**Abstract.** Ischemic postconditioning (IPoC) has been demonstrated to prevent myocardial ischemia-reperfusion injury (MIRI), but its cardioprotective effect is abrogated by hypercholesterolemia. The aim of the present study was to determine whether lycopene (LP), a type of carotenoid, can restore the cardioprotective effect of IPoC in hypercholesterolemic rats. Male Wistar rats were fed a cholesterol-enriched diet for 12 weeks to establish a hypercholesterolemic model. The rat hearts were isolated and subjected to 30 min ischemia and 60 min reperfusion using a Langendorff apparatus. LP was administered to the rats intraperitoneally for 5 consecutive days prior to ischemia and reperfusion. Myocardial pathological changes, infarct size and cell apoptosis were measured by hematoxylin and eosin, triphenyltetrazolium chloride and TUNEL staining, respectively. The changes in endoplasmic reticulum (ER) stress markers, the reperfusion injury salvage kinase (RISK) pathway and mitochondrial apoptosis-related proteins were detected by western blotting. Overall, the results demonstrated that low-dose LP in combination with IPoC ameliorated myocardial histopathological changes, reduced the infarct size and release of cardiac enzymes, and decreased cardiomyocyte apoptosis in hypercholesterolemic rats, but no beneficial effects were achieved by the same dose of LP or IPoC treatment were used alone. Furthermore, the combination of LP and IPoC inhibited the expression of glucose-regulated protein 78 and C/EBP homologous protein, increased the phosphorylation levels of AKT, ERK1/2 and glycogen synthase kinase-3β, repressed mitochondrial permeability transition pore opening, and reduced the expression of cytochrome c, cleaved caspase-9 and cleaved caspase-3. Collectively, these findings demonstrated that LP can restore the cardioprotective effects of IPoC on MIRI in hypercholesterolemic rats, and this restoration by LP was mediated by inhibition of ER stress and reactivation of the RISK pathway in hypercholesterolemic rat myocardium.

**Introduction**

Preventing myocardial ischemia-reperfusion injury (MIRI) is a frequent problem during the application of reperfusion therapy, such as percutaneous coronary intervention in patients with ST-segment elevation myocardial infarction (1). Accumulating evidence has demonstrated that a short series of repetitive cycles of reperfusion and ischemia performed at the onset of sustained reperfusion, namely ischemic postconditioning (IPoC), attenuated MIRI in several animal models (2-6) and certain human studies (7,8). However, recent studies have reported that the cardioprotective effect of IPoC is usually abrogated by pathological conditions, including hypercholesterolemia (9), although the mechanism underlying the loss of cardioprotection in hypercholesterolemic animals remains elusive.

It is widely accepted that hypercholesterolemia is one of the most important risk factors for the development of atherosclerosis, but there is evidence that hypercholesterolemia also exerts direct negative effects on the myocardium itself (10), including increased endoplasmic reticulum (ER) stress (11), altered activation of signaling pathways (12,13) and induction of apoptosis (13), and some researchers believe that this may partly explain why the cardioprotective effect of IPoC is suppressed by hypercholesterolemia (14). For example, Andreadou et al (15) investigated whether hypercholesterolemia suppresses the protective effect of IPoC through a decrease in the phosphorylation of endothelial nitric oxide synthase (eNOS) and the production of nitric oxide (NO). Our previous studies demonstrated that the repression of IPoC by hypercholesterolemia is associated with inactivation of the...
reperfusion injury salvage kinase (RISK) pathway (13) and excessive mitochondrial permeability transition pore (mPTP) opening (12). Of note, it was observed that fasudil, a Rho-kinase inhibitor, restored the cardioprotective effect of IPoC in hypercholesterolemic rats by activating the PI3K/AKT/eNOS signaling pathway (16). Furthermore, it was demonstrated that blocking of mPTP opening with cyclosporine A could restore the benefits of IPoC in hypercholesterolemic rat hearts (17). Therefore, ameliorating the negative impact of hypercholesterolemia on the myocardium by pharmacological intervention appears to be a novel method for preserving or restoring the cardioprotection of IPoC under hypercholesterolemic conditions.

