


Cardiovascular Protection Variables Based on Exercise Intensity in Stable Coronary Heart Disease Patients After Coronary Stenting: A Comparative Study

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Purpose: Our study aimed at determining and comparing the mechanism of cardiovascular protection variables in moderate-intensity continuous training (MICT) and high-intensity interval training (HIIT) in patients with stable coronary heart disease (CHD) after coronary stenting.

Participants and Methods: This experimental study used the same subject and cross-over design, involving eleven stable CHD patients after coronary stenting. These were randomly divided into two groups; MICT for 29 minutes at 50–60% heart rate reserve and HIIT with 4x4 minute intervals at 60–80% heart rate reserve, each followed by three minutes of active recovery at 40–50% heart rate reserve. These were conducted three times a week for two weeks. The participants' levels of adrenaline, noradrenaline, endothelial nitric oxide synthase (eNOS), extracellular superoxide dismutase (EC-SOD) activity assayed, and flow-mediated dilatation (FMD) were examined before and after treatments were completed.

Results: The HIIT significantly increased the levels of noradrenaline and eNOS compared with MICT ($p < 0.05$). Also, HIIT was better in maintaining EC-SOD activity and FMD compared with MICT ($p < 0.05$). Through the noradrenalin pathway, HIIT had a direct and significant effect on eNOS and FMD ($p < 0.05$) but MICT, through the noradrenaline pathways, had a direct and significant effect on eNOS ($p < 0.05$), and through the EC-SOD activity pathways had a direct and significant effect on FMD ($p < 0.05$). MICT reduced EC-SOD activity and also decreased the FMD value.

Conclusion: HIIT is superior to MICT in increasing cardiovascular protection by increasing the concentrations of noradrenalin and eNOS, maintaining EC-SOD activity, and FMD in stable CHD patients after coronary stenting.

Keywords: coronary heart disease, high-intensity interval training, catecholamine, eNOS, SOD, FMD

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Introduction

Coronary heart disease (CHD) is currently the leading cause of death worldwide.¹ However, regular exercise is considered as one of the preventive and therapeutic measures for CHD.² High-intensity interval training (HIIT) has some positive effects on the heart such as increasing the blood catecholamine concentration, shear stress, and relative hypoxia.^{3,4} It also increases the activities of sympathetic nerve, which stimulate the production of higher levels of catecholamine such as

adrenaline and noradrenaline, compared with moderate-intensity continuous training (MICT).⁵ Additionally, increase in the concentration of catecholamine hormone due to HIIT activates β 2-adrenergic receptors (β 2-ARs) and β 3-adrenergic receptors (β 3-ARs) which induce the production of nitric oxide (NO) in coronary and peripheral blood vessels, through endothelial nitric oxide synthase (eNOS) activation and expression.^{6,7} Enhancement of the expression and activity of extracellular superoxide dismutase (EC-SOD) also increase the concentration of NO.⁸ However, this increase in the expression of NO causes a vasodilating effect on the coronary arteries, shown through flow-mediated dilatation (FMD) examination.⁹ Based on this background, the purpose of this experimental study was to determine and compare the mechanism of cardiovascular protection variables in HIIT and MICT patients with stable coronary heart disease (CHD) after coronary stenting.

Participants and Methods

Participants

Based on eligibility, 11 participants were enrolled in our study in September 2017, which was completed in December 2018. All the participants had clinical evidence of stable CAD with significant stenosis treated with a percutaneous coronary intervention (PCI) at the Department of Cardiology, National Cardiovascular Center, Harapan Kita, and Siloam Hospital, TB Simatupang, Jakarta, and were screened after successful PCI, defined as complete revascularization.

The inclusion criteria include; age <60 years, left ventricular ejection fraction (LVEF) > 50% (measured within six months of enrolment by echocardiography), sinus rhythm, resting heart rate <100 beats/minute, and willingness to sign the informed consent form.

Then, the exclusion criteria include; patients with orthopedic or neurological limitations to exercise, acute systemic illness (thyrotoxicosis, electrolyte abnormalities, uncontrolled diabetes mellitus, severe anemia Hb < 9 g/dL) or fever, pregnant, severe cardiac valve diseases, fixed-rate pacemaker or implantable cardioverter-defibrillator device, uncontrolled atrial or ventricular dysrhythmias, recent sudden cardiac death syndrome, creatinine serum > 2.5 mg/dL, mental or physical impairment leading to an inability to exercise adequately.

Also, the study protocol was reviewed and approved by the National Cardiovascular Center, Harapan Kita Institutional Review and Ethics Board (No. LB 01/02/

VII/KEP. 012/2017), in conformation to the Nuremberg Code and the Declaration of Helsinki on the use of human subjects, and written informed consent was obtained from all participants before being enrolled in the study. The characteristics, medical history, and medications of the participants are shown in Table 1.

Exercise Stress Test

A continuous progressive stress test was performed on a treadmill (GE T2100; idsMED, Jakarta, Indonesia) with the aid of Bruce's protocol, one day before and one day after the exercise program. The estimated workload in metabolic equivalents (Mets) and ST segments were collected and entered on-line. Also, prescriptions of the exercise and the target heart rates were determined based on the results of the treadmill test, according to the ability of each participant at that time, which was a day before the training. The MICT and HIIT were determined based on

Table 1 Baseline Characteristics of the Participants

| Patient Characteristics | n = 11 |
|--|-----------------|
| Gender [n (%)] | |
| Male | 8 (72.7) |
| Female | 3 (27.3) |
| Age (mean \pm SD) (years) | 48.5 \pm 6.6 |
| Weight (mean \pm SD) (kg) | 72.4 \pm 9.1 |
| Height (mean \pm SD) (cm) | 164.1 \pm 5.2 |
| BMI (mean \pm SD) (kg/m ²) | 27.0 \pm 3.9 |
| Ejection fraction (mean \pm SD) (%) | 61.5 \pm 5.4 |
| Risk factors [n (%)] | |
| Hypertension | 3 (27.3) |
| Smoking | 9 (81.8) |
| Diabetes mellitus | 1 (9.1) |
| Dyslipidemia | 6 (54.5) |
| Family history | 3 (27.3) |
| Menopause | 3 (27.3) |
| Baseline ECG [n (%)] | |
| Normal | 7 (63.6) |
| Ischemia | 0 |
| Old infarct | 4 (36.4) |
| Medication [n (%)] | |
| Acetylsalicylic acid | 11 (100) |
| Clopidogrel | 9 (81.8) |
| ACE inhibitor/ARB | 8 (72.7) |
| β -blocker | 10 (90.9) |
| Statin | 11 (100) |
| Isosorbide dinitrate | 2 (18.2) |

Abbreviations: BMI, body mass index; ECG, electrocardiogram; ACE, angiotensin-converting enzyme; ARB, angiotensin II receptor blocker.

the Borg rating scale and the participants' heart rate reserve (HRR), was calculated using the formula:¹⁰ Target heart rate = [(Maximum heart rate - Resting heart rate) × % Intensity desired] + Resting heart rate.

