protein.
**Statistical analysis:**

Statistical Package For The Social Sciences (SPSS) 22.0 software system was used in present study and expressed as data x±s. Analysis of Variance (ANOVA) was used to compare the data more than three groups and t test was used for pairwise comparison between two groups. p<0.05 was considered to be statistically significant.

**RESULTS AND DISCUSSION**

Compared with the control group, LVDP, LVEDP, ±dP/dt\(_{\text{max}}\), heart rate (HR) and (LVSP-LVDP)×HR were significantly lower in sham, CEPO and CEPO+PD98059 groups (p<0.05). Compared with the sham group, the LVDP, LVEDP, ±dP/dt\(_{\text{max}}\), HR and (LVSP-LVDP)×HR were significantly higher in CEPO group (p<0.05) and significantly lower in CEPO group, as shown in fig. 1.

Compared with the control group, the percentage of myocardial infarct size in sham, CEPO and CEPO+PD98059 groups were significantly higher (p<0.05). However, the myocardial infarction size in CEPO and CEPO+PD98059 groups were significantly lower than that in sham group (p<0.05). In addition, compared with the control group, the MVD in sham, CEPO and CEPO+PD98059 groups were significantly lower (p<0.05) and the MVD in CEPO and CEPO+PD98059 groups were significantly higher than that in sham group (p<0.05). As shown in fig. 2.

Compared with the control group, the levels of PGI2, ET-1, Ang II and vWF in sham, CEPO and CEPO+PD98059 groups were significantly increased (p<0.05). Compared with the sham group, the levels of PGI2, ET-1, Ang II, vWF in CEPO and CEPO+PD98059 groups were significantly decreased (p<0.05), suggesting that CEPO can improve the coronary microcirculation function of diabetic rats. The detection results of PGI2, ET-1, Ang II, vWF levels in peripheral serum of rats in each group are shown in fig. 3.

Compared with the control group, the expression of VEGF and Ang-1 in sham, CEPO and CEPO+PD98059 groups were significantly decreased (p<0.05). Compared with the sham group, the expression of VEGF and Ang-1 were significantly increased in CEPO and CEPO+PD98059 groups (p<0.05). The detection results of VEGF and Ang-1 expression in muscle tissue of rats in each group are shown in fig. 4.

Compared with the control group, the expression of p-FGFR2, p-PLCγ, p-ERK1/2 proteins in CEPO and CEPO+PD98059 groups were significantly increased (p<0.05), but there is no significantly difference in sham group (p>0.05). Compared with the sham group, the expression of p-FGFR2, p-PLCγ, p-ERK1/2 proteins were further increased in CEPO and CEPO+PD98059 groups (p<0.05). As shown in fig. 5.

**Fig. 1: Hemodynamic parameters of rats in four groups**

Note: *p<0.05, compared with the control group; ’p<0.05, compared with the sham group. (■) Control; (□) Sham; (▲) CEPO; (▲) CEPO+PD98059
Fig. 2: Myocardial infarct size and MVD of rats in four groups
Note: *p<0.05, compared with the control group; †p<0.05, compared with the sham group

Fig. 3: Changes of microcirculation markers in four groups
Note: *p<0.05, compared with the control group; †p<0.05, compared with the sham group. (□) Control; (△) Sham; (■) CEPO; (□□) CEPO+PD98059

Fig. 4: The expression of VEGF and Ang-1 in four groups
Note: *p<0.05, compared with the control group; †p<0.05, compared with the sham group. (□) Control; (△) Sham; (■) CEPO; (□□) CEPO+PD98059

Fig. 5: The protein expression levels of p-FGFR2, p-PLCγ, p-ERK1/2 in four groups
Note: *p<0.05, compared with the control group; †p<0.05, compared with the sham group. (□) Control; (△) Sham; (■) CEPO; (□□) CEPO+PD98059
Diabetes, as one of the major diseases threatening human health, its complications are an important factor to reduce the prognosis of patients, among which cardiovascular and cerebrovascular complications are the most common cause of death in patients with diabetes.[14]. Although oral hypoglycemic drugs and insulin can greatly alleviate the patient's condition, there are still many side effects. Therefore, it is of great significance to formulate effective strategies for the prevention and treatment of diabetes. Diabetic cardiomyopathy is mainly caused by high blood glucose and coronary microcirculation hypoperfusion. In view of the fact that myocardial activity largely depends on the supply of microcirculation oxygen, nutrients and the transport of metabolites, restoring coronary microcirculation is an important way to improve myocardial activity.[15-16]