Lycopene (LP), a type of carotenoid present in tomatoes and other red fruits and vegetables, has potent antioxidant properties, and has been reported to reduce the risk of multiple diseases, including coronary heart disease (18). More importantly, several recent studies have demonstrated that LP protects against MIRI by inhibiting the development of ER stress (19) and blocking mPTP opening (20). Given that LP can alleviate the negative impact of hypercholesterolemia on the myocardium, we hypothesized that LP treatment may preserve or restore the protective effect of IPoC against MIRI in the presence of hypercholesterolemia.

The aim of the present study was to determine whether LP treatment could restore the cardioprotective effect of IPoC in hypercholesterolemic rat hearts, and elucidate the underlying mechanism.

Materials and methods

Animals and establishment of a hypercholesterolemic rat model. A total of 110 male Wistar rats, weighing 100±10 g, were purchased from Changsheng Biotechnology Co., Ltd. All rats were housed in an environment of constant temperature (22-25°C) and relative humidity (50-60%) on a 12/12 h light/dark cycle, and food and water were provided ad libitum. To establish the hypercholesterolemic rat model, the rats were divided in two groups as follows: i) High-cholesterol diet (HCD) group (n=90), rats were consecutively fed normal feedstuff for 12 weeks; and ii) normal diet (ND) group (n=20), rats were consecutively fed normal feedstuff for 12 weeks. Moreover, the period of LP treatment through intraperitoneal injection in rats ranged between 2.5 and 20 mg/kg/day; therefore, 5 mg/kg/day was selected as the low dose and 20 mg/kg/day as the high dose of LP treatment. Moreover, the period of LP treatment through intraperitoneal injection ranged between 1 and 10 days in previous studies. Therefore, an intermediate treatment period of 5 days was selected for this study. LP was administered intraperitoneally at a dose of 5 mg/kg/1 h prior to ischemia, then IR experimental procedures were as described for the IR group.

According to previous studies (23-25), the dose of LP used for intraperitoneal injection in rats ranged between 2.5 and 20 mg/kg/day; therefore, 5 mg/kg/day was selected as the low dose and 20 mg/kg/day as the high dose of LP treatment. Moreover, the period of LP treatment through intraperitoneal injection ranged between 1 and 10 days in previous studies. Therefore, an intermediate treatment period of 5 days was selected for this study. LP was administered intraperitoneally at a dose of 5 mg/kg/day for 5 consecutive days. The dose of 4-PBA was based on a previous study (26).

Measurement of changes in serum lipids after LP treatment. To assess whether LP has an effect on the level of serum lipids, 45 rats in the LP-5, IPoC + LP-5 and LP-20 group (n=15 per group) were performed lipid profile as described earlier, after received LP administration at a dose of 5 or 20 mg/kg/day for 5 consecutive days.

Observation of myocardial mitochondria by electron microscopy. At the end of the reperfusion period, a 1x1x1-mm piece of cardiac tissue was isolated from the left ventricle and electron microscopic sections were prepared as previously described (27). The morphological changes of the mitochondria were
observed by transmission electron microscopy (JEM-1200EX; JEOL, Ltd.).

**Hematoxylin and eosin staining.** The hearts were removed from the perfusion apparatus at the end of the reperfusion period. The left ventricular heart tissues were fixed in 4% paraformaldehyde for 24 h at room temperature, dehydrated and embedded in paraffin. The paraffin blocks were then cut into 5-µm sections. The paraffin sections were dewaxed, stained with hematoxylin for 5 min, followed by 1% hydrochloric acid alcohol differentiation for 3 sec and eosin staining for 3 min at room temperature. The pathological changes in heart tissues were observed under a light microscope and imaged. To evaluate myocardial damage quantitatively, the sections were scored by two independent observers blinded to the experimental protocol, as previously described (28): 0, no damage; 1, interstitial edema and focal necrosis; 2, diffuse myocardial cell swelling and necrosis; 3, necrosis with the presence of contraction bands and neutrophil infiltrate; and 4, widespread necrosis with the presence of contraction bands, neutrophil infiltrate and hemorrhage.

