Surgically Metabolic Resection of Pericardial Fat to Ameliorate Myocardial Mitochondrial Dysfunction in Acute Myocardial Infarction Obese Rats

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ABSTRACT

Background: Pericardial fat (PF) is highly associated with cardiovascular disease but the effectiveness of surgical resection of PF is still unknown for myocardial mitochondrial structure and function in acute myocardial infarction (AMI) with obesity. The aim of this study was to demonstrate the difference in myocardial mitochondrial structure and function between obese AMI with additionally resected PF and those without resected PF.

Methods: Obese rats with 12-week high fat diet (45 kcal% fat, n = 21) were randomly assigned into 3 groups: obese control, obese AMI and obese AMI with additionally resected PF. One week after developing AMI and additional resection of PF, echocardiogram, myocardial mitochondrial histomorphology, oxidative phosphorylation system (OXPHOS), anti-oxidative enzyme and sarcoplasmic reticulum Ca\(^{2+}\) ATPase 2 (SERCA2) in the non-infarcted area were assessed between these groups.

Results: There was significant improvement of systolic function in AMI with PF resection compared with the AMI group in the echocardiogram. Even though the electron microscopic morphology for the mitochondria seems to be similar between the AMI with PF resection and AMI groups, there was an improved expression of PGC-1α and responsive OXPHOS including NDUFB3, NDUFB5 and SDHB are associated with the ATP levels in the AMI with PF resection compared with those in the AMI group. In addition, the expression levels of antioxidant enzymes (MnSOD) and SERCA2 were improved in the AMI with PF resection compared with those in the AMI group.

Conclusion: Surgical resection of PF might ameliorate myocardial mitochondria dysfunction in obese AMI.

Keywords: Acute Myocardial Infarction; Heart Failure; Pericardial Fat; Surgical Resection
INTRODUCTION

Cardiovascular disorders are the leading cause of death representing all global deaths\(^1\) and acute myocardial infarction (AMI) is a common cause of cardiovascular disorders.\(^2\) Ischemia with inflammation in the cardiomyocyte is the main pathophysiology of AMI.\(^3\)

In addition, obesity is strongly associated with an increased risk of developing AMI or heart failure.\(^4,5\) Especially, pericardial fat (PF) has been shown to have a diverse role\(^6\) in locations not classically associated with adipose tissue storage,\(^7\) but with the occurrence of cardiovascular events.\(^8\) A large amount of PF has been known to be significantly associated with cardiovascular diseases,\(^2,4,5\) and decrease anti-inflammatory adiponectin production and increase the synthesis of pro-inflammatory adipokines in obesity.\(^6\) In response to AMI, mesenchymal cells from the PF could migrate into the myocardium to be transformed into fibroblasts\(^7,8\) and shift to a nidus for inflammation and oxidative stress subsequent to aggravating cardiovascular disease,\(^7,8\) which demonstrates the paracrine effect of PF as a pathological transformation for cardiac damage and dysfunction. A recent study demonstrated that surgically resected PF improved the inflammatory status.\(^9-11\)

In addition, mitochondrial oxidative phosphorylation system (OXPHOS) and adenosine triphosphate (ATP) in the cardiomyocyte were mainly suppressed in the animal with cardiac dysfunction\(^12,13\) and PGC-1\(\alpha\) with oxidative stress was also significantly associated with heart failure.\(^14-16\)

Therefore, we investigated whether surgical resection of PF ameliorates myocardial mitochondrial structure, and dysfunction in the AMI obese rat model.

METHODS

Rats and housing

Animal female Wistar rats (6 weeks old, 160 g) were purchased from Harlan (Madison, WI, USA) and housed in the Animal Care Center of Eulji University in Daejeon under controlled conditions. The rats were housed in pairs at 23 ± 1°C with a 12 hours light/dark cycle and given access to water and rat chow consisting of a ‘high fat diet (HFD)’ (45% calories from fat).