In this study, we used Med Lab-U/8C, a biological signal acquisition and processing system, to directly monitor the physical indicators of cardiac work and evaluate the real-time cardiac function. LVDP and the maximum rate of rise of left ventricular pressure (+dP/dt max) are important indicators to reflect cardiac systolic function. The decrease of LVDP and +dP/dt_max indicates the decrease of myocardial systolic function. In addition, the LVEDP and the maximum rate of decrease of left ventricular pressure (-dP/dt_min) are important indicators to reflect cardiac diastolic function. The increase of LVEDP or the decrease of -dP/dt_min indicates the decrease of cardiac diastolic function; (Left Ventricular Systolic Pressure-Left Ventricular Diastolic Pressure)×heart rate (LVSP-LVDP)×HR reflects the contractile force of myocardium. The results showed that after ligation of left anterior descending coronary artery, the systolic and diastolic functions of the heart decreased to varying degrees, which was reflected in the decrease of left ventricular development pressure and the maximum rise rate of left ventricular pressure, while the end diastolic pressure increased and the maximum decline rate of left ventricular pressure decreased, which was consistent with the previous research results.[17]. Pretreatment and postconditioning with ecpo can significantly improve LVDP, LVEDP, +dP/dt_max and other indicators of the heart, that is, significantly improve the cardiac systolic and diastolic function, which is similar to the findings of Apstein.

ERK are a class of serine/threonine protein kinases, whose main function is to transmit mitogen signal transduction proteins.[18,19] ERK is the key to transmit signal from surface receptor to nucleus. Phosphorylated ERK 1/2 is transferred from cytoplasm to nucleus, which mediates transcription activation of various cytokines, regulates cell proliferation and differentiation, cell morphology maintenance, cytoskeleton construction, cell apoptosis and cell malignant transformation and other biological processes. Hyperglycemia cannot only cause myocardial injury through the formation of advanced glycation end products, but also cause a variety of vascular complications by stimulating the proliferation of vascular smooth muscle.[20], which can be attenuated by the activation of ERK signaling pathway. In diabetic rats with myocardial infarction, the activation of ERK cell signaling pathway is closely related to myocardial cell anti apoptosis.[21]. The effect of hyperglycemia on ERK phosphorylation in cardiomyocytes is also time-dependent and short-term hyperglycemia can enhance ERK phosphorylation.[22,23]. Recent studies have shown that CEPO can activate ERK cell signaling pathway to protect myocardium from ischemia/reperfusion injury.[24,25]. In chronic hypoxia, CEPO may reduce the levels of PGII2, ET-1, Ang II, vWF in peripheral serum to near normal level through ERK signal pathway.[10]. In this study, we demonstrated that CEPO could protect myocardium against myocardial fibrosis and apoptosis by activating p-ERK signaling pathway.

Myocardial infarction can lead to hypoxia in body tissues and cause changes in local cytokine levels, induce the proliferation, migration, lumen formation and formation of new blood vessels of vascular endothelial cells with existing vessels, so as to adaptively increase blood supply and oxygen supply. During this process, abnormal secretion of cytokines, especially angiogenesis factor VEGF and basic fibroblast growth factor (bFGF), occurred. bFGF, a basic protein, has the effect of promoting cell mitosis and promoting angiogenesis. After binding with its specific receptor FGFR, bFGF can activate tyrosine kinase in vascular endothelial cells, thus activating PLCγ.[6]. PLCγ is the most important member of the four phosphoinositol specific phospholipase C (PLC β, PLC γ, PLC δ and PLC ε), which are widely expressed in vascular endothelial cells. Phosphorylation is the key factor to regulate PLCγ activity. In the process of bFGF and other factors activating vascular endothelial cells, PLCγ can form receptor complex with FGFR2, further activate ERK1/2 signaling pathway protein downstream, start endothelial cell proliferation and generate neovascular endothelial cells. In this study, we found that CEPO can significantly promote bFGF induced phosphorylation of receptor and downstream
cell signaling pathway proteins, which may be one of the mechanisms of CEPO promoting angiogenesis.

In conclusion, carbamoyl erythropoietin can improve the hemodynamics of diabetic myocardial infarction rats and regulate ERK 1/2 pathway in promoting neovascularization of diabetic myocardial infarction rats, which provides a new target for clinical medication to reduce myocardial injury caused by myocardial infarction.

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Conflict of interests:
The authors declared no conflicts of interest.

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