The Effects of Exercise Training Intensity on the Expression of C/EBPβ and CITED4 in Rats with Myocardial Infarction

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Abstract

Background: Exercise training (ET) has a cardioprotective effect and can induce new cardiomyocyte formation in physiological cardiac remodeling. However, following a large myocardial infarction (MI), the changes in cardiac regeneration and the subsequently different intensities of ET remain unexplored. We investigated the effect of different intensities of ET on the expression of C/EBPβ and CITED4 after an MI.

Methods: Adult male Wistar rats were exposed to left coronary artery ligation (LAD). Following an echocardiographic evaluation at four weeks after surgery, MI-operated animals were randomly assigned to either the MI-sedentary (MI-Sed) or the MI-trained groups (low intensity interval training (LIIT), moderate intensity interval training (MIIT), and high intensity interval training (HIIT)). MI-trained animals performed six weeks ET protocols (five days/week).

Results: Ten weeks after surgery, HIIT and MIIT animals had significantly higher LV ejection fraction compared to MI-Sed animals. HIIT animals had significantly diminished mRNA levels of C/EBPβ compared to MI-Sed. The mRNA level of CITED4 in HIIT and MIIT were significantly high compared to MI-Sed.

Conclusions: The present study demonstrates that cardiac functions are ET intensity-dependent, following a MI. It seems that higher intensities of ET are most effective agents in promoting the expression of C/EBPβ and CITED4. Thus, ET has emerged as an important variable in preclinical and clinical investigations.

Keywords: Myocardial Infarction, Exercise Training Intensity, C/EBPβ and CITED4

1. Background

Cardiovascular diseases (CVD) are the main cause of death in developed communities (1, 2). The exceedingly terrible clinical situation of CVD is represented by an intense myocardial infarction (MI) (3, 4). MI brings about a huge loss of cardiomyocytes and increases fibrocytes proliferation resulting in scar tissue formation and cardiac disorders (3). A few years ago, the idea that the heart is a post-mitotic organ with no regenerative limit had grabbed attention (5, 6). Nonetheless, information from animal models and human examinations recommended that the heart additionally has a constrained ability to produce new cardiomyocytes from ancestor cells and existing cardiomyocytes (7, 8). In fact, portions of the cardiomyocytes are supplanted within a typical human longevity (9, 10).

Exercise is a well confirmed intercession for the prevention and treatment of CVD (7, 11). Exercise applies its advantageous impacts by decreasing cardiovascular risk factors, and influencing the cellular and molecular remodeling of the heart (11, 12). Generally, increment in cardiomyocyte size is probably the central mechanism of exercise actuated cardiovascular development (11). Of late, it has been demonstrated that exercise training (ET) determines cardiac growth through the generation of new cardiomyocytes (13-17). It has been clinically proven that exercise is useful in averting and treating CVD. Past investigations have shown that ET could decrease the unfavorable remodeling of the harmed heart and enhance cardiac performance following a MI (12, 13).

The importance of the intensity (MIIT & HIIT) of exercise and its relationship with cardiovascular effects has been investigated in many studies (14-17). Still, the optimal...
dosage, intensity, frequency, and duration of exercise have remained unclear. Although a few weeks of vigorous ET can significantly augment cardiomyocyte formation (18), alternations in cardiac regeneration capacity following various intensities of ET after MI are yet to be elucidated.

2. Objectives

In this study, we have investigated the effects of low, moderate, and high intensity of ET on cardiac regeneration markers (C/EBPβ and CITED4) after a MI.

3. Methods

3.1. Study Design and Animals

Fifty-five adult male Wistar rats (230 g - 300 g ~ 10 weeks of age) were utilized for this investigation. Food and water were accessible ad libitum. All animal procedures were performed with the ethics committee of Rajaie Cardiovascular Medical and Research Center (RHC.A.C.IR.REC.1393.28). The investigation conformed to the guide for the care and use of laboratory animals published by the United States National Institutes of Health (NIH Publication, 8th edition, 2011).