Study design

After 12 weeks of HFD, all rats (n = 21) were randomly assigned into 3 groups: control, AMI, and AMI with PF resection. Subjects were anesthetized with 10% chloral hydrate by peritoneal injection and ventilated with a rodent respirator. The chest was opened through the left thoracotomy so that the left anterior descending coronary could be visualized and permanently ligated with a 7-0 silk suture at the site of its emergence from the left ventricle or so that a sham operation could be performed without ligation as described previously.\(^15\) The control group (n = 7) had the sham operation among the rats (n = 14) who had a permanent ligation of the left anterior descending artery, the AMI with PF resection group (n = 7) had an additional PF resection (Supplementary Fig. 1). In the AMI with PF resection group, the PF covering the anterior aspect of the heart was removed during the AMI operation.

Sampling, histology and transmission electron microscopy (TEM)

The blood samples and myocardial tissues of all rats were collected after the echocardiogram and sacrifice. Serum was stored at ~80°C until further analyses, and myocardial tissues were snap-frozen and stored at ~80°C, and some of the non-infarct myocardial tissues were fixed for
histologic examination or TEM. Transverse sections in each group were fixed in 10% buffered formalin (Sigma Aldrich, Co., St Louis, MO, USA) and embedded in paraffin. Thick transverse sections of 1 μm were cut from the tissue block and stained with hematoxylin and eosin, and myocardial tissues were fixed in 2.5% paraformaldehyde-glutaraldehyde at 4°C for 24 hours and then washed with 0.1 M phosphate buffered solution at 4°C. After washing two times, the tissue was post-fixed with 1% OsO4 buffered solution (pH 7.4) for 1 hour and 30 minutes. Tissue samples were dehydrated by serial ethanol and propylene oxide treatment and embedded in Poly/EM Bed812 embedding medium, and the resin was then polymerized in a vacuum drying oven (Yamato Scientific, Tokyo, Japan) at 60°C for 48 hours. Non-infarct myocardial tissue was sectioned using an EM Ultra-microtome LKB-2088 and stained with 1% toluidine blue, and then, ultra-thin sections were double-stained with uranyl acetate and lead citrate and examined using a Hitachi H-7600 electron microscope (Hitachi, Tokyo, Japan).

**Echocardiogram**

An echocardiogram was undertaken one week after the AMI operation as described previously. All rats were anesthetized by intraperitoneal injection with a mixture of ketamine (50 mg/kg) and xylazine (1 mg/kg), and the hair of the chest wall of all rats was carefully removed, and warm ultrasound transmission gel was liberally applied to ensure an optimal image quality for the echocardiogram. A high-frequency 15-MHz linear transducer (Entos CL15-7; Philips Medical Systems, Bothell, WA, USA) connected to an ultrasound system (ATL-HDI5000; Philips Medical Systems) was used for the acquisition of echocardiographic images. All cardiac structure and function traces were manually measured with a caliper by an operator using the leading-edge method according to the American Society of Echocardiography.

**Real time polymerase chain reaction**

The acquired myocardial tissue was homogenized using a polytron homogenizer (Fisher Scientific Inc., Pittsburgh, PA, USA), and the homogenate was stored to permit the complete dissociation of nucleoprotein complexes. Homogenates containing 0.2 mL of chloroform per 1 mL of TRI Reagent (Molecular Research Center Inc., Cincinnati, OH, USA) were shaken, and total RNA from the homogenates was isolated according to the manufacturer's instructions, and equal amounts of total RNA were reverse transcribed into cDNA using the iScript cDNA Synthesis Kit (Bio-Rad, Hercules, CA, USA) (The primer sequences are listed in the Supplementary Table 1). Real-time polymerase chain reaction was done in a 20 μL reaction mixture containing 1 μg cDNA, 10 pmol of forward primer and reverse primer each, and 10 μL of SYBR Green Supermix (Bio-Rad) using a CFX96 Real-Time PCR Detection System (Bio-Rad Laboratories [Singapore] Pte Ltd., Singapore). The threshold cycle values for each target mRNA were normalized to the mRNA of glyceraldehyde-3-phosphate dehydrogenase (GAPDH), and the relative expression level of each target gene was calculated using the CFX Manager software version 1.5 (Bio-Rad).

