Plasma Pentraxin 3 Concentration Increases in Endurance-Trained Men

Miyaki Asako, Maeda Seiji, Otsubi Takeshi, Ajisaka Ryuichi

Journal or Publication Title: Medicine and science in sports and exercise
Volume: 43
Number: 1
Page Range: 12-17
Year: 2011-01

©2011 The American College of Sports Medicine

This is a non-final version of an article published in final form in Medicine & Science in Sports & Exercise: January 2011 - Volume 43 - Issue 1 - pp 12-17

URL: http://hdl.handle.net/2241/115893
doi: 10.1249/MSS.0b013e3181e84bce
Plasma pentraxin3 concentration increases in endurance-trained men

Asako Miyaki¹, Seiji Maeda¹, Takeshi Otsuki², and Ryuichi Ajisaka¹

¹)Division of Sports Medicine, Graduate School of Comprehensive Human Sciences, University of Tsukuba, Tsukuba, Ibaraki, Japan, ²)Graduate School of Health and Sport Sciences, Ryutsu Keizai University, Ryugasaki, Ibaraki, Japan

Running title: Exercise training and PTX3

This work was supported by Grants-in-Aid for Scientific Research 21300234 and 21650179 and Grant-in-Aid for JSPS Fellows 21·692 from Japan Society for the Promotion of Science.

Address for Correspondence:
Seiji Maeda, Ph.D.
Division of Sports Medicine
Graduate School of Comprehensive Human Sciences
University of Tsukuba
Tsukuba, Ibaraki 305-8577
Japan
TEL +81 29-853-2683
FAX +81 29-853-2986
E-mail: maeda@taiiku.tsukuba.ac.jp
ABSTRACT

Background: Pentraxin3 (PTX3), which is mainly produced by endothelial cells, macrophages, and smooth muscle cells in the atherosclerotic region, has a cardioprotective effect. Endurance exercise training has also been known to offer cardioprotection. However, the effect of regular endurance exercise on PTX3 is unknown. This study aimed to investigate whether plasma PTX3 concentrations increase in endurance-trained men. Ten young endurance-trained men and 12 age- and gender-matched sedentary controls participated in this study. Methods: We measured plasma PTX3 concentrations of the participants in each group. We also determined systemic arterial compliance (SAC) by using simultaneous M-mode ultrasound and arterial applanation tonometry of the common carotid artery and used high-density lipoprotein cholesterol (HDLC) as an index of cardioprotective effect. Results: Maximal oxygen uptake was significantly higher in the endurance-trained men than in the sedentary controls. SAC and HDLC were significantly higher in the endurance-trained men than in the sedentary controls (SAC: 1.74 ± 0.11 vs. 1.41 ± 0.09 ml/mmHg; p < 0.05, HDLC: 70 ± 5 vs. 57 ± 4 mg/dl; p < 0.05). Plasma PTX3 concentrations were markedly higher in the endurance-trained men than in the sedentary controls (0.93 ± 0.11 vs. 0.68 ± 0.06 ng/ml; p < 0.05). Relationships between plasma PTX3 concentrations and SAC and HDLC were linear. Conclusion: This is the first study revealing that endurance-trained individuals had higher levels of circulating PTX3 than sedentary controls. PTX3 may play a partial role in endurance exercise training-induced cardioprotection.

Keywords: endurance training, cardioprotection, systemic arterial compliance, high-density lipoprotein cholesterol
INTRODUCTION

Paragraph Number 1 It is generally accepted that an increase in regular physical activity, especially habitual endurance exercise, reduces cardiovascular risk factors (7, 15, 16). Endurance exercise training induces increase in both high-density lipoprotein cholesterol (HDLC) (2, 30) and arterial compliance (5, 6). Increased HDLC and arterial compliance have been recognized as having beneficial cardioprotective effects (20, 29).