3.2. Coronary Artery Ligation Procedure

Rats were orally intubated with a 16-gauge intravenous catheter in the wake of being anesthetized with intraperitoneal infusion of a blend of ketamine (75 mg/kg) and xylazine (5 mg/kg). A positive pressure ventilation was kept up with room air and oxygen utilizing a Harvard rodent ventilator (model 683, USA). The site of operation was shaved, and then disinfected with either Betadine or alcohol. Once the animal was unresponsive to toe-pinch, left-sided thoracotomy was performed and the left anterior descending coronary artery (LAD) was for all time ligated utilizing a 6-0 prolene (Ethicon Inc., USA) suture. Legitimate ligation of the LAD was affirmed by watching the whitening of the myocardial tissue distal to the suture and dyskinesia of the anterior wall (19). Sham-operated rats experienced a similar operation without coronary artery ligation to serve as the control group (Sham; n = 6). At the end of the operation, cefazolin (25 mg/kg IM), flunixin (1-2 mg/kg IM), and warm sterile saline (0.5-1 mL, SC) were infused and the rats were left on the warming cushion until the effect of anesthesia wore off. Postoperative care was kept utilizing analgesia and hemodynamic monitoring for 24 h (1, 20).

3.3. Echocardiographic Measurements

Transthoracic echocardiography was performed using a VIVID 7 dimension system (General Electric-Vingmed Ultrasound, Horten Norway), equipped with a 12-MHz electronic transducer. Echocardiographic assessments were performed a month after surgery and towards the end of the examination protocols. Under a similar anesthesia as for the infarct surgery which was kept up all through the echocardiographic examination, echocardiographic parameters had been recorded according with the American Society of Echocardiography guidelines. Left parasternal short-axis views of the left ventricle at the level of the papillary muscles was used to measure the following parameters: Left ventricular end-diastolic diameter (LVEDD), left ventricular end-systolic diameter (LVESD), posterior wall thickness in diastole (PWTd) and posterior wall thickness in systolic (PWTs). LV ejection fraction (EF) and fractional shortening (FS), was calculated according to the following formulas (21, 22):
\[
\%FS = \frac{LVEDD - LVESD}{LVEDD} \times 100
\]
\[
\%EF = \frac{LVEDD^2 - LVESD^2}{LVEDD^2} \times 100
\]

Echocardiographic appraisal was used a month after surgery. This was to assess the degree of MI and measure the cardiac contractile performance before initiation of further experimental procedures. Rats with high level of myocardial harm and created MI, assessed at FS \(\leq 35\%\), were incorporated into the examination (23).

3.4. Training Protocol

Out of 55 MI-operated rats, 25 rats which were survived from MI met the inclusion criteria as training protocol. The rats were assigned randomly to MI-trained groups high intensity interval training (HIIT, n = 6), moderate intensity interval training (MIIT, n = 6), and low intensity interval training (LIIT, n = 6). Familiarization with the motorized treadmill (Iranian Danesh Salar Company, Iran) was performed for MI-trained groups in the third and fourth week after surgery by walking for 5 m/min at slow pace five days per week for 5 min/day. The animals had 2 days rest before taking maximal oxygen consumption (VO2max). VO2max measured and applied at maximal stress test to evaluate the running speed of each rat (24). High, medium and low intensity interval training rats were exercised up to 70 min, five days a week. The training protocol included 10 minutes of warm-up (40%-50% VO2max) and 60 minutes of interval training (25, 26). Four minutes
running protocol was assessed with different intensities such as HIIT (85% - 90%), MIIT (65% - 70%) and LIIT (55% - 60%) of measured VO2max, and 2 minutes of active recovery between each interval HIIT and MIIT (50% - 60%) and LIIT (45% - 50%) of measured VO2max. No ET was gained by MI-Sed and Sham animals during this experiment procedure (25, 26). In previous studies the relationship between maximum VO2 and running speed in similar post-MI rat models was varied for every week (24). In particular, the running speed was expanded continuously for six weeks of training by 0.02 m/s every week (25). Treadmill inclination during training and testing was 0° (26).