**Western blot analysis**

Acquired myocardial tissues were homogenized in the Pro-Prep Protein Extraction Solution (Intron Biotechnology, Seoul, Korea) with protease inhibitor (Roche Applied Science, Penzberg, Germany) and phosphatase inhibitor (Roche Applied Science). All the homogenates were incubated overall for 30 minutes and centrifuged at 13,000 rpm (4°C) for 5 minutes, and the supernatant was transferred into a 1.5 ml tube. Lysates were run on a 4–20% MP TGX Precast Gel (Bio-Rad) and transferred to PVDF membranes using a wet, vertical Criterion Blotter (Bio-Rad). Membranes were blocked with 5% skim milk in 50 mmol/L
Tris-HCl (pH 7.4) and 150 mmol/L NaCl (TBS) and then incubated overnight at 4°C with antibodies. The anti-GAPDH antibody (1:1,000) was obtained from Cell Signaling (Beverly, MA, USA), and the NDUFB3, NDUFB5 and SDHB antibodies (1:1,000) were obtained from Novus Biologicals (Littleton, CO, USA).

Antibodies directed against PGC-1α (1:500) were obtained from Cell Signaling. All antibody dilutions were made in 0.5% bovine serum albumin [BSA] and 0.1% sodium azide-Tris-buffered saline-Tween 20 [TBST], and the membranes were incubated with the appropriate horseradish peroxidase-conjugated secondary antibody. Secondary antibodies (goat anti-rabbit IgG) were obtained from Cell Signaling, and the signal was developed using an enhanced chemiluminescence detection system (Millipore, Billerica, MA, USA).

**ATP measurements**

The acquired myocardial tissues were homogenized in mammalian cell ATP lysis solution (Perkin-Elmer, Waltham, MA, USA). The whole homogenates were centrifuged at 12,000 rpm (4°C), and the supernatants were transferred into 1.5 mL tubes. Then, the ATPlite assay was performed according to the manufacturer’s instructions (Perkin-Elmer), and the luminescence emitted from the ATP-dependent luciferase reaction was calculated using a 2030 Multi-label Reader (Perkin-Elmer).

**Statistical analysis**

All parameters were expressed as the mean ± standard deviation and analyzed by one-way analysis of variance with Tukey’s post hoc test. Statistical analyses were performed using SPSS (version 15.0; SPSS Inc., Chicago, IL, USA), and a P value < 0.05 was considered statistically significant.

**Ethics statement**

All experiments were approved by the Animal Welfare Committee of Eulji University (EUIACUC19-28) in Daejeon, Republic of Korea.

**RESULTS**

The comparison of the echocardiographic parameters between AMI and AMI with PF resection

Systolic interventricular septum (IVSs), diastolic interventricular septum (IVSd), and diastolic left ventricular posterior wall (LVPWd) were similar between the AMI with PF resection and AMI groups (Table 1). However, systolic left ventricular intraventricular dimension (LVIDs)

### Table 1. The comparison of echocardiographic parameters between groups

| Echocardiographic parameters | Control group | AMI group | AMI + PF resection group | P value |
|-----------------------------|---------------|-----------|--------------------------|---------|
| LVPWs (cm)                  | 0.15 ± 0.02   | 0.12 ± 0.02 | 0.12 ± 0.02               | 0.999   |
| LVIDs (cm)                  | 0.42 ± 0.04   | 0.64 ± 0.09 | 0.53 ± 0.11               | 0.243   |
| IVSs (cm)                   | 0.14 ± 0.03   | 0.12 ± 0.03 | 0.11 ± 0.02               | 0.536   |
| LVPWd (cm)                  | 0.13 ± 0.02   | 0.13 ± 0.02 | 0.13 ± 0.02               | 0.999   |
| LVIDd (cm)                  | 0.59 ± 0.07   | 0.72 ± 0.11 | 0.69 ± 0.12               | 0.490   |
| IVSd (cm)                   | 0.10 ± 0.02   | 0.12 ± 0.03 | 0.10 ± 0.02               | 0.131   |

AMI = acute myocardial infarction, PF = pericardial fat, LVPWs = systolic left ventricular posterior wall, LVIDs = systolic left ventricular intraventricular dimension, IVSs = systolic interventricular septum, LVPWd = diastolic left ventricular posterior wall, LVIDd = diastolic left ventricular intraventricular dimension, IVSd = diastolic interventricular septum.
and diastolic left ventricular intraventricular dimension (LVIDd) were slightly decreased in the AMI with PF resection group compared to the AMI group which shows that the left ventricular systolic function was significantly recovered in the AMI with PF resection group compared to the AMI group (34.3 ± 3.8% vs. 28.5 ± 7.3%, $P = 0.041$) (Fig. 1).

