1 Introduction

The heart is the most energy-consuming organ in the human body, but the concentration of ATP in myocardial cells is relatively low, which means that myocardial cells must continuously synthesize ATP to maintain normal diastolic and contractile functions. Mitochondria are the most important place for intracellular oxidative phosphorylation and ATP synthesis. Cardiomyocytes contain a large number of mitochondria, a double-layer membrane-coated organelle that exists in most cells. It is the main structure for energy production in cells and the main place for cells to carry out aerobic respiration. In addition, mitochondria are also involved in cell apoptosis and signal transduction, cell proliferation, cell metabolism and other processes. Mitochondrial injury is the main pathogenesis that causes myocardial ischemia/reperfusion injury, which in turn leads to myocardial death, apoptosis and contractile failure. Therefore, mitochondrial dysfunction is a determinant of myocardial ischemia/reperfusion injury. Studies have found that autologous mitochondria transplantation, in which a person’s autologous healthy mitochondria is delivered to the ischemic area of damaged mitochondria through mitochondria in coronary arteries, can significantly improve myocardial function (Giorgio et al., 2018; Shin et al., 2019; Chiva-Blanch & Badimon, 2017; Espinosa-Moncada et al., 2018; Vetrani et al., 2018).

Lycopene has a long-chain polyunsaturated olefin molecular structure, which gives it a strong capability to eliminate free radicals and antioxidant capacity. The current research on its biological effects mainly focuses on anti-oxidation, reducing the risk of cardiovascular disease, reducing genetic damage, and inhibiting the occurrence and development of tumors. Our research is to explore whether Lycopene can mitigate cardiac ischemia/reperfusion injury, and to further explore whether its underlying mechanism plays a role by inhibiting the opening of MPTP.

2 Methods

2.1 Cell grouping and CCK-8 to detect the viability of H9c2 cells

H9c2 cells were obtained. Cell viability was determined according to the operation manual, and the OD value at 450 nm was measured. The groupings are as follows: A, the Control group,
in which the cells were grown in an incubator, without receiving sugar-free Earle's medium replacement; B, the H/R group, in which the cells received sugar-free Earle's medium and normal medium in sequence, meaning that the cells were treated with hypoxia for 12 h and reoxygenation for 1 h respectively; C, the low-dose drug group. After pretreatment with 5 μM Lycopene for 4 h, the cells received sugar-free Earle's medium and normal medium, and then received 12 h hypoxia and 1 h reoxygenation treatment; D, the middle-dose drug group, the cells received 4 h pretreatment with 10 μM Lycopene, other procedures are the same as group C; E, the high-dose drug group, the cells received pretreatment with 20 μM Lycopene for 4 h, with other procedures the same as group C.

2.2 Trypan blue staining method to determine the survival rate of H9c2 cells.

H9c2 cells were prepared. According to the operation procedure and principle of trypan blue staining: normal cardiomyocytes are not stained, only cardiomyocytes with loss of cell membrane integrity can be stained blue by trypan blue. Therefore, the calculation formula of cell survival rate is: cell survival rate = total number of unstained cells/total number of cells × 100%, and the grouping is the same as above.

2.3 LDH activity detection of H9c2 cell damage.

Following the instructions, the activity of LDH was calculated according to the formula: LDH (U/L) = (measured OD value - control OD value) / (standard OD value - blank OD value) × standard concentration × 1000.

2.4 Flow cytometry to measure the apoptosis rate of H9c2 cells

The groups are as follows: A, the Control group; B, the H/R group; C group, the cells were pretreated with 20 μM Lycopene for 4 h; and D group, the cells were pretreated with 20 μM Lycopene + 20 μM Atractylodes carboxylate for 4 h. Flow cytometry was used for apoptosis detection.

2.5 Rhodamine 123 method to measure the changes of mitochondrial membrane potential of H9c2 cardiomyocytes.

According to the operation and dyeing principle of Rhodamine 123, the stronger fluorescence intensity and denser fluorescence mean the less reduction of mitochondrial membrane potential. The weaker fluorescence intensity and sparser fluorescence suggest more reduction of mitochondrial membrane potential.

2.6 Calcium induction method to measure the opening degree of mitochondria of H9c2 cardiomyocytes.

H9c2 cardiomyocyte mitochondria were extracted. Buffer A of the 190 μL of MPTP detection kit was added to the mitochondrial pellets of each group. The microplate reader was used for continuous detection of the A540 from 0 min to 10 min. The samples were added with Buffer B of the 10 μL of MPTP detection kit. A microplate reader was used to measure once at A540 at an interval of 1 min, and continuously for 10 min.