3.5. Quantitative RT PCR (qRT-PCR)

The total RNA isolated from cardiac tissue was routinely used for reverse transcription polymerase chain reaction (RT-PCR) or quantitative real-time PCR (qPCR). The total RNA was extracted using EasyPure RNA Kit (TransGen Biotech, China) following the manufacturer’s instructions. For RT-PCR, the cDNAs were synthesized using Transcript First-Strand cDNA Synthesis SuperMix (TransGen Biotech, China) with total RNA as templates. qRT-PCR was performed using TransStart Top Green qPCR SuperMix (TransGen Biotech, China) on Rotor-Gene 6000 system (Corbett, Concorde, NSW, Australia). The PCR-reaction mixtures contained 5 µL TransStart Top Green qPCR SuperMix, 1 µL of template cDNA, 1 µL forward and reverse primers, and 3 µL of sterile distilled water. The thermal cycling conditions comprised an initial denaturation step at 95°C for 2 min, followed by 40 cycles of 95°C for 10 s, 60°C for 30 s. β-actin was used for normalization and 2^-∆∆Ct method was applied for the evaluated mRNA expression of the target gene (27). All the reactions were carried out in triplicate. All evaluations were performed by specialists who were unaware of the group assignment. Primer sequences (forward and reverse) are as follows: Cbp/p300-interacting trans-activator, with Glu/Asp-rich carboxy-terminal domain, 4 (CITED4), acgaggtgttggcagctct and caacctagcagcagagga; CCAAT/enhancer-binding protein beta (C/EBPβ), caacctagcagcagaggaac and acagctgtccaccttcct; and β-actin, cggtggtctcatcacttegg and atggccagaggcttacacc. The present study demonstrates that both moderate intensity (MIIT, 65% - 70% of VO2max) and high intensity (HIIT, 85% - 90% of VO2max) ET for up to six weeks can improve cardiac function capacity as measured by ejection fraction and fractional shortening in infarcted heart. As predicted, the HIIT exercise-trained group demonstrated the highest increase. Also six weeks of ET significantly up-regulated mRNA levels of CITED4 and C/EBPβ genes. These genes are known to induce and inhibit cardiomyocyte proliferation (CITED4 and C/EBPβ) following exercise. HIIT animals significantly diminished mRNA levels C/EBPβ compared to MI-Sed. The mRNA level of CITED4 in HIIT and MIIT was significantly more compared to MI-Sed.

4. Results

Echocardiographic data - from Sham and MI-operated animals at four and 10 weeks after surgery - are presented in Table 1. Four weeks after surgery, we compared echocardiographic data between Sham and MI-operated animals. This comparison showed that fractional shortening, ejection fraction, IVSD, and LVPWs decreased dramatically four weeks after MI induction. In contrast, LVIDs increased significantly in MI-operated compared with Sham (P < 0.05). As found in Table 1, and additionally outlined in Figure 1A, in comparison with sham group, an enlarged ventricular cavity and thinned myocardium was seen in MI-operated group. Ten weeks after surgery, ejection fraction in HIIT and MIIT animals and fractional shortening in HIIT animals increased significantly compared to MI-sed animals. In addition, MI-operated animals (MI-sed, LIIT, MIIT, and HIIT) had significantly lower ejection fraction and fractional shortening compared to Sham animals. Also, LVIDd in HIIT animals and LVIDs in MI-sed, LIIT, and HIIT animals increased significantly compared to MI-sed animals (P < 0.05). As shown in Figure 1B, HIIT exercise-trained group showed the greatest increases in ejection fraction and fractional shortening after MI. These progressions were joined by a 32% and 53% increase in ejection fraction and fractional shortening compared to MI-sed, respectively (Table 1).

In the present study, we have evaluated the expression of mRNA levels of the CITED4 and C/EBPβ genes. These genes are known to induce and inhibit cardiomyocyte proliferation (CITED4 and C/EBPβ) following exercise. HIIT animals significantly diminished mRNA levels C/EBPβ compared to MI-Sed. The mRNA level of CITED4 in HIIT and MIIT was significantly more compared to MI-Sed.

5. Discussion

The present study demonstrates that both moderate intensity (MIIT, 65% - 70% of VO2max) and high intensity (HIIT, 85% - 90% of VO2max) ET for up to six weeks can improve cardiac function capacity as measured by ejection fraction and fractional shortening in infarcted heart. As predicted, the HIIT exercise-trained group demonstrated the highest increase. Also six weeks of ET significantly up-regulated mRNA levels of CITED4 and down-regulated mRNA level of C/EBPβ. Collectively, data from this study suggests that higher intensities of ET are the most effective agents in increasing cardiac function and regeneration capacity in an infarcted heart.

Growing evidences have demonstrated that ET usefully adjusts the clinical course of post MI myocardial remodeling and enhances cardiovascular capacity (28, 29). Post-
MI training may be associated with weakened ventricular hypertrophy and contractile decay, increased maximal cardiac output, ameliorated fibrosis, and an improvement in the quality of life in these patients (28, 30). Our study also confirmed previous findings that ET for up to six weeks improved cardiac function and cardiac regeneration capacity markers.

Exercise applies its gainful impacts by not simply decreasing the weight of established cardiovascular risk factors, but also straightforwardly influencing the cellular and molecular structure and performance of the heart (11, 18). Various studies demonstrated that ET may stimulate proliferation of preexisting cardiomyocytes and resident cardiac stem cells in vivo (11, 18, 26, 31, 32). Recently, a study proposed that four weeks of ET on myocardial infarcted rats stimulates cardiomyocytes proliferation, hinders cardiomyocytes apoptosis, and enhances cardiac function and remodeling (33).