Blood Sampling and Analysis

The blood samples were obtained at rest, one day before and one day after the exercise program using a 19-gauge needle by direct vein puncture. This was drawn into 6-mL vacuum tubes containing EDTA at room temperature. Then, the resulting platelet-poor plasma was collected in 1.5-mL Eppendorf tubes and frozen at -80°C for biomarker assays. However, all the samples were drawn and analyzed by blinded technicians. Also, the plasma adrenaline and noradrenaline concentrations were determined by enzyme immunoassay method (Adrenaline/noradrenaline ELISA kit; Catalog Number KA1877, Abnova Corporation, Taipei, Taiwan). The endothelial nitric oxide synthase (eNOS) concentrations were measured on the same plasma samples using the Quantitative Sandwich ELISA method (Human endothelial nitric oxide ELISA kit; Catalog Number MBS030177, MyBiosource, Inc., CA, USA). Then, the extracellular superoxide dismutase (SOD) activity was measured using colorimetric based assay, superoxide ions generated from the conversion of xanthine, and oxygen to uric acid and hydrogen peroxide by xanthine oxidase (SOD activity kit; Catalog Number ADI-900-157, Enzo Life Sciences, Inc., New York, USA).

Flow-Mediated Dilatation

The flow-mediated dilatation (FMD) was determined a day before and a day after the exercise program. The participants were allowed to rest for 10 min before initiating the FMD study in the brachial artery, to ensure stable conditions during scanning. Rapid inflation and pneumatic deflation cuff (big ben[®] sphygmomanometers, Riester, Jungingen, Germany) were positioned on the right forearm, 2 inches below the antecubital fossa. A 10-MHz multi-frequency linear array probes, attached to a high-resolution ultrasound machine (Aloka ProSound Alpha 10, Virginia, USA), was then used to image the brachial artery in the distal third of the upper arm. When an optimal image was obtained, the probe was held stable, and the ultrasound parameters were set to optimize the longitudinal, B-mode image of the lumen-arterial wall interface. After baseline images of the right brachial artery were obtained for 1 minute, the cuff was inflated to 50 mm Hg above the patient's systolic blood pressure to occlude

the right brachial artery and the cuff was kept inflated for 5 minutes. Then, the longitudinal image of the artery was continuously recorded from 30 s before to 2 min after cuff deflation.¹¹ Images of the brachial artery diameters were captured in diastole (gated with electrocardiogram R wave). The FMD is calculated as a percentage change of peak diameter with respect to reactive hyperemia baseline diameter, as shown by this equation:¹²

$$\text{FMD(\%)} = \frac{\text{Peak diameter} - \text{baseline diameter}}{\text{Baseline diameter}} \times 100\%$$

Study Protocol

The study subjects were all receiving different treatments at certain time intervals. Also, the cross-over design was used to avoid the residual effects of treatment. The study subjects were 11 stable CHD patients after the implantation of coronary artery stents who met the inclusion criteria, each of which received two treatments; the HIIT and the MICT programs with the same total workload.

All the participants were asked to be in good hydration condition, get enough sleep to avoid fatigue, fever or flu, take the last meal at least two hours before exercise, continue taking their medications, stop taking alcohol and avoid smoking before partaking in the exercise. Additionally, the participants were instructed not to get involved in additional exercise during the study period. Then, the physical examination, measurement of vital signs, and resting ECG recording were carried out before the exercise. Only participants with physical stable condition such as no new complaints, systolic blood pressure <160 mmHg, diastolic blood pressure <100 mmHg, resting heart rate <100 beats/min, respiratory <25 times/min, body temperature 36–37°C, and no change from the ECG compared to the baseline ECG at the beginning of the examination, were allowed to start the exercise.

The treadmill test procedures were performed before each training to determine its prescriptions, based on the resting and maximum heart rates, according to each patient's ability. In addition, the treadmill tests were conducted to examine participants' aerobic capacity before and after exercise program, although not shown in the results of our study. Examinations of the blood for adrenaline, noradrenaline, eNOS, EC-SOD activity, and FMD were also conducted before and after the patient followed the specified training program.

The participants met for training three times weekly for two weeks, amounting to six sessions, for each training

program, under the supervision of a physician, trained nurse, and a trainer. Based on previous studies, daily MICT sessions for about 6–10 days could increase the aerobic capacity and mitochondrial enzyme activity of the participants, while seven sessions of HIIT for two weeks could increase subjects' skeletal muscle capacity for fatty acid oxidation during the exercise.^{13,14}

The training was conducted using a motorized treadmill (Matrix MX-T5x, Wisconsin, USA) where participants walked on it with no inclination. The MICT group went through a 5 minute warm-up period at an intensity corresponding to 40% of HRR before walking for 29 minutes continuously at an intensity 40–60% of HRR. The training session ended with a 3 minute cooling down period at 40% of HRR. Similarly, the HIIT group went through a 5 minute warm-up period at an intensity of 40%, followed by walking on the treadmill four times with 4 minute interval at 60–80% with three active recovery of 3 minute walking at 40–50%, and a 3 minute cool down at 40% of HRR. Each participant had the exercise with an electrocardiography (ECG) telemetry and a heart rate monitoring device (Nihon Kohden ECG Gateway, Tokyo, Japan). Also, the speed of the treadmill was continually adjusted as the training progressed to ensure that all sessions were carried out at the desired heart rate throughout the period. The BORG 6–20 scale was used to measure the rate of perceived exertion after each training session.¹⁵

In addition, the participants were detrained for two weeks after going through the first training to eliminate the effects of the previous exercise. Furthermore, each participant was subjected to a physical exercise different from the previous one; the participants who previously received MICT got HIIT, and vice versa.

The flow of participants through each period of the training program is shown in [Figure 1](#).

Statistical Analysis

Participants' characteristics were subjected to descriptive statistical analysis, with the categorical variables expressed as percentages and continuous variables expressed as mean \pm standard deviation (SD). Non-parametric statistics were used due to the small groups involved in the study and to avoid assumptions of normal distributions. Also, changes within each parameter were assessed using the Wilcoxon signed ranks test with SPSS software, version 23.0 (IBM, New York, USA).

Additionally, path analysis was conducted to investigate the effects of each variable, which made it possible

for simultaneous conduction of multiple and linear regression analyses. Therefore, it permitted the specification of complex models with multiple variable influences.¹⁶ Path analyses were conducted with AMOS software, version 25.0 (IBM, New York, USA) and the level of significance was set at $p < 0.05$ for all tests.