Histomorphology and transmission electron microscopic images of mitochondria in the myocardium

The electron microscopic images of the non-infarct area between the AMI and AMI with PF resection groups seem to be similar at first look (Fig. 2). However, the myocardial mitochondria in the AMI with PF resection group showed more electro-dense cristae and membrane than those in the AMI group.

Fig. 1. The comparison of cardiac function between AMI and AMI with PF resection. EF was measured by echocardiography. Values are expressed as mean ± standard deviation. AMI = acute myocardial infarction, PF = pericardial fat, EF = ejection fraction. $^*P < 0.05$.

Fig. 2. The electron microscopic image of the mitochondria in the peri-infarction area between AMI and AMI with PF resection. (A) AMI; (B) AMI with PF resection. Transmission electron microscopic images ($\times$3,000; scale bar 1,000 nm) in the myocardia. AMI = acute myocardial infarction, PF = pericardial fat.
mRNA expressions of some OXPHOS subunits, antioxidant enzyme and sarcoplasmic reticulum Ca\(^{2+}\) ATPase 2 (SERCA2)

Among the genes of complexes I and II implicated in cardiovascular diseases,\(^{20}\) the expressions of OXPHOS in the real-time quantitative RT-PCR showed that the NADH dehydrogenase (ubiquinone) 1 beta subcomplex (NDUFB) 5 was significantly decreased in the AMI group but was significantly recovered in the AMI with PF resection group as previously reported.\(^{21}\) However, the expression of NDUFB3, cytochrome c1 (CYC1), NADH dehydrogenase (ubiquinone) flavoprotein 1 (NDUFV1) and NADH dehydrogenase (ubiquinone) Fe-S protein 1 (NDUFS1) were not different between the AMI with PF resection and AMI groups (data not shown). The expression of complex II/succinate dehydrogenase B subunit (SDHB) was also significantly decreased in the AMI group but was recovered in the AMI with PF resection group (Fig. 3). In addition, the expressions of anti-oxidant enzymes such as superoxide dismutase (SOD) and SERCA2 were decreased in the AMI group but were recovered in the AMI with PF resection group (Figs. 3 and 4).

![Fig. 3. The comparison of OXPHOS subunit mRNA expression between AMI and AMI with epicardial adipose resection. mRNA expression levels of the following OXPHOS subunits are depicted: NDUFB3 of complex I (A); NDUFB5 of complex I (B); SDHB of complex II (C). Representative blots for OXPHOS subunit: NDUFB5 level (D). Values are expressed as mean ± standard deviation. OXPHOS = oxidative phosphorylation system, AMI = acute myocardial infarction, NDUFB = NADH dehydrogenase (ubiquinone) 1 beta subcomplex, SDHB = succinate dehydrogenase B subunit, PF = pericardial fat. *P < 0.05 vs. control; #P < 0.05 vs. AMI.](https://doi.org/10.3346/jkms.2022.37.e55)
ATP and western blot analysis of NDUFB 5 and PGC-1α
The levels of ATP synthesis were significantly decreased in the AMI group but were recovered in the AMI with PF resection group. PGC-1α was significantly decreased in the AMI group but was recovered in the AMI with PF resection group (Fig. 5, Supplementary Fig. 2).

DISCUSSION
In the present study, OXPHOS subunit NDUFB5, SDHB and PGC-1α with ATP were decreased in AMI group but recovered in AMI with PF resection group compared with those in the AMI group. SERCA2 expression and MnSOD were also significantly decreased in AMI group but recovered in AMI with PF resection group compared with those in the AMI group.
A large amount of PF lead to decline release of adiponectin and instead synthesizes proinflammatory adipokines and the mesenchymal cells from the epicardium could migrate to be transformed into fibroblasts. The accumulation of PF is accompanied by inflammation and fibrosis and the release of proinflammatory adipocytokines from PF depots may thus contribute to the comorbidities that are characteristically seen in heart failure. Regardless of the source of systemic inflammation, PF itself leads to inflammation and fibrosis of the neighboring myocardium to development of heart failure.