2.7 Detection of protein expression by WB method.

After SDS-PAGE gel electrophoresis, membrane transfer, western blotting, chemiluminescence operation, the expressions of Bcl-2, Bax, cytochrome C, APAF-1, caspase 9 and caspase-3 proteins were determined.

2.8 Statistical analysis.

The data were analyzed using SPSS 17.0 software. All data were expressed as mean ± standard deviation. Multi-sample comparisons of 3 or more groups were conducted using one-way analysis of variance, and the least significant difference test was used for multiple comparison pairwise. If P < 0.05, it is considered a statistically significant difference.

3 Result

3.1 The effect of Lycopene on the vitality of hypoxia/reoxygenation H9c2 cardiomyocytes.

Compared with the Control group, the A450 value of the H/R group was significantly reduced, and different concentrations of Lycopene can effectively improve the viability of H9c2 cardiomyocytes, as shown in Figure 1.

3.2 The effect of Lycopene on the survival rate of hypoxia/reoxygenation H9c2 cardiomyocytes.

Compared with the Control group, the survival rate of the H/R group was significantly reduced. Different concentrations of Lycopene can significantly improve the survival rate of H9c2 cardiomyocytes, as shown in Figure 2.

3.3 The effect of Lycopene on hypoxia/reoxygenation H9c2 cardiomyocyte damage.

Compared with the Control group, the LDH value of the H/R group was significantly higher. Different concentrations of Lycopene can effectively mitigate the damage of H9c2 cardiomyocytes, as shown in Figure 3.

** Figure 1. H9c2 myocardial cell viability results. **

\[ \text{Cell viability (\% of Control)} \]

\[ * * p < 0.01 \text{ vs H/R} \]
\[ \# \# p < 0.01 \text{ vs 5μM} \]
\[ \# p < 0.05 \text{ vs 5 μM} \]
\[ ! p < 0.05 \text{ vs 10μM} \]
3.4 The effect of Lycopene on the apoptosis rate of hypoxia/reoxygenation H9c2 cardiomyocytes.

Compared with the Control group, the apoptosis rate of the H/R group was significantly increased. Compared with the Lycopene group, the apoptosis rate measured in the H/R group was also significantly increased. Compared with the Control group, the protein expression level of the Lycopene + Atractylodes carboxylate group was significantly increased, as shown in Figure 4.

![Figure 2. Detection of H9c2 cardiomyocyte viability.](image)

** p < 0.01 vs H/R  * p < 0.05 vs H/R  ## p < 0.01 vs 5μM  # p < 0.05 vs 5μM

3.5 The effect of Lycopene on the changes of mitochondrial membrane potential of hypoxia/reoxygenation H9c2 cardiomyocytes.

Rhodamine 123 is a fluorescent dye that can penetrate the mitochondrial membrane. The amount of Rhodamine 123 entering the mitochondria is indeed closely related to the level of the mitochondrial membrane potential. If the mitochondrial membrane potential is high, more Rhodamine 123 will enter and retain, with strong fluorescence, and vice versa. Therefore, the fluorescence intensity of Rhodamine 123 reflected the level of mitochondrial membrane potential. Compared with the Control group, the mitochondria ΔΨm of the H/R group was significantly reduced. Compared with the Lycopene group, the mitochondria ΔΨm of the H/R group was significantly reduced, and the mitochondria ΔΨm of the Lycopene + Atractylodes carboxylate group was also significantly reduced, as shown in Figure 5.

![Figure 3. Detection of H9c2 cardiomyocyte damage.](image)

** p < 0.01 vs H/R  ## p < 0.01 vs 5μM  # p < 0.05 vs 5μM ! p < 0.01 vs 10μM  & p < 0.05 vs 20μM

3.6 The effect of Lycopene on the expression of pro-apoptotic factor protein in the mitochondrial apoptosis pathway.

Compared with the Control group, the protein expression level of the H/R group was significantly higher. Compared with the Lycopene group, the protein expression level of the H/R group was significantly increased, and the protein expression level of Lycopene + Atractylodes carboxylate was also significantly increased, as shown in Figure 6.

Compared with the Control group, the Bcl-2 expression of the H/R group was significantly reduced, the Bax expression was significantly increased, thus the Bcl-2/Bax protein ratio was greatly reduced. Compared with the Lycopene group, the expression level of Bcl-2 in the H/R group was significantly reduced, the expression level of Bax was significantly increased, so the ratio of Bcl-2/Bax was obviously reduced. The Bcl-2 expression level of the Lycopene + Atractylodes carboxylate group was significantly reduced, and the Bax expression level was also significantly increased, as shown in Figure 7.