Results

The average age of the participants was 48.5 ± 60.6 -years, majority of whom had normal ECG, overweight, and smoking as the most common risk factors for CHD. The average left ventricular ejection fraction based on echocardiography was $61.5 \pm 5.4\%$, and all participants received optimal medical therapy, as shown in [Table 1](#).

Eleven participants completed the two weeks of MICT or HIIT sessions and the average heart rate during the period is shown in [Table 2](#). The HIIT group did not show any statistical significant differences in adrenaline concentration compared to the MICT group, after four weeks of training, as shown in [Table 3](#). There was a significant increase of noradrenalin in the HIIT group compared with MICT at $p < 0.05$; as shown in [Table 4](#). However, there was a significant decrease in EC-SOD activity in the MICT group compared with HIIT at $p < 0.05$; as shown in [Table 6](#). Also, the concentration of eNOS significantly varied before and after HIIT at $p < 0.05$; as shown in [Table 5](#), while a greater improvement was observed in FMD of the HIIT group compared with MICT at $p < 0.05$; as shown in [Table 7](#).

[Table 8](#) shows the standardized regression coefficients of the initial result of path analysis extracted for MICT and HIIT; models from AMOS software shown in [Figures 2A](#) and [3A](#). Positive coefficients mean that the increase in an independent variable leads to a rise in a dependent variable and vice versa. The initial path analysis showed that adrenaline concentration did not affect eNOS concentration since $p > 0.05$, and did not affect FMD as well, since $p > 0.05$. However, the concentration of noradrenaline significantly affected eNOS since $p < 0.001$, with a positive coefficient; but the concentration of noradrenaline did not affect FMD since $p > 0.05$. Similarly, level of eNOS did not have any effect on FMD since $p > 0.05$, but the activity of EC-SOD played a significant role on FMD at $p < 0.05$, with a positive coefficient.

Results in the HIIT group showed that adrenaline concentration did not affect eNOS and FMD since $p > 0.05$ in both cases. Noradrenalin concentration, on the other hand, influenced eNOS since $p < 0.001$, with a positive

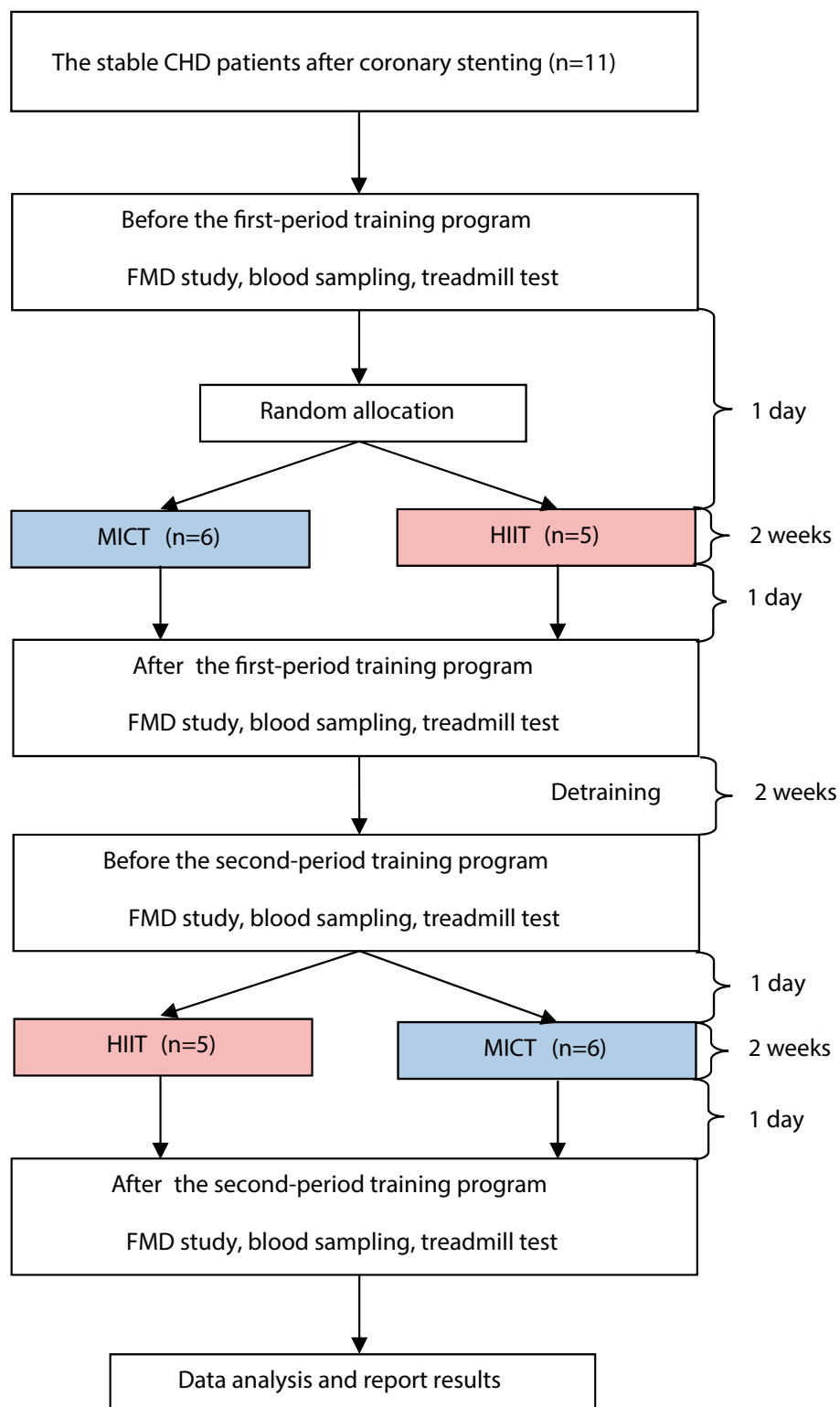


Figure 1 Flow of participants through each period of training program.

Abbreviations: MICT, moderate-intensity continuous training; HIIT, high-intensity interval training; FMD, flow-mediated dilatation.

coefficient; however, it did not affect FMD since $p > 0.05$. Furthermore, eNOS concentration and the activity of EC-SOD did not affect FMD since $p > 0.05$.