Paragraph Number 2 Recently, pentraxin 3 (PTX3), which is mainly produced by endothelial cells, macrophages and smooth muscle cells in the atherosclerotic region (21, 24), has been identified as a substance playing an important role in cardioprotection and atheroprotection. It has been reported in mice that after transient ischemia in the left anterior descending coronary artery, the area of necrotic heart tissue expanded in PTX3-deficient mice compared to that in the control mice (25), suggesting that PTX3 can prevent ischemic tissue from necrotizing. Furthermore, a previous study demonstrated that PTX3 heals vascular injury via activation of tissue factor (17). Recently, it has been revealed that the mice lacking PTX3 promotes vascular inflammatory response and atherosclerosis (19). These findings suggest that PTX3 has cardioprotective and atheroprotective effects.

Paragraph Number 3 Since PTX3 is implicated in cardioprotection, it is reasonable to hypothesize that PTX3 participates in the mechanisms underlying endurance exercise training-induced cardioprotective effect. However, the relationship between plasma PTX3 concentrations and exercise training-induced cardioprotective effect remains unclear. We hypothesized that endurance trained
individuals have higher levels of plasma PTX3 than sedentary controls and this increase in PTX3 would partly participate in the mechanism underlying endurance exercise training-induced cardioprotection. To test our hypothesis, we measured plasma PTX3 concentrations; plasma HDL-C concentrations; and systemic arterial compliance (SAC) in endurance-trained men. We measured HDL-C and SAC as indices of endurance exercise training-induced cardioprotective effect.

**METHODS**

*Paragraph Number 4  Subjects.* All participants in this study were Japanese. Ten young endurance-trained men (19-26 years) and 12 age- and gender-matched sedentary controls (19-25 years) participated in this study. All of endurance-trained men’s careers were longer than 2 years. The training mainly consisted of some kind of running training, such as long-distance running and interval training, and which volume and intensity were $5.5 \pm 0.3$ sessions/wk ($2.4 \pm 0.3$ h/session) and the rating of 15–17 in the Borg’s scale (i.e., hard-very hard). On the other hand, control men had a sedentary lifestyle (no regular physical activity) for at least 2 years. All subjects were free of signs, symptoms, and history of any overt chronic diseases. None of the participants had a history of smoking, and none were currently taking any medications. Additionally, none of the subjects were NSAIDs or aspirin users. Before all measurements, the subjects refrained from alcohol consumption and intense physical activity (exercise) for 24 h and fasted overnight (12 h), without water. All measurements were performed after a resting period of at least 20 min at a constant room temperature (25°C).

*Paragraph Number 5  This study was reviewed and approved by the institutional review board at the University of Tsukuba. The study conformed to the
principles outlined in the Helsinki Declaration. All potential risks and procedures involved in the study were explained to the subjects, and written informed consent to participate in the study was obtained from all subjects.

**Paragraph Number 6  Maximal Oxygen Uptake.** The maximal oxygen uptake was determined during incremental cycling to exhaustion (3 min at 80 W, with a 30-W increase every 3 min) by monitoring breath-by-breath oxygen consumption and carbon dioxide production (AE280S; Minato Medical Science, Osaka, Japan), heart rate, and ratings of perceived exertion (Borg scale). The values of maximal oxygen uptake were accepted if subjects met at least 2 of the following criteria: a VO2 plateau (<150 ml O2/min with an increased work rate), highest respiratory exchange ratio >1.15, peak heart rate within 5 beats of the age-predicted maximum (220 minus the age in years), rating of perceived exertion >19, or extreme fatigue such that the pedaling rate on the bicycle ergometer was <50 rpm.