**Measurement of infarct size.** Myocardial infarct size was measured by triphenyltetrazolium chloride (TTC) staining. The hearts were harvested at the end of the reperfusion period and the heart samples were frozen at -80°C for 30 min. The frozen hearts were cut into 1-2-mm sections from the apex to the bottom and incubated in 1% TTC solution at 37°C for 30 min. Subsequently, the heart sections were washed with 1X PBS and fixed in 4% paraformaldehyde overnight at room temperature. The stained sections were photographed by a digital camera and analyzed using ImageJ 2x analysis software (National Institutes of Health). The severity of myocardial infarction was reflected by the ratio of infarct size to total size.

**Measurement of lactate dehydrogenase (LDH) and creatine kinase muscle/brain isoenzyme (CK-MB) activity.** LDH and CK-MB are released from cardiomyocytes into the coronary effluent during perfusion (29). The activities of LDH and CK-MB in the coronary effluent were determined using the LDH and CK-MB assay kits (Nanjing Jiancheng) using 5% skimmed milk for 1 h at room temperature, and then incubated with specific antibodies, including anti-glucose-regulated protein 78 (GRP78; 1:1,000; Abcam; cat. no. ab108613), anti-β/EBP homologous protein (CHOP; 1:1,000; Abcam; cat. no. ab11419), anti-β-AKT (1:1,000; Abcam; cat. no. ab179463), anti-phospho (p)-AKT (phospho S473; 1:1,000; Abcam; cat. no. ab81283), anti-ERK1/2 (1:1,000; Abcam; cat. no. ab17942), anti-p-ERK1/2 (phospho T202 + Y204; 1:1,000; Abcam; cat. no. ab214362), anti-glycogen synthase kinase-3β (GSK-3β; 1:1,000; Abcam; cat. no. ab93926), anti-p-GSK-3β (phospho S9; 1:1,000; Abcam; cat. no. ab107166), anti-cytochrome c (1:1,000; Abcam; cat. no. ab133504), anti-cleaved caspase-9 (1:1,000; Abcam; cat. no. ab2324), anti-cleaved caspase-3 (1:1,000; Abcam; cat. no. ab2302) and anti-β-actin (1:1,000; Zhongshan Jinqiao Biotechnology; cat. no. TA-09), at 4°C overnight. On the following day, the PVDF membrane was incubated with horseradish peroxidase-conjugated goat anti-rabbit IgG (1:4,000; EarthOx Life Science; cat. no. E030120-01) or goat anti-mouse IgG (1:4,000; EarthOx Life Science; cat. no. E030110-01) at room temperature for 30 min. Detection of protein bands was performed using an ECL for western blotting kit (Beyotime Institute of Biotechnology) according to the manufacturer's instructions. Relative densitometry was calculated with Image J2x analysis software.

**Statistical analysis.** All data are presented as mean ± SD and were analyzed using SPSS version 17.0 (SPSS, Inc., Chicago, IL, USA). Differences among three or more groups were first evaluated using one-way analysis of variance, and if the differences were significant, multiple comparison analysis was further performed with Fisher's least significant difference test. Differences between two groups were evaluated using independent samples t-test. P<0.05 was considered to indicate a statistically significant difference.

**Results**

**Changes in serum lipids.** The serum levels of TC and LDL were significantly increased in the HCD group as compared with the ND group after 12 weeks (Table I). The result suggested that the hypercholesterolemic rat model was successfully established. Moreover, there was no significant difference in the serum level of TC and LDL after a 5-day LP treatment either at the dose of 5 or 20 mg/kg/day (Table II), suggesting...
that short-term LP treatment did not affect the serum levels of TC and LDL.