Table 9 summarizes the standardized regression coefficients after non-significant paths in MICT and HIIT were deleted; the final path models are shown in

Table 2 The Average of Participants Achieved Improvements in Heart Rate During the Training Program

| Participants | MICT | | | HIIT | | | | |
|--------------|------------------------|---------------------------|-------------------------|------------------------|---------------------------|--|--|---|
| | HR Resting (Mean ± SD) | Target HR (40–60% of HRR) | HR Achieved (Mean ± SD) | HR Resting (Mean ± SD) | Target HR (60–80% of HRR) | High-Intensity HR Achieved (Mean ± SD) | Active Recovery Target HR (40–50% HRR) | Active Recovery HR Achieved (Mean ± SD) |
| 01 | 66.67 ± 5.92 | 87–89 | 86.83 ± 1.17 | 70.50 ± 5.89 | 83–88 | 87.17 ± 1.72 | 78–81 | 79.50 ± 1.05 |
| 02 | 68.83 ± 7.08 | 99–105 | 99.17 ± 4.58 | 69.00 ± 4.60 | 96–107 | 98.33 ± 5.61 | 86–91 | 89.17 ± 1.72 |
| 03 | 86.00 ± 7.97 | 115–119 | 116.83 ± 1.33 | 88.50 ± 5.54 | 114–123 | 117.17 ± 3.19 | 105–110 | 107.17 ± 2.48 |
| 04 | 71.83 ± 4.88 | 84–88 | 83.33 ± 1.03 | 68.00 ± 5.18 | 88–98 | 92.83 ± 3.82 | 80–84 | 82.00 ± 2.10 |
| 05 | 68.33 ± 3.50 | 86–90 | 90.00 ± 2.83 | 69.50 ± 4.37 | 92–99 | 97.17 ± 1.72 | 84–88 | 86.67 ± 1.63 |
| 06 | 78.17 ± 8.47 | 109–112 | 109.67 ± 0.82 | 82.50 ± 4.64 | 99–106 | 103.50 ± 3.21 | 92–96 | 93.33 ± 1.51 |
| 07 | 82.00 ± 7.35 | 96–100 | 99.83 ± 2.40 | 78.33 ± 5.61 | 99–109 | 105.67 ± 6.47 | 90–94 | 92.50 ± 2.74 |
| 08 | 56.50 ± 1.97 | 105–115 | 114.33 ± 0.52 | 54.83 ± 5.34 | 120–141 | 121.17 ± 7.88 | 99–110 | 107.33 ± 4.32 |
| 09 | 78.50 ± 6.69 | 118–126 | 119.67 ± 2.07 | 70.50 ± 8.22 | 126–139 | 126.00 ± 6.66 | 113–119 | 116.33 ± 3.01 |
| 10 | 74.67 ± 3.39 | 109–113 | 109.83 ± 5.53 | 83.83 ± 7.94 | 116–127 | 122.00 ± 2.00 | 105–110 | 108.17 ± 2.14 |
| 11 | 86.67 ± 3.08 | 92–96 | 95.50 ± 2.35 | 84.17 ± 5.88 | 105–114 | 111.00 ± 3.52 | 95–100 | 95.83 ± 1.47 |

Note: Data are expressed as beats per minute.

Abbreviations: MICT, moderate-intensity continuous training; HIIT, high-intensity interval training; HR, heart rate; HRR, heart rate reserve; SD, standard deviation.

Table 3 Changes in Adrenaline Concentration After and Before MICT and HIIT

| Adrenaline (ng/mL) | Before | After | P-value (One-Tailed) | Δ After-Before |
|---------------------|---------------|---------------|----------------------|----------------|
| MICT | 0.022 ± 0.014 | 0.027 ± 0.015 | 0.316 | 0.005 ± 0.018 |
| HIIT | 0.023 ± 0.013 | 0.032 ± 0.024 | 0.080 | 0.008 ± 0.019 |
| p-value(one-tailed) | 0.380 | 0.337 | | 0.480 |

Note: Data are expressed as mean ± SD.

Abbreviations: MICT, moderate-intensity continuous training; HIIT, high-intensity interval training; SD, standard deviation.

Table 4 Changes in Noradrenaline Concentration After and Before MICT and HIIT

| Noradrenaline (ng/mL) | Before | After | P-value (One-Tailed) | Δ After-Before |
|-----------------------|---------------|---------------|----------------------|----------------|
| MICT | 0.070 ± 0.131 | 0.038 ± 0.082 | 0.343 | −0.032 ± 0.146 |
| HIIT | 0.064 ± 0.120 | 0.160 ± 0.214 | 0.014* | 0.096 ± 0.143 |
| p-value(one-tailed) | 0.297 | 0.014* | | 0.014* |

Note: Data are expressed as mean ± SD. *Indicates statistical significance (p<0.05).

Abbreviations: MICT, moderate-intensity continuous training; HIIT, high-intensity interval training; SD, standard deviation

Table 5 Changes in eNOS Concentration After and Before MICT and HIIT

| eNOS (ng/mL) | Before | After | P-value (One-Tailed) | Δ After-Before |
|----------------------|---------------|---------------|----------------------|----------------|
| MICT | 1.309 ± 0.797 | 1.235 ± 0.350 | 0.267 | −0.074 ± 0.702 |
| HIIT | 1.249 ± 0.507 | 1.302 ± 0.478 | <0.001* | 0.053 ± 0.184 |
| p-value (one-tailed) | 0.380 | 0.212 | | 0.439 |

Notes: Data are expressed as mean ± SD. *Indicates statistical significance (p<0.001).

Abbreviations: eNOS, endothelial nitric oxide synthase; MICT, moderate-intensity continuous training; HIIT, high-intensity interval training; SD, standard deviation.

Figures 2B and 3B. The final path analysis showed that in MICT, noradrenalin concentration affected eNOS concentration since p<0.001, and EC-SOD activity affected eNOS since p<0.05, with a positive coefficient.

Results in the HIIT group showed that noradrenaline concentration affected eNOS since p<0.001, with a positive coefficient, and affected FMD since p<0.05, but with a negative coefficient.

Table 6 Changes in EC-SOD Activity After and Before MICT and HIIT

| EC-SOD Activity (units/ μ L) | Before | After | P-value (One-Tailed) | Δ After-Before |
|----------------------------------|-------------------|-------------------|----------------------|-----------------------|
| MICT | 0.378 \pm 0.047 | 0.362 \pm 0.046 | 0.023* | -0.016 \pm 0.024 |
| HIIT | 0.357 \pm 0.034 | 0.368 \pm 0.037 | 0.229 | 0.010 \pm 0.044 |
| p-value (one-tailed) | 0.035* | 0.378 | | 0.048* |

Notes: Data are expressed as mean \pm SD. *Indicates statistical significance ($p < 0.05$).

Abbreviations: EC-SOD, extracellular superoxide dismutase; MICT, moderate-intensity continuous training; HIIT, high-intensity interval training; SD, standard deviation.