There is previous report on the relationship between the resection of PF and cardiac dysfunction. The accumulation of PF is also clinically associated with cardiac dysfunction, and structural remodeling in obese people, and surgical resection for PF in AMI could induce beneficial effects for ischemic heart disease.

Mitochondria have been found to play an important role in the cardiac energetics in the energy production of ATP and mitochondria occupy 30–40% of the total area of a cardiomyocyte, where it is indispensable for normal cardiac energetics. The ob/ob and db/db mice and fa/fa rats with systolic dysfunction which show a decreased rate of intracellular Ca²⁺ reduction via SERCA2 in myocytes. Mitochondria are also responsible to maintain in tricellular Ca²⁺ turnover and the constant energy demands by oxidative phosphorylation (gene set as OXPHOS), and decreased calcium handling was found in the heart failure. In addition, mitochondrial genes such as OXPHOS and PGC1-α, a key metabolic regulator that coordinates in mitochondrial biogenesis and PGC1-α also acts as a crucial regulator of oxidative metabolism in cardiovascular disease. PGC1-α-null mice develop heart failure due to reduced mitochondrial OXPHOS in myocardial energy metabolism and oxidative stress has been known to be another axis in heart failure, and deficiencies of anti-oxidant enzyme contribute to mitochondrial damage in AMI as well.

In the present study, the myocardial mitochondria in the AMI with PF resection group showed more electro-dense cristae and membrane than those in the AMI group. The mitochondrial outer membrane has specific components for communication with other intra-cellular organelles and for the recognition and import of mitochondrial proteins while the inner membrane, the tubular-like cristae is the main machinery for respiratory chain complex containing OXPHOS for oxidative phosphorylation in the form of ATP. In our present study, it is speculated that more electro-dense crista might be related to more respiratory capacity in the mitochondria energy metabolism in the PF resection group. The expression of SERCA2 was recovered in the AMI with PF resection group compared with that in the AMI group, and the expressions of the OXPHOS subunit NDUFB5, SDHB, anti-oxidant enzyme and PGC-1α were also recovered in the AMI with PF resection group compared with those in the AMI group which were subsequently associated with increased ATP levels. Improved mitochondrial ATP synthesis in the non-infarct area might be related to increased systolic function in the AMI with PF resection group. However, further basic study is needed to support the beneficial effect for improvement of cardiac systolic function in the resection of PF.

There are some limitations in this study. First, this study included a relatively small number of animals which may limit the generalization of the results. Second, the volume and area of resected PF could be dependent on the operator and cardiac anatomy. Especially, pericardial or epicardial fat in the small animal was indistinguishable during the AMI operation. Third, a more detailed targeted experiment is needed to investigate the relationship between PF.
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resection and the improvement of cardiac systolic function. Forth, the experiment was a prospective study, but unmeasured selection bias may exist. Collectively, PF resection might be a beneficial effect to ameliorate myocardial mitochondrial dysfunction in the obese AMI rat model.

In conclusion, even though the myocardial mitochondrial morphology in the non-infarction area seemed to be similar between the AMI and AMI with PF resection groups, AMI with PF resection showed improved expressions of PGC-1α, antioxidant enzyme and OXPHOS subunit associated with ATP synthesis compared to AMI. Therefore, surgical resection of PF in the obese AMI might ameliorate myocardial mitochondria dysfunction.

SUPPLEMENTARY MATERIALS

Supplementary Table 1
Primer sequences of real-time polymerase chain reaction

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Supplementary Fig. 1
Study design and comparison of weight gain between AMI and AMI with PF resection. Data are expressed as mean ± standard error (n = 7).

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Supplementary Fig. 2
The example for comparison of western blot analysis of NDUFB5 and PGC-1α between AMI and AMI with PF resection. (A) NDUFB5 level; (B) PGC1-α level.

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