**Figure 1.** Histopathological changes of the ischemic myocardium. (A) Myocardial submicroscopic structure examined by transmission electron microscopy; scale bar, 20 μm. The swelling of the mitochondria was reduced and mitochondrial cristae were denser, as indicated by arrows. (B) Myocardium was stained with hematoxylin and eosin and examined under an optical microscope; scale bar, 50 μm. Myocardial degeneration and inflammatory infiltration are indicated by arrows. (C) Myocardial damage was evaluated quantitatively. The data are expressed as mean ± SD; n=6 per group. *P<0.05 vs. IR group. IR, ischemia-reperfusion; IPoC, ischemic postconditioning; LP-5/LP-20, 5 or 20 mg/kg/day LP treatment for 5 days prior to induction of ischemia; LP, lycopene; 4-PBA, 4-phenylbutyric acid.

**Table I.** Analysis of serum lipid profile.

| Group                        | TC (mg/dl) | HDL-C (mg/dl) | LDL-C (mg/dl) |
|------------------------------|------------|---------------|---------------|
| Normal diet (n=20)           | 65.27±6.76 | 26.61±5.87    | 32.28±6.59    |
| High cholesterol diet (n=90) | 132.08±17.42 | 27.19±5.23   | 92.96±19.08   |

The data are expressed as means ± standard deviation. *P<0.05 vs. normal diet group. TC, total cholesterol; HDL-C, high-density lipoprotein cholesterol; LDL-C, low-density lipoprotein cholesterol.

**Table II.** Analysis of the change in the serum lipid profile following lycopene (5 or 20 mg/kg/day) treatment.

| Lycopene treatment | TC (mg/dl) | HDL-C (mg/dl) | LDL-C (mg/dl) |
|--------------------|------------|---------------|---------------|
| Before             | 132.08±17.42 | 27.19±5.23   | 92.96±19.08   |
| After              | 131.37±16.21 | 26.97±4.27   | 92.40±16.83   |

The data are expressed as the mean ± standard deviation (n=45). TC, total cholesterol; HDL-C, high-density lipoprotein-cholesterol; LDL-C, low-density lipoprotein cholesterol.

LP restores the protective effect of IPoC in hypercholesterolemic rats. In the IPoC + LP-5 group, the swelling of the mitochondria
was reduced and mitochondrial cristae were denser, as observed by electron microscopy (Fig. 1A); myocardial degeneration and inflammatory infiltration were also less extensive, as observed by light microscopy (Fig. 1B and C); myocardial infarct size (Fig. 2A and B) and the release of cK-MB and LdH (Fig. 2C and D) were reduced and the apoptosis rate was decreased (Fig. 3), whereas cardiac function, as reflected by LVDevP and ±dp/dt, was improved (Fig. 4) in the IPoc + LP group compared with the IR group under conditions of hypercholesterolemia, although no beneficial effects were observed in the LP‑5 and IPoc only groups. These results suggest that LP pretreatment may preserve or restore the cardioprotective effect of IPoc in hypercholesterolemic rats. Moreover, the LP-20 and 4-PBA groups displayed beneficial effects similar to the effects in the IPoc + LP-5 group, suggesting that high-dose LP treatment or pharmacological ER stress inhibition may protect hypercholesterolemic rats against MIRI.

Figure 2. Protective effect of IPoc in combination with LP treatment on hypercholesterolemic rat hearts. (A) The ischemic myocardium was stained by triphenyltetrazolium chloride solution. The viable myocardium was stained red and the infarct area remained unstained. Scale bar, 2 mm. (B) Statistical analysis of myocardial infarct size. (C) Release of CK-MB in the coronary effluent. (D) Release of LDH in the coronary effluent. The data are expressed as the mean ± SD; n=6 per group. *P<0.05 vs. IR group. IR, ischemia reperfusion; IPoc, ischemic postconditioning; LP-5/LP-20, 5 or 20 mg/kg/d LP treatment for 5 days prior to induction of ischemia; LP, lycopene; 4-PBA, 4-phenylbutyric acid; CK-MB, creatine kinase muscle/brain isoenzyme; LDH, lactate dehydrogenase.

Effects of LP, alone or combined with IPoc, on ER stress in hypercholesterolemic rats. As shown in Fig. 5, the protein expression of GRP78 and CHOP (markers of ER stress) was markedly decreased in the IPoc + LP-5 and LP-20 groups compared with IR and IPoc groups, which was similar to the effect of the ER stress inhibitor 4-PBA. However, no significant change was observed in the IPoc and LP-5 only groups. Therefore, high-dose LP or the combination of low-dose LP and IPoc were successful in reducing ER stress in hypercholesterolemic rats.