Table 7 Changes in FMD After and Before MICT and HIIT

| FMD (%) | Before | After | P-value (One-Tailed) | Δ After-Before |
|----------------------|--------------------|--------------------|----------------------|-----------------------|
| MICT | 10.555 \pm 4.436 | 10.009 \pm 3.994 | 0.254 | -0.546 \pm 3.849 |
| HIIT | 8.809 \pm 2.744 | 9.664 \pm 3.452 | 0.324 | 0.855 \pm 3.645 |
| p-value (one-tailed) | 0.133 | 0.439 | | 0.042* |

Notes: Data are expressed as mean \pm SD. *Indicates statistical significance ($p < 0.05$).

Abbreviations: FMD, flow-mediated dilatation; MICT, moderate-intensity continuous training; HIIT, high-intensity interval training; SD, standard deviation.

The final diagram of the cardiovascular protection mechanism for MICT and HIIT are shown in Figure 4. MICT reduced EC-SOD activity and the lower the activity, the lower the FMD assay. The higher the noradrenaline in the MICT group, the higher the concentration of eNOS. However, the HIIT further increased noradrenaline and eNOS concentrations compared with MICT. Hence, the higher the noradrenaline concentration in HIIT, the higher the eNOS and the lower the FMD assay, and vice-versa.

Safety Assessment

There was no adverse event in any of the groups.

Table 8 The Initial Results of Path Analysis for MICT and HIIT

| Model | MICT | | HIIT | |
|-----------------------------------|---------|----------|---------|----------|
| | β | p-value | β | p-value |
| Adrenaline \rightarrow eNOS | 0.136 | 0.388 | 0.122 | 0.487 |
| Noradrenaline \rightarrow eNOS | 0.680 | <0.001** | 0.585 | <0.001** |
| Adrenaline \rightarrow FMD | -0.236 | 0.191 | 0.284 | 0.097 |
| Noradrenaline \rightarrow FMD | -0.087 | 0.721 | -0.346 | 0.099 |
| eNOS \rightarrow FMD | -0.081 | 0.742 | -0.304 | 0.151 |
| EC-SOD Activity \rightarrow FMD | 0.503 | 0.005* | 0.036 | 0.833 |

Notes: Single-headed arrows (\rightarrow) indicate a path. β indicate standardized regression coefficients. *Indicates statistical significance ($p < 0.05$). **Indicates statistical significance ($p < 0.001$).

Abbreviations: MICT, moderate-intensity continuous training; HIIT, high-intensity interval training; eNOS, endothelial nitric oxide synthase; EC-SOD, extracellular superoxide dismutase; FMD, flow-mediated dilatation.

Discussion

Based on the result, there was no significant difference in the change in adrenaline concentration in both groups. Stimulation of the sympathetic nerves increased during exercise with higher intensity, thereby stimulating the adrenal medulla gland to produce adrenaline hormone. The concentration of this hormone in plasma is also influenced by its clearance but it reduces when the concentration of adrenaline increases more than ten times its basal value.¹⁷

This clearance is reduced due to HIIT causing high secretion of adrenaline, which requires a longer time to be eliminated from the body. During MICT also, the clearance of adrenaline concentration is reduced due to its distribution in the organs being more evenly distributed due to the longer duration of exercise (29 minutes). Hence, it takes a long time to eliminate adrenaline in the plasma. Also, there was no significant increase in adrenaline concentrations, presumably because its concentration was more related to psychological stress, hypoglycemia, caffeine consumption, and Valsalva maneuver, compared to noradrenaline concentrations, which are more related to physical exercise and body position (orthostatic).¹⁸

The results showed that the level of noradrenaline increased more with HIIT compared with MICT activities. Similarly, a study by Greiwe et al, showed that exercise on a treadmill for 15 minutes in 6 women and 3 men aged 28 years, resulted in higher noradrenaline at rest compared to post-exercise, with an intensity of 60% VO_{2max} . However,

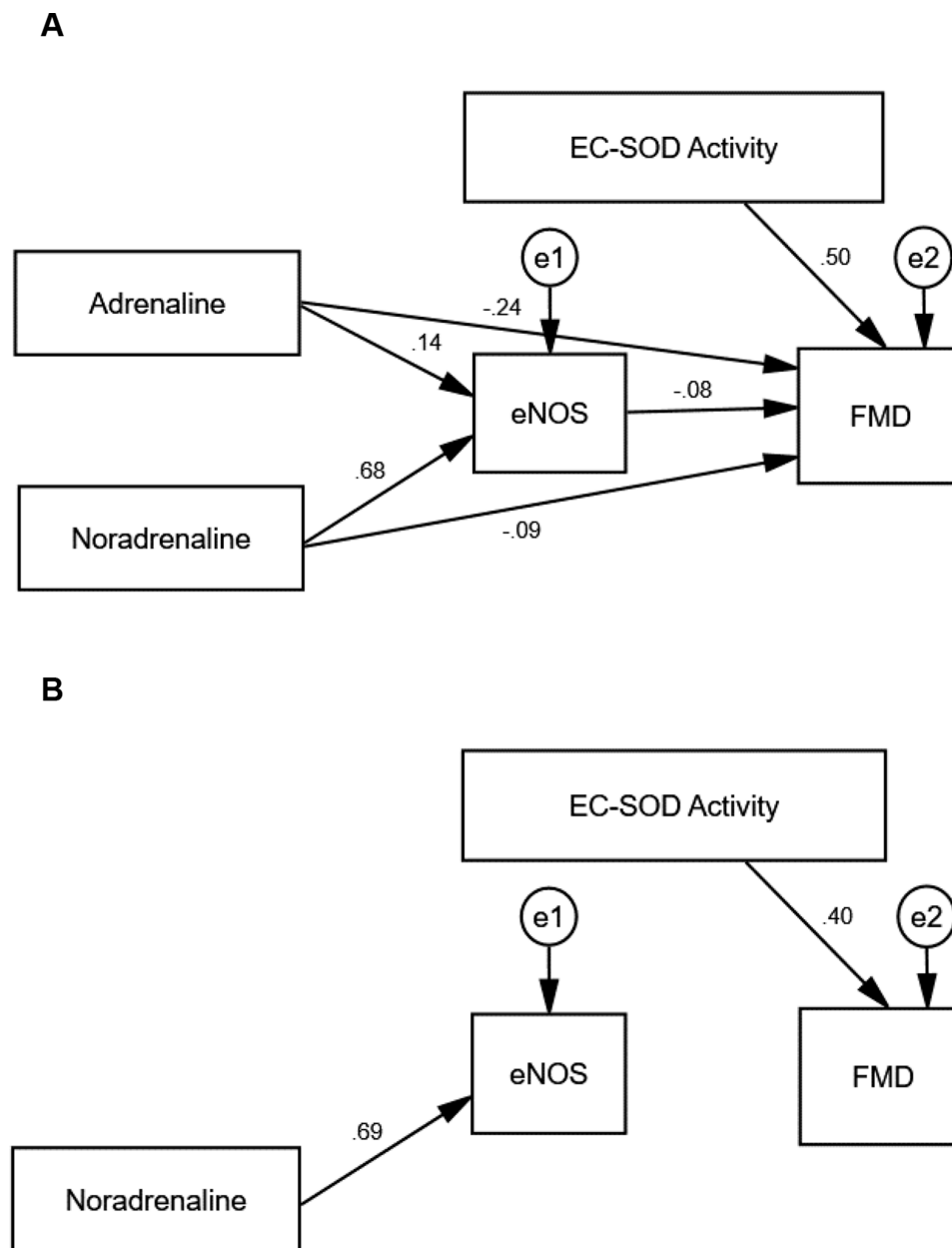


Figure 2 The path model of MICT.

