Effect of acupuncture on salivary immunoglobulin A after a bout of intense exercise

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ABSTRACT

In the field of athletics, acupuncture has been used for treatment of injury, reduction of fatigue and management of physical condition. However, there is little information on the effect of acupuncture on the immune function in response to exercise.

Purpose

The aim of this study was to examine the effect of acupuncture treatment on the mucosal immune function after a single period of intense exercise by measuring salivary immunoglobulin A (SIgA).

Methods

12 healthy men (23.6 ± 5.3 years) participated in this study with a crossover design. The subjects exercised on a bicycle equipped with an ergometer at 75% VO2max for 60 min. Acupuncture treatment was applied at LU6, LI4, ST36 and ST6, for 30 min after the exercise. The control treatment was rest without acupuncture and that the order of the treatment was randomised. We measured parameters including saliva flow rate, SIgA concentration, SIgA secretion rate, heart rate and plasma catecholamine concentration all before the exercise and at 1 h, 2 h, 3 h, 4 h and 24 h after the exercise. The visual analogue scale for self-perceived tiredness and mood state questionnaires were recorded before the exercise and at 24 h after the exercise.

Result

Intense exercise-induced decrease of SIgA levels was attenuated by the acupuncture treatment. In contrast, the subjective fatigue score and psychological measurement were not affected by the acupuncture.

Conclusion

Acupuncture treatment may attenuate the decrease in SIgA level induced by intense exercise.

INTRODUCTION

Athletes have to maintain and even improve high-fitness levels and skills to perform well at competitions. Therefore, they need to do high-frequency, high-intensity training. Among elite athletes and coaches it is well known that athletes have a higher risk of upper respiratory tract infection (URTl) than non-athletes.1 It has been reported that acute high-intensity exercise and long-term heavy training lead to a decrease in immune functions.2-3 Additionally, Nieman has reported that people who habitually do moderate exercise have less risk of URTl than those people who do not; however, he also has reported that high-intensity exercise increases the risk of URTl.4 Since it is known that URTl adversely