**Paragraph Number 7  SAC.** SAC was measured by carotid artery applanation tonometry and Doppler echocardiography as described previously (22). Briefly, carotid artery pressure waveforms were obtained by applanation tonometry (formPWV/ABI; Colin Medical Technology, Komaki, Japan) after a resting period of at least 20 min. At the time of waveform recording, brachial arterial systolic, diastolic, and mean blood pressure (SBP, DBP, and MBP, respectively) were measured by oscillometry (form PWV/ABI; Colin Medical Technology). The pressure signal obtained by tonometry was calibrated by equating the carotid MBP and DBP to brachial artery values. SAC was calculated as follows: SAC = \( \frac{Ad}{dP \times R} \), where \( Ad \) is the area under an arbitrary portion of the diastolic pressure waveform, \( dP \) is the pressure change in this portion, and \( R \) is systemic vascular resistance given as MBP divided by mean blood flow. The calculation of SAC is based on the assumption that the diastolic pressure decay is a mono-exponential function of time. Mean blood flow
was obtained using a Doppler echocardiographic system (EnVisor; Koninklijke Philips Electronics, Eindhoven, Netherlands) as described previously by our laboratory (22). The insertion point of the aortic valve tips at the end of diastole was defined by two-dimensional imaging in the parasternal long-axis view with a 3.5-MHz transducer, and the M-mode echocardiogram at that level was recorded with the computer. Doppler ultrasonographic flow velocity curves in the ascending aorta were simultaneously obtained using a 1.9-MHz probe held in the suprasternal notch. Mean blood flow was calculated as a product of the aortic cross-sectional area and the mean flow velocity (ImageJ; National Institutes of Health, Bethesda, MD).

**Paragraph Number 8  Plasma PTX3 Concentration.** All the blood samples were obtained from the antecubital vein with using a 21-gauge needle. Each blood sample was placed in a chilled tube containing ethylenediaminetetraacetic acid (EDTA) (2 mg/mL) and was then centrifuged at 2,000 g for 15 min at 4°C. The plasma was stored at –80°C until the assay. Plasma concentrations of PTX3 were determined using a commercial enzyme-linked immunosorbent assay (ELISA) kit (Quantakine DPTX 30; R&D Systems Inc., Minneapolis, USA). The PTX3 assay was carried out according to the manufacturer’s instructions. Briefly, standard or plasma samples assayed in duplicate, and 20 μl of which were added to microtiter plate wells coated with a monoclonal antibody specific for PTX3, followed by incubation at room temperature for 2 hours. The wells were then washed 4 times with a buffered surfactant solution, and thereafter, 200 μl of anti-PTX3 polyclonal antibody conjugated to alkaline phosphatase were added to each well and incubation for 2 hours at room temperature. After appropriate washing, 200 μl of substrate solution were added to each well and incubated again for 30 min at room temperature. The reaction was then stopped by the addition of 2N sulfuric acid to the wells, and absorbance was measured at 450 nm with corrections set at 540 nm using a
microplate reader. The values of plasma PTX3 levels were extrapolated from a curve drawn using standard PTX3. The intra- and inter-assay coefficients of variation were 3.8% and 6.1%, respectively (values provided by Quantakine DPTX 30; R&D Systems Inc.). The intra-assay coefficient of variation in this study was 5.6%. No significant cross-reactivity or interference with other factors related to PTX3 or other cytokines was observed (information provided by Quantakine DPTX 30; R&D Systems Inc.).

Paragraph Number 9 **Blood Biochemistry.** The serum concentrations of total cholesterol (TC), low-density lipoprotein cholesterol (LDLC), HDLC, and triglycerides (TG) and the plasma concentrations of glucose (BG) were determined using standard enzymatic techniques. Briefly, TG and TC concentrations were determined by the cholesterol dehydrogenase and glycerol kinase methods, respectively (1, 13). LDLC and HDLC concentrations were measured by a direct method (9, 31). The BG concentration was assayed by the hexokinase and glucose-6-phosphate dehydrogenase methods (26).

Paragraph Number 10 **Statistical Analysis.** Student’s t test for unpaired values was used to evaluate the statistical differences between the endurance-trained men and the sedentary controls. Relationships between SAC or HDLC and plasma PTX3 concentrations were analyzed using Pearson’s correlation. Data were expressed as means ± SE. Values of $P < 0.05$ were accepted as significant.