Notes: (A) The hypothesized model with path coefficients of MICT. (B) The trimmed model with path coefficients of MICT. All variables (dependent and independent) are shown in rectangles. Variables not included in the models are shown in circles with (e) sign. Single-headed arrows with standardized coefficients link independent variables to the dependent variable. The diagrams are plotted and optimized in Amos (version 25.0).

Abbreviations: eNOS, endothelial nitric oxide synthase; EC-SOD, extracellular superoxide dismutase; FMD, flow-mediated dilatation.

the noradrenaline concentration increased when the intensity of the exercise was within 60–85% VO_{2max} .¹⁹ In our study, the intensity of MICT was within 50–60% HRR, which is equivalent to 50–60% VO_{2max} , and the intensity of HIIT was within 60–80% HRR (60–80% VO_{2max}), which was almost similar to this previous study. Our study assumes that the higher the intensity of exercise, the stimulation of the sympathetic nervous system

increases, thereby resulting in higher noradrenaline. Likewise, Greiwe et al demonstrated the increment in the spillover of noradrenaline during moderate to high-intensity exercise, and increased continuously with time, while the systemic noradrenaline clearance reduced.¹⁹

According to Padilla et al, there was a significant relationship between the intensity of aerobic exercise training and the flow-mediated dilatation response. The

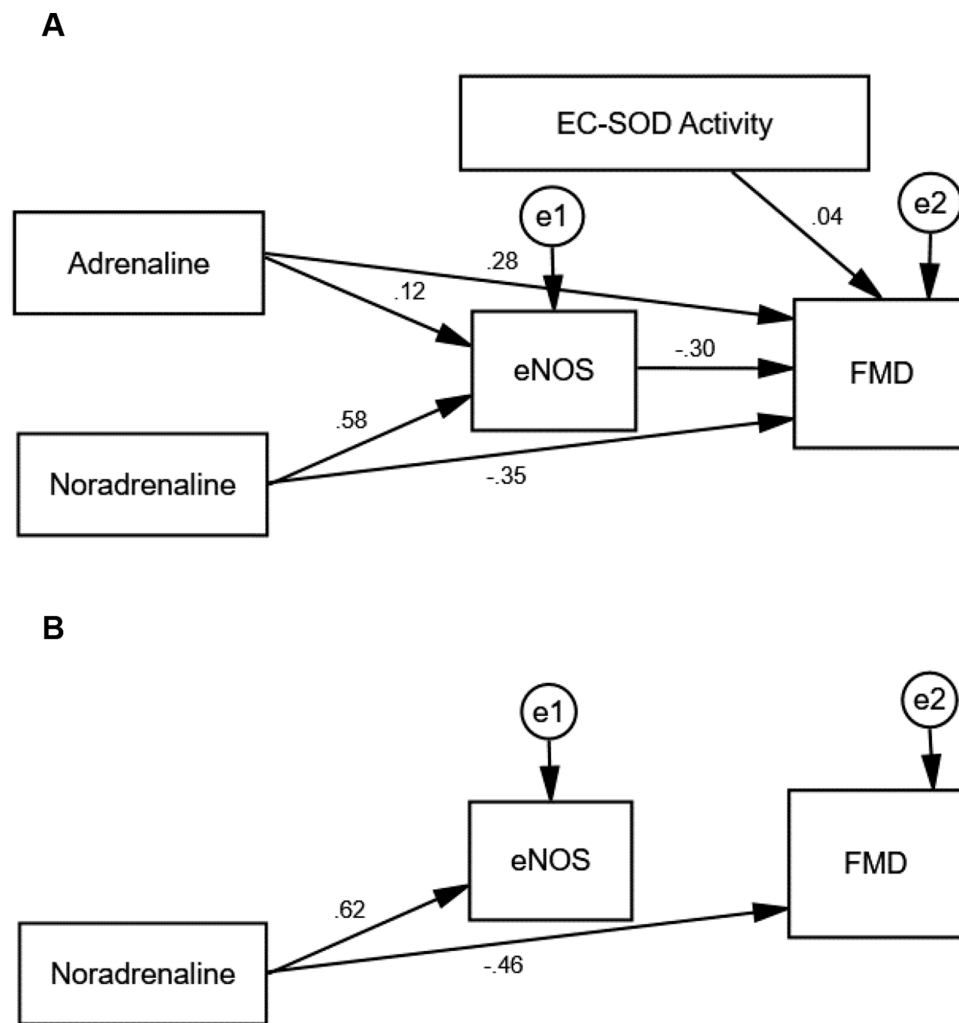


Figure 3 The path model of HIIT.

Notes: (A) The hypothesized model with path coefficients of HIIT. (B) The trimmed model with path coefficients of HIIT. All variables (dependent and independent) are shown in rectangles. Variables not included in the models are shown in circles with (e) sign. Single headed arrows with standardized coefficients link independent variables to dependent variable. The diagrams are plotted and optimized in Amos (version 25.0).

Abbreviations: eNOS, endothelial nitric oxide synthase; EC-SOD, extracellular superoxide dismutase; FMD, flow-mediated dilatation.

study showed that the brachial artery shear stress was greatest following high-intensity exercise.²⁰ In vivo exercise has a similar effect as laminar shear stress on

endothelial function. Normally, laminar shear stress regulates endothelial structure and function by controlling the expression of mechanosensitive genes and production of vasoactive factors. This shear stress affects eNOS acutely and chronically, thereby activating the enzyme to produce NO within seconds. It also stimulates intracellular calcium, increases calmodulin to bond with eNOS, and increases eNOS activity acutely and transiently.^{21,22} In addition, shear stress stimulates the phosphorylation of eNOS at serine 635 and 1177 by protein kinase-A. Laminar shear stress which occurs for long time increases eNOS messenger ribonucleic acid (mRNA) and protein; the eNOS expression therefore increases due to increased transcription of temporary eNOS mRNA and sustained improvement in eNOS mRNA stability.²²

Table 9 The Final Results of Path Analysis for MICT and HIIT

| | Model | β | p-value |
|------|------------------------------------|---------|----------|
| MICT | Noradrenaline \rightarrow eNOS | 0.692 | <0.001** |
| | EC-SOD Activity \rightarrow eNOS | 0.404 | 0.043* |
| HIIT | Noradrenaline \rightarrow eNOS | 0.616 | <0.001** |
| | Noradrenaline \rightarrow FMD | -0.456 | 0.019* |

Notes: Single-headed arrows (\rightarrow) indicate a path. β indicate standardized regression coefficients. *Indicates statistical significance ($p < 0.05$). **Indicates statistical significance ($p < 0.001$).