Effects of LP, alone or combined with IPoc, on the RISK pathway in hypercholesterolemic rats. Considering that the RISK pathway is suppressed in hypercholesterolemic rat hearts (12,13), the effect of LP, alone or combined with IPoc, on the activation of AKT, ERK1/2 and gSK-3β was investigated. The results revealed that the phosphorylation of AKT (Fig. 6A), ERK1/2 (Fig. 6B) and gSK-3β (Fig. 6C) were enhanced in the IPoc + LP-5, LP-20 and 4-PBA groups, compared with the IR group. However, no significant change in the phosphorylation levels of AKT, ERK1/2 or GSK-3β was observed in the IPoc and LP-5 only groups. Thus, high-dose LP or the combination of low-dose LP and IPoc successfully activated the RISK pathway in hypercholesterolemic rats.

Effects of LP, alone or combined with IPoc, on mPTP opening in hypercholesterolemic rats. mPTP is the end effector of IPoc and is regulated by the RISK pathway (30). Thus, the effect of LP, alone or combined with IPoc, on mPTP opening was investigated. The results demonstrated that the sensitivity of mPTP to calcium, an inducer of mPTP opening (31), was markedly decreased in the IPoc + LP-5, LP-20 and 4-PBA
Figure 3. Effect of IPoC in combination with LP treatment on myocardial cell apoptosis. (A) Myocardial cell apoptosis was measured in a TUNEL assay. The apoptotic cell nuclei were stained brown and the living cell nuclei were stained blue. (B) Statistical analysis number of TUNEL-positive cells. The data are expressed as the mean ± SD; n=6 per group. *P<0.05 vs. IR group. IR, ischemia reperfusion; IPoC, ischemic postconditioning; LP-5/LP-20, 5 or 20 mg/kg/d lycopene treatment for 5 days prior to induction of ischemia; LP, lycopene; 4-PBA, 4-phenylbutyric acid.

Figure 4. Effect of IPoC in combination with LP treatment on cardiac function in hypercholesterolemic rat hearts. (A) Effect on heart rate. (B) Effect on LVDevP. (C) Effect on +dp/dt. (D) Effect on -dp/dt. The data are expressed as means ± standard deviation; n=6 per group. *P<0.05 vs. IR group. IR, ischemia reperfusion; IPoC, ischemic postconditioning; LP-5/LP-20, 5 or 20 mg/kg/day lycopene treatment for 5 days prior to induction of ischemia; LP, lycopene; 4-PBA, 4-phenylbutyric acid; LVDevP, left ventricular developed pressure; +dp/dt, positive first-order derivative of ventricular pressure; -dp/dt, negative first-order derivative of ventricular pressure.
Figure 5. Effect of IPoC in combination with LP treatment on endoplasmic reticulum stress in hypercholesterolemic rat hearts. (A) Protein expressions of GRP78 and CHOP were detected by western blotting. (B) Densitometry analysis of the expression of GRP78. (C) Densitometry analysis of the expression of CHOP. The data are expressed as mean ± SD; n=3 per group. *P<0.05 vs. IR group. IR, ischemia reperfusion; IPoC, ischemic postconditioning; LP-5/LP-20, 5 or 20 mg/kg/day lycopene treatment for 5 days prior to induction of ischemia; LP, lycopene; 4-PBA, 4-phenylbutyric acid; GRP78, glucose-regulated protein 78; CHOP, C/EBP homologous protein.