**RESULTS**

Paragraph Number 11 Table 1 summarizes the characteristics of the endurance-trained men and the sedentary controls. There were no significant differences in age, height, weight, BMI, TG, TC, LDLC, and BG between the
endurance-trained men and the sedentary controls. Table 2 shows the hemodynamics in the endurance-trained men and the sedentary controls. Diastolic blood pressure and resting heart rate were significantly lower in the endurance-trained men than in the sedentary controls. There was no significant difference in systolic blood pressure and pulse pressure between the two groups. Maximal oxygen uptake was higher in the endurance-trained men than in the sedentary controls (Table 1). HDLC in the endurance-trained men was markedly higher than the sedentary controls (Fig. 1). SAC was significantly higher in the endurance-trained men than in the sedentary controls (Fig. 2). Figure 3 shows plasma PTX3 concentrations in the two groups. Plasma PTX3 concentrations were higher in the endurance-trained men than in the sedentary controls. The relationships between plasma PTX3 concentrations and HDLC and SAC were linear (Fig. 4). However, no significant relationship was detected between maximal oxygen uptake and plasma PTX3 concentrations. In the trained group, we found a significant positive correlation between plasma PTX3 concentrations and HDLC (Fig. 5). However, there was no relation between plasma PTX3 concentrations and SAC in the trained group (Fig. 5). In the sedentary controls, plasma PTX3 concentrations were not related to HDLC or SAC (Fig. 5).

DISCUSSION

Paragraph Number 12 In the present study, we determined plasma PTX3 concentrations in endurance-trained men. It was first demonstrated that plasma PTX3 concentrations were markedly higher in the endurance-trained men than in the sedentary controls. The endurance-trained men also showed clearly higher maximal oxygen uptake, HDLC, and SAC than the sedentary controls. Furthermore, the
relationships between plasma PTX3 concentrations and HDLC and SAC were linear. An increase in PTX3 may play a role in the endurance exercise training-induced increase in HDLC and SAC, i.e., the cardioprotective effects induced by exercise training.

Paragraph Number 13 PTX3 is mainly produced by endothelial cells, macrophages, and smooth muscle cells in the local atherosclerotic region (21, 24). However, the role of PTX3 in the cardiovascular system is unclear. Circulating PTX3 concentrations were reported to increase in patients with cardiovascular disease (23, 27). Napoleone et al. (17) reported that PTX3 could repair vascular wounds by promoting activation of tissue factor. Peri et al. (23) demonstrated that PTX3 was produced from dying cardiomyocytes but not from necrotic cells in patients with acute myocardial infarction. Recently, it has been demonstrated that PTX3 functions at the crossway between pro-inflammatory and anti-inflammatory stimuli to balance the over activation of a pro-inflammatory, pro-atherogenic cascade (19). Namely, the increased levels of PTX3 in cardiovascular disease could reflect a protective physiological response (19). Salio et al. (25) demonstrated that after acute myocardial infarction, the exacerbated heart tissue area in PTX3-deficient mice had expanded compared to that in the control mice. Thus, PTX3 plays a role of repair in cardiovascular injury. Moreover, the recent report showed that the double-knockout mice lacking PTX3 and apolipoprotein E (ApoE) gene developed larger atherosclerosis than the mice lacking only ApoE (19). Taken together, it is thought that PTX3 has a cardioprotective and atheroprotective effects.

Paragraph Number 14 The benefit of habitual endurance exercise is recognized as a lifestyle modification worldwide. In epidemiological studies,
physically inactive subjects were reported have significantly higher risks of cardiovascular disease, and mortality rates in these subjects were reported to be high (7, 15, 16). Endurance exercise training produces beneficial cardioprotective effects. Increased HDLC and arterial compliance have been recognized as beneficial cardioprotective effects (20, 29). Habitual endurance exercise induces the increase in HDLC and SAC (2, 5, 6, 30). In the present study, HDLC, SAC, and plasma PTX3 concentrations were significantly higher in the endurance-trained men than in the sedentary controls. Furthermore, we demonstrated that there was a significant positive correlation between plasma PTX3 concentrations and SAC or HDLC. These findings suggest that endurance-trained men have beneficial cardioprotective effects and PTX3 may partly participate in the mechanism underlying endurance exercise training-induced cardioprotective effect.