Abbreviations: MICT, moderate-intensity continuous training; HIIT, high-intensity interval training; eNOS, endothelial nitric oxide synthase; EC-SOD, extracellular superoxide dismutase; FMD, flow-mediated dilatation.

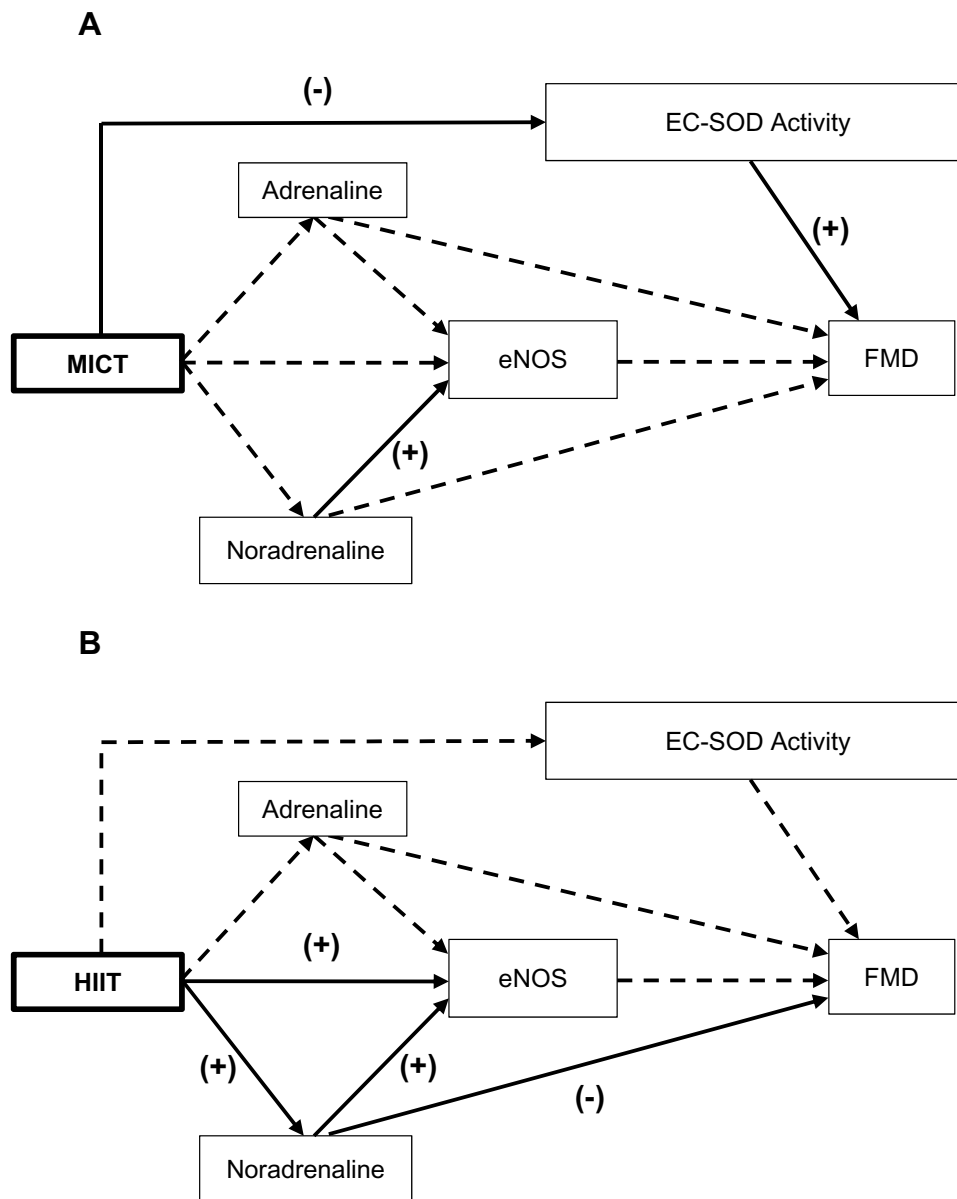


Figure 4 The best diagram of cardiovascular protection mechanism for MICT (A) and HIIT (B).

Notes: All variables (dependent and independent) are shown in rectangles. Arrows represent links, with significant links shown as solid lines and nonsignificant paths shown as broken lines. (+), positive influence; (-), negative influence.

Abbreviations: MICT, moderate-intensity continuous training; HIIT, high-intensity interval training; eNOS, endothelial nitric oxide synthase; EC-SOD, extracellular superoxide dismutase; FMD, flow-mediated dilatation.

The HIIT in our study might result in more significant shear stress in the endothelial layer, thereby producing a higher concentration of eNOS compared with MICT. In general, the concentration of eNOS in plasma is an indication of high or low NO synthesis and also equivalent to an improved endothelial function.²³

MICT was performed three times a week for two weeks in our study, resulting in decreased EC-SOD activity. This has shown the inability to maintain or increase EC-SOD activity. It is assumed that MICT is unable to elicit an

adaptive response to oxidative stress through the activation of nuclear erythroid 2-related factor (Nrf2) signal. Nrf2 is a redox transcription factor and a major regulator of antioxidants. It is also a cytoprotective cofactor responsible for increasing the defense mechanism of an antioxidant.²⁴ Acute exercise could activate Nrf2 signaling through increased reactive oxygen species (ROS) production, thereby increasing the transactivation of antioxidant genes and myocardial cardioprotection.²⁴ According to Gounder et al, increase in the regulation of Nrf2 expression occurs during

high-intensity training.²⁵ However, there were only few studies on the adaptive response of Nrf2 to exercise in humans.²⁶

Landmesser et al showed that EC-SOD activity decreased in CHD patients, which was related to endothelial dysfunction.²⁷ The HIIT in our study was performed three times a week for two weeks in stable CHD patients and was able to maintain the EC-SOD activity, compared with MICT. Acute HIIT could increase oxidative stress, which stimulates ROS, including superoxide anions. However, repeated exercise could cause changes in the regulation of the body's defense system by increasing antioxidants with the capacity of turning pro-oxidants into inactive ones.²⁸

HIIT could create unfavorable conditions with a continuous increase in its intensity, because it could lead to increased ROS and decreased circulating antioxidants.²⁹ This condition could however be prevented with intervals, in the form of active or passive recovery, so as to eliminate its adverse effects on blood vessels.^{29,30}

Also, HIIT could provoke more blood flow in the large blood vessels during exercise where shear stress increases and induces NO bioavailability.³¹ Moreover, repetitive shear stress due to HIIT could initiate changes at the molecular level. The potassium canal of endothelial cells becomes more sensitive, thereby facilitating the influx of calcium into the cells. This increased in intracellular calcium in endothelial cells increases the activity and expression of eNOS, and NO production, hence resulting in vasodilation.³² The HIIT in our study increased the mean change in FMD such that it was better to stay well maintained, compared with MICT. In general, HIIT is more effective in improving brachial artery endothelial function (FMD) compared with MICT.