Figure 6. Effect of IPoC in combination with LP treatment on the reperfusion injury salvage kinase pathway in hypercholesterolemic rat hearts. (A) Phosphorylation of AKT was measured by western blotting and T-AKT was used as the internal reference. (B) Phosphorylation of ERK1/2 was measured by western blotting and T-ERK1/2 was used as the internal reference. (C) Phosphorylation of GSK-3β was measured by western blotting and T-GSK-3β was used as the internal reference. The data are expressed as the mean ± SD; n=3 per group. *P<0.05 vs. IR group. IR, ischemia reperfusion; IPoC, ischemic postconditioning; LP-5/LP-20, 5 or 20 mg/kg/day lycopene treatment for 5 days prior to induction of ischemia; LP, lycopene; 4-PBA, 4-phenylbutyric acid; p-, phospho-; T-, total; GSK-3β, glycogen synthase kinase.
groups, compared with the IR group (Fig. 7A and B), but no significant change was observed in the IPoC and LP-5 groups. mPTP opening may lead to the activation of the mitochondrial apoptosis pathway. As expected, the expressions of mitochondrial apoptosis-related proteins, including cytochrome c, cleaved caspase-9 and cleaved caspase-3, were also reduced...
in the LP-5, LP-20 and 4-PBA groups, compared with the IR group (Fig. 7C-E), but no significant change was observed in the IPoC and LP-5 only groups, which was in accordance with the extent of mPTP opening. Taken together, these results confirmed that high-dose LP or the combination of low-dose LP and IPoC inhibit mPTP opening in hypercholesterolemic rat hearts.

Discussion

Whether the cardioprotective effect of IPoC is suppressed by hypercholesterolemia remains controversial (14). The results of the present study demonstrated that the cardioprotection of IPoC is lost in rats fed a high-cholesterol diet for 12 weeks, which is consistent with previous findings in mini-swine fed a cholesterol-enriched diet for 4 weeks (32) and rabbits fed a cholesterol-enriched diet for 6 weeks (33). However, Donato et al. (34) reported that the cardioprotection of IPoC was preserved in rabbits after 4 weeks on a cholesterol-enriched diet. It is hypothesized that this discrepancy may be partly attributed to the differences in the duration of the cholesterol-enriched diet and the animal species.

Statins, a widely used class of lipid-lowering drugs, were found to restore the loss of IPoC benefits in hypercholesterolemic animals. Iliodromitis et al. (35) first reported that the loss of the IPoC benefits may be reversed in hypercholesterolemic rabbits that received a 3-week simvastatin treatment due to the decrease in TC and LDL plasma levels and attenuation of the oxidative and nitrosative stress in the ischemic myocardium. Moreover, Andreadou et al. (15) reported that a 3-day pravastatin treatment preserved the cardioprotective effect of IPoC in hypercholesterolemic rabbits, independently of any lipid-lowering effects, potentially through eNOS activation and the attenuation of nitro-oxidative stress. In the present study, a 5-day LP treatment achieved no decrease in serum lipids, but the combination of low-dose LP and IPoC attenuated MIRI in hypercholesterolemic rats. Although LP has antihyperlipidemic properties, its lipid-lowering effect usually requires long-term treatment. Zeng et al. (36) found that the values of TC and LDL had not significantly changed when measured after 1 week of LP treatment, but exhibited a significant decrease after ≥3 weeks of LP treatment. This may explain why blood lipid levels had not decreased after a 5-day LP treatment in the present study. Given that the cardioprotective effect of IPoC is abolished by hypercholesterolemia, LP administration may preserve or restore the protective effect of IPoC in hypercholesterolemic rats, independently of any lipid-lowering effects. Interestingly, it was also observed that hypercholesterolemic rats treated with high-dose LP alone exhibited similar cardioprotection as those receiving a combination of low-dose LP and IPoC. Thus, it may be further inferred that LP in combination with IPoC exerts synergistic or additive protective effects against MIRI in hypercholesterolemic rats. Although high-dose LP treatment exerted protective effects in IR hearts under conditions of hypercholesterolemia, its side effects on other organs could not be determined due to the isolated rat heart model used in the present study, which is a potential limitation for its use in clinical practice. In contrast to high-dose LP treatment, low-dose LP combined with IPoC exerted similar cardioprotective effects, but was associated with fewer potential side effects due to the lower LP dose; for instance, high doses of LP may increase the risk of lung cancer in smokers (37,38). Therefore, combined treatment may represent a safer method for preventing MIRI in patients with hypercholesterolemia.