In confirming these results, our study indicated that six weeks of ET induced markers of CITED4 and C/EBPβ. Also, our result demonstrated that higher intensity of ET (MIIT and HIIT) leads to further up and down-regulation markers of CITED4 and C/EBPβ compared to MI-sed. These results were similar to Waring’s et al. study, which showed high intensity exercise trained animals had the greatest number of newly formed cardiomyocytes after four weeks (26). Various mechanisms and signaling pathways regulate cardiac regeneration which is an intricate behavior (33). In this regard, Bostrom et al. reported that C/EBPβ stifles cardiomyocyte development and proliferation in...
Table 1. Echocardiographic Data from MI-Operated and Sham Rats 4 and 10 Weeks After Surgery

|                      | Sham, N = 6 | MI-Operated, N = 24 |
|----------------------|-------------|---------------------|
| **Four Weeks After Surgery** |             |                     |
| IVSD (cm)            | 0.17 ± 0.04 | 0.14 ± 0.03b        |
| IVSS (cm)            | 0.27 ± 0.04 | 0.20 ± 0.04         |
| LVIDd (cm)           | 0.50 ± 0.11 | 0.55 ± 0.09         |
| LVIDs (cm)           | 0.22 ± 0.07 | 0.40 ± 0.07b        |
| LVPWd (cm)           | 0.35 ± 0.05 | 0.17 ± 0.07         |
| LVPWs (cm)           | 0.23 ± 0.04 | 0.19 ± 0.03b        |
| EF (%)               | 90.49 ± 2.44| 58.30 ± 10.79b      |
| FS (%)               | 55.40 ± 3.94| 27.06 ± 6.58b       |
| **10 Weeks After Surgery** |             |                     |
| IVSD (cm)            | 0.17 ± 0.02 | 0.16 ± 0.02         |
| IVSS (cm)            | 0.27 ± 0.03 | 0.26 ± 0.05         |
| LVIDd (cm)           | 0.51 ± 0.11 | 0.63 ± 0.07         |
| LVIDs (cm)           | 0.39 ± 0.05b| 0.32 ± 0.09         |
| LVPWd (cm)           | 0.37 ± 0.02 | 0.14 ± 0.02         |
| LVPWs (cm)           | 0.22 ± 0.04 | 0.21 ± 0.07         |
| EF (%)               | 91.59 ± 3.30| 72.87 ± 8.95b       |
| FS (%)               | 56.44 ± 3.74| 37.72 ± 8.11b       |

Abbreviations: EF, left ventricular ejection fraction; FS, left ventricular fractional shortening; IVSD and IVSS, interventricular septal end diastolic and end systolic; LVIDd and LVIDs, left ventricular internal diameter end diastolic and end systolic; LVPWd and LVPWs, left ventricular posterior wall end diastolic and end systolic.

a Values are expressed as mean ± SD.

b P < 0.05 vs. Sham.

c P < 0.05 vs. MI-Sed.

the grown-up mammalian heart and decreases C/EBPβ in swim trained mice. This seems to be an important signal of physiological cardiomyocyte hypertrophy and proliferation (31). Our results demonstrated the down-regulation of mRNA expression of C/EBPβ in HIIT animals compared to MI-sed. The impact of exercise on cardiomyocyte hypertrophy and proliferation includes decrease in the expression of C/EBPβ, and a connected increment in the expression of CITED4. Down-regulation of C/EBPβ, in turn, releases serum response factor (SRF) to bind target gene promoters. They are also involved in the activation of an exercise gene set (GATA-4, Tbx5, Nkx2.5, and Mef2c), and ultimately cardiac growth. According to results of previous and present studies, the reduction in C/EBPβ expression is sufficient to improve cardiac function and up-regulation of cardiomyocyte hypertrophy and proliferation markers after MI (31). Taken together, these data indicate that higher intensities of ET-induced cardiac regeneration following MI.

5.1. Conclusions

In conclusion, we demonstrate that cardiovascular functions are ET intensity-dependent, following MI. It seems that higher intensities of ET are most effective agents in increasing cardiac regeneration markers (CITED4 and C/EBPβ) in MI rats. Therefore, ET intensity has emerged as a crucial variable in future preclinical and clinical studies.

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Footnotes

Conflict of Interests: None declared.
Ethical Considerations: All animal procedures were performed in accordance with the Ethics Committee of Rajaee Cardiovascular Medical and Research Center (RHC.AC.IR.REC.1393.28).

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