Furthermore, the path analysis showed that adrenaline concentration in MICT and HIIT has no significant effect on the concentration of eNOS. This was due to changes in the adrenaline which occurred as the exercise has not been able to activate β 2-ARs and β 3-ARs in blood vessel as the stimulation and activation on these receptors could induce eNOS activation. Research conducted by Banquet et al, showed that the stimulation in β 2-ARs could activate eNOS through Src kinase-PI3K/Akt-dependent.³³

Adrenaline affects blood vessel dilatation through its affinity on adrenergic receptors. It has a magnitude of affinity for α 1 adrenergic receptors (α 1-ARs) and α 2 adrenergic receptors (α 2-ARs), which is almost the same as noradrenaline, but with a higher affinity for β 2-ARs and

lower for β 3-ARs compared to noradrenaline. Low adrenaline concentrations results to vasodilation, while high adrenaline causes vasoconstriction.³⁴ The concentration of adrenaline in our study, caused by MICT and HIIT had not significant effect on FMD, presumably because its concentration was not up to the level needed to stimulate adrenergic receptors in blood vessels.

Vasodilation of blood vessels is influenced by several mechanisms, including adenosine through the hyperpolarization of calcium-activated potassium channel membrane, or the endothelium-derived hyperpolarizing factor (EDHF) pathway,³⁵ prostacyclin through the cyclic adenosine monophosphate (cAMP) pathway,³⁶ and NO through the cyclic guanosine monophosphate (cGMP) pathway.³⁷ Since eNOS expression increases the bioavailability of NO, this results in vasodilation of blood vessels which increases FMD. In our study, eNOS had no effect on FMD presumably due to the involvement of other pathways which are more dominant in influencing FMD, such as the EDHF or prostacyclin pathway. Hence, there is need for further research in both exercises.

The noradrenaline concentration influences eNOS concentration in both MICT and HIIT. This is a catecholamine hormone more associated with exercise compared with adrenaline. Circulating noradrenaline usually shows a metabolic effect with concentration more than 1.8 ng/mL.¹⁸ In our study, the average increase in noradrenaline concentration after the MICT and HIIT programs was 0.038 ± 0.082 and 0.160 ± 0.214 ng/mL respectively, which was able to increase the concentration of eNOS. It is assumed that increasing the concentrations of noradrenaline in both exercise were able to stimulate α -ARs and β 2-ARs. Then, the stimulation of these adrenergic receptors resulted in an increase in Akt phosphorylation, which in turn mediates eNOS phosphorylation in Serine 1177, thereby increasing the concentration of eNOS.^{38,39}

The noradrenaline stimulation of β 3-ARs is also thought to be the underlying cause of the increase in eNOS after the exercise. Stimulation of β 3-ARs has the capacity of activating eNOS through the Rac1-PKA-Akt pathway.⁴⁰ The results in our study showed that the increase in the concentration of noradrenaline was more significant after performing HIIT than MICT. Similarly, HIIT increased eNOS concentrations compared to MICT. This is due to the fact that the noradrenaline concentration in HIIT was more significant in stimulating α -ARs, β 2-ARs, and β 3-ARs.

Noradrenaline is a protector against oxidants and known to be a superoxide free radical scavenger, with

the capacity of inducing SOD expression.⁴¹ The increase in noradrenaline concentration, which is more significant in HIIT, and its function as an antioxidant which induces EC-SOD expression, is thought to be the reason why HIIT is more able to maintain EC-SOD activity than MICT.

Our study showed that the noradrenaline concentration of HIIT affected FMD negatively. This means that increase in noradrenaline results in decrease in FMD, and vice-versa. Noradrenaline stimulation of cardiac β 1-ARs is comparable to adrenaline, its effect on peripheral α -ARs is equivalent, but its effect on β 2-ARs for peripheral vasodilation is less than that of adrenaline. Noradrenaline significantly increases vascular resistance and reduces blood flow in the muscles compared with adrenaline.⁴² Although noradrenaline stimulates β 2-ARs, its effect on the vasoconstriction of blood vessels is greater. This is because the anatomical location of α 1-ARs is closer to the point where noradrenaline is released from terminal neurons compared to the location of β 2-ARs. Also, the number and activity of vascular α 1-ARs are greater compared with β -ARs, thereby resulting in vasoconstriction.⁴³ This is thought to be the reason why a significant increase in noradrenaline could reduce the value of FMD.

However, noradrenaline has a greater affinity for β 3-ARs compared to adrenaline.³⁴ The new β 3-ARs is stimulated when the plasma catecholamines are very high. This is considered a “safety valve” when intense adrenergic stimulation occurs at this receptor. In addition, the stimulation of β 3-ARs has a cardioprotective effect on the ventricles and atrium of the heart in the form of a positive feedback mechanism against inotropic and chronotropic which increase due to the increase in this catecholamine hormone. Furthermore, the stimulation of β 3-ARs reduces the incidence of arrhythmias due to the presence of the catecholamine hormone in β 1-ARs.⁴⁴ Thus, it is assumed that a significant increase in the concentration of noradrenaline in HIIT is expected to stimulate β 3-ARs in heart, hence providing a better cardioprotection effect with the capacity of maintaining the balance of oxygen consumption in the heart muscle when there is a high increase in the catecholamine hormone.

Conclusion

Based on the results, HIIT increased the concentrations of noradrenaline and eNOS, as well as maintained the EC-SOD activity and FMD. Although, it was unable show an increase in adrenaline concentration more than MICT. Also, HIIT provided a better effect on cardiovascular protection compared with MICT, as a result of increased

in noradrenaline and eNOS, as well as through its ability to maintain EC-SOD activity and FMD in stable CHD patients after coronary stenting.

However, there is need for future research on the effect of HIIT on cardiovascular protection using other means.

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Disclosure

The authors report no conflicts of interest in this work.

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