ER is responsible for the biosynthesis of all secreted and membrane proteins. The accumulation and aggregation of unfolded proteins in the ER leads to a condition referred to as ER stress (39). The unfolded protein response is initially activated to degrade unfolded proteins and restore normal ER function. If ER stress is prolonged, or if the adaptive response fails, the cells will die through apoptosis (40). Thus, ER stress is crucial for most cellular activities and survival. GRP78 and CHOP, two major ER stress-related proteins, reflect the activation of ER stress. The levels of GRP78 and CHOP were found to increase in the ischemic myocardium by the combination of low-dose LP and IPoC, which is similar to the effects of ER stress inhibitors, such as 4-PBA. Considering that hypercholesterolemia increases ER stress in the IR myocardium (11) and LP reduces ER stress in hypoxic-reoxygenated cardiomyocytes (19), it may be deduced that the restoration of IPoC in hypercholesterolemic rat by LP treatment may be mediated via inhibition of ER stress.

IPoC protects against MIRI through activation of the RISK pathway (AKT and ERK1/2) (41), and subsequently inhibiting GSK-3β and mPTP opening (42); however, hypercholesterolemia suppresses the activation of the RISK pathway and induces the activation of GSK-3β and mPTP opening (12,13). In the present study, the combination of LP and IPoC activated AKT and ERK1/2, subsequently inhibiting GSK-3β and mPTP opening in hypercholesterolemic rat hearts, suggesting that the restoration of IPoC by LP may be partly attributed to the reactivation of the RISK pathway. ER stress has been demonstrated to impair the activation of the RISK pathway (43). Miki et al. (43) provided evidence that ER stress may suppress the phosphorylation of AKT and ERK1/2, and further impair GSK-3β-mediated suppression of mPTP opening. Notably, the combination of LP and IPoC exerted the same effect on the RISK pathway as that of an ER stress inhibitor. Thus, it may be inferred that LP reactivates AKT and ERK1/2 in hypercholesterolemic rat hearts partly through inhibition of ER stress.

There were certain limitations to the present study. First, although LP restored the cardioprotection of IPoC in hypercholesterolemic rats by inhibition of ER stress and activation of the RISK pathway, whether activating ER stress or inhibiting the RISK pathway by a pharmacological or genetic approach could reverse the restoration of IPoC in hypercholesterolemic rats was not determined. Therefore, these conclusions should be further confirmed in future studies. Second, MIRI was mimicked in isolated rat hearts using a Langendorff apparatus. In this model, the ischemic heart was only subjected to 60-120 min reperfusion and then harvested (44). Therefore, cardiac fibrosis or remodeling, including changes in histology and protein biomarkers (matrix metallopeptidase-2/9 and cathepsins S/K) (45-48), could not be observed in this model within this short period of time. Finally, considerable evidence indicates that reactive oxygen species (ROS) have a predominant role in myocardial damage induced by reperfusion (49,50).
However, the present study mainly focused on the role of the regulation of ER stress and the RISK signaling pathway in the restoration of IPOc by LP under hypercholesterolemic conditions and the lack of ROS detection is a limitation of the present study. Additional mechanisms that may be involved in this process, such as oxidative stress mediated by ROS, require further investigation.

Collectively, to the best of our knowledge, the findings of the present study are the first to demonstrate that LP restores the cardioprotective effect of IPOc in hypercholesterolemic rats. Furthermore, the restoration of IPOc by LP in hypercholesterolemic rat hearts was found to be associated with the inhibition of ER stress and activation of the RISK pathway. Given that LP is non-toxic and has fewer adverse effects compared with statins, the combination of LP and IPOc may be a novel therapy for preventing MIRI in patients with hypercholesterolemia that receive percutaneous coronary intervention.

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Availability of data and materials

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

Authors’ contributions

DJ and NW conceived and designed the experiments. LD conducted the experiments. CL, XL, ZH and SL participated in the completion of the experiments. LD and NW analyzed the data. LD wrote the paper. All authors read and approved the final version of this manuscript.

Ethics approval and consent to participate

The experimental protocol was approved by the Institutional Ethics Committee of China Medical University.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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