Paragraph Number 15 It is known that high physical activity and/or endurance exercise training is effective for good health. On the other hand, exercise causes increase in inflammatory factors in various tissues, such as circulating blood, fat, and skeletal muscle (3, 10, 12, 18). A previous study reported that PTX3 is expressed and secreted in vascular walls as a result of the inflammatory response (11). Furthermore, it has been reported that PTX3 is produced via the myeloid differentiation protein 88-interleukin-1 receptor [MyD88-IL1R] pathway, which induces initial factors for starting inflammatory response (e.g., nuclear factor-kappa B [NF-kB]) (25). MyD88 is also known as a necessary factor for vascular remodeling (28). Tumor necrosis factor-α (TNF-α) and interleukin-1β (IL-1β) are known as triggers of PTX3 production (4, 14). On the other hand, anti-atherogenic IL-10 stimulates PTX3 production from dendric cells and monocytes (8). Additionally, PTX3 inhibits pro-atherogenic cytokines interferon-γ production (21). Norata et al. (19) recently
reported that PTX3 is a molecule for finely tuning vascular inflammatory response by both pro- and anti-inflammatory factors. Thus, PTX3 is modulated by both pro-atherogenic and anti-atherogenic factors. PTX3 may be a necessary substance for maintaining vascular homeostasis. Taken together, PTX3 participates in a part of inflammation and plays a role in cardioprotection and atheroprotection. However, the precise roles of PTX3 remain to be elucidated.

Paragraph Number 16 There are several limitations of this study that should be emphasized. First, this was a cross-sectional study. Therefore, the results suggesting a role for PTX3 in cardioprotection are preliminary. These findings need to be confirmed in a longitudinal study. Second, the small sample size is clearly one of the limitations of this study. We have demonstrated that plasma PTX3 concentrations, HDLC, SAC and maximal oxygen uptake were increased in endurance-trained men. Furthermore, the relationships between plasma PTX3 concentrations and HDLC and SAC were linear. However, there was no relation between plasma PTX3 concentrations and maximal oxygen uptake. This may be the influence of a small sample size in the present study. Furthermore, the subjects in this study were young Japanese men. Therefore, these results may not generalize to other populations.

Paragraph Number 17 In conclusion, the present study revealed for the first time that circulating PTX3 concentrations are markedly higher in endurance-trained men than in sedentary controls. We also demonstrated that SAC and HDLC, which are cardioprotective factors, were elevated by the regular endurance exercise. It is possible that PTX3 may partly participate in the mechanism underlying endurance exercise training-induced cardioprotection.
ACKNOWLEDGMENTS

This work was supported by Grants-in-Aid for Scientific Research 21300234 and 21650179 and Grant-in-Aid for JSPS Fellows 21·692 from Japan Society for the Promotion of Science. And, the results of the present study do not constitute endorsement by ACSM.

CONFLICTS OF INTEREST

The authors have no financial, consultant, institutional, or other relationships that might lead to bias or a conflict of interest.
REFERENCES


FIGURE LEGENDS

Figure 1. High-density lipoprotein cholesterol (HDLC) in endurance-trained men and in sedentary controls. Data are expressed as means ± SE.

Figure 2. Systemic arterial compliance (SAC) in endurance-trained men and in sedentary controls. Data are expressed as means ± SE.

Figure 3. Plasma pentraxin 3 (PTX3) concentrations in endurance-trained men and in sedentary controls. Data are expressed as means ± SE.

Figure 4. Relationships between plasma PTX3 concentrations and HDLC (A) and SAC (B) were linear. Endurance-trained men (△) and sedentary controls (○) are shown.

Figure 5. Relationships between plasma PTX3 concentrations and HDLC (A) and SAC (B) in endurance-trained men and between plasma PTX3 concentrations and HDLC (C) and SAC (D) in sedentary controls.
Table. 1 Characteristics of sedentary control men and endurance-trained men.