![Figure 4. Detection of H9c2 cardiomyocyte apoptosis.](image)

A: Control  B: H/R  C: 20 μM Lycopene  D: 20 μM Lycopene + 20 μM Atractylodes carboxylate
3.7 The effect of Lycopene on the opening degree of mitochondria of hypoxia/reoxygenation H9c2 cardiomyocytes.

The samples of each group were continuously measured with a microplate reader at the A540 from 0 min to 10 min. The lower measured value means the higher sensitivity of MPTP to calcium, suggesting the higher openness of MPTP. The sensitivity of MPTP to calcium, that is, the opening degree of MPTP, which is reflected by the ratio of A540 (10 min) / A540 (0 min), is quantified. Because the value measured at A540 (0 min) is the highest, while the value measured at A540 (10 min) is the lowest, the ratio is expressed as min/max A540. The lower ratio of min/max A540 suggests the higher sensitivity of MPTP to calcium and the higher opening degree of MPTP, and vice versa. Our results showed that, compared with the Control group, the A540 value and min/max A540 value measured in the H/R group were significantly reduced. Compared with the administration group, the A540 value and min/max A540 value measured in the H/R group were significantly reduced, See Figure 8.

4 Discussion.

At present, substantive progress has been made in cellular events that may lead to fatal I/R injury following reperfusion after ischemia, and mitochondrial integrity is the core of this work. A number of mitochondrial-centric mechanisms are believed to play a role, such as the generation of reactive oxygen species (ROS), the opening of the mitochondrial permeability transition pore (MPTP), and the activation of endogenous apoptosis (Fan et al., 2017; Costabile et al., 2019; Brown et al., 2017). The apoptotic pathway inherent in mitochondria is characterized by increased mitochondrial membrane permeability. This loss of mitochondrial outer membrane integrity is widely regarded as the "irreversible point" of apoptosis, and Bcl-2 family proteins (including Bax, Bak, and bID) play an important role in this process. The translocation of DRP1 to mitochondria promotes Bax/Bak recruitment, oligomerization and pore formation at OMM, which triggers the release of cytochrome c. In addition, the recruitment of OMM by Bax/Bak is accompanied by the release of a new apoptotic factor (DDP/TIMM8a), which promotes the recruitment of DRP1 to the Bax/Bak site. DRP1 then passes through Bax/baki-dependent ubiquitination and stabilizes.

Mitochondria are the core participants in regulating cell homeostasis. They are essential for energy production. However, reactive oxygen species accumulate as by-products of the electron transport chain, causing mitochondrial damage. Oxidative stress-mediated ROS leads to rapid depolarization of
MPTP is a multi-protein giant channel complex that spans the inner and outer mitochondrial membranes, which basically allows small molecule communication between the matrix and the cytosol. The structural composition of pores is not well understood, which is a topic of ongoing debate. Preliminary studies have shown that the pore is composed of a voltage-dependent anion channel (VDAC) located in the outer mitochondrial membrane, adenine nucleotide transporter (ANT) in the inner mitochondrial membrane, and cyclophilin D (Cyp-D) in the matrix. However, subsequent gene deletion studies have shown that the formation of MPTP does not require ANT and VDAC, partly due to the presence of different subtypes of these putative pore components. Despite these observations, it is generally believed that Cyp-D is the main factor in the formation of MPTP (Zhou et al., 2019; Badi et al., 2019; Medak & Townsend, 2019).

In our research, it was found that Lycopene can inhibit the decrease of ΔΨm and reduce the sensitivity of MPTP to Ca2+, thus clarifying that Lycopene exerts an anti-apoptotic mechanism by inhibiting the opening of MPTP. On this basis, it is confirmed that Lycopene can achieve the regulation of MPTP through inhibiting BAX expression and promoting Bcl-2 expression. In order to further prove that Lycopene exerts myocardial protection by acting on MPTP and inhibiting the opening of MPTP, this study applied the MPTP opening agonist: Atractylidin. The experimental results showed that Atractylidin counteracts the effect of Lycopene in inhibiting cardiomyocyte apoptosis, and it was further found that Atractylidin counteracted the protective effect of Lycopene to stabilize mitochondria ΔΨm and reduce the sensitivity of MPTP to Ca2+.

5 Conclusion

In summary, it is concluded that Lycopene plays a protective role in mitigating MIRI by acting on MPTP and inhibiting the opening of MPTP.

Reference


Lycopene on Ischemia-reperfusion Myocardium