<table>
<thead>
<tr>
<th></th>
<th>Sedentary</th>
<th>Endurance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, years</td>
<td>20.8 ± 0.8</td>
<td>20.7 ± 0.6</td>
</tr>
<tr>
<td>Height, cm</td>
<td>173.2 ± 1.5</td>
<td>173.3 ± 1.9</td>
</tr>
<tr>
<td>Weight, kg</td>
<td>66.3 ± 2.1</td>
<td>62.5 ± 1.9</td>
</tr>
<tr>
<td>BMI, kg/m²</td>
<td>22.1 ± 0.6</td>
<td>20.8 ± 0.4</td>
</tr>
<tr>
<td>TG, mg/dL</td>
<td>75 ± 16</td>
<td>72 ± 14</td>
</tr>
<tr>
<td>TC, mg/dL</td>
<td>178 ± 9</td>
<td>183 ± 9</td>
</tr>
<tr>
<td>LDLC, mg/dL</td>
<td>107 ± 10</td>
<td>101 ± 8</td>
</tr>
<tr>
<td>BG, mg/dL</td>
<td>88 ± 3</td>
<td>86 ± 2</td>
</tr>
<tr>
<td>Maximal oxygen uptake, ml/min/kg</td>
<td>44.7 ± 1.0</td>
<td>60.3 ± 0.8</td>
</tr>
</tbody>
</table>

Data are expressed as means ± SE. Sedntary, sedentary control men; Endurance, endurance-trained men.
Table. 2 Hemodynamics of sedentary control men and endurance-trained men

<table>
<thead>
<tr>
<th></th>
<th>Sedentary</th>
<th>Endurance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Systolic blood pressure, mmHg</td>
<td>118 ± 3</td>
<td>114 ± 3</td>
</tr>
<tr>
<td>Diastolic blood pressure, mmHg</td>
<td>65 ± 2</td>
<td>59 ± 1**</td>
</tr>
<tr>
<td>Pulse pressure, mmHg</td>
<td>53 ± 1</td>
<td>55 ± 2</td>
</tr>
<tr>
<td>Heart rate, bpm</td>
<td>63 ± 3</td>
<td>53 ± 3*</td>
</tr>
</tbody>
</table>

Data are expressed as means ± SE.  *P < 0.05 vs. Sedentary, **P < 0.01 vs. Sedentary
Figure 1 (Miyaki et al.)

HDL-cholesterol (mg/dL)

<table>
<thead>
<tr>
<th></th>
<th>Sedentary</th>
<th>Endurance</th>
</tr>
</thead>
<tbody>
<tr>
<td>P</td>
<td></td>
<td>&lt; 0.05</td>
</tr>
<tr>
<td>Value</td>
<td>57 ± 4</td>
<td>70 ± 5</td>
</tr>
</tbody>
</table>

Miyaki et al. Page 21
Figure 2 (Miyaki et al.)

- **Systemic arterial compliance**
- **(ml/mmHg)**
- **P < 0.05**

- **Sedentary**
  - 1.41 ± 0.09

- **Endurance**
  - 1.74 ± 0.11
Figure 3 (Miyaki et al.)

Plasma PTX3 concentration (ng/ml)

- **Sedentary**: 0.68 ± 0.06
- **Endurance**: 0.93 ± 0.11

P < 0.05
Figure 4 (Miyaki et al.)

A

Endurance-trained men
Control

\[ y = 26.63x + 41.61 \]
\[ r = 0.54 \text{ (p<0.01)} \]

HDL cholesterol

PTX3 (ng/ml)

B

\[ y = 0.52x + 1.15 \]
\[ r = 0.44 \text{ (p<0.05)} \]

Systemic arterial compliance

PTX3 (ng/ml)
Figure 5 (Miyaki et al.)

A

\[ y = 27.86x + 43.88 \]
\[ r = 0.65 \quad (p < 0.05) \]

B

\[ y = 0.464x + 1.31 \]
\[ r = 0.46 \quad (p = 0.18) \]

C

\[ y = 6.40x + 52.48 \]
\[ r = 0.11 \quad (p = 0.74) \]

D

\[ y = 0.089x + 1.35 \]
\[ r = 0.06 \quad (p = 0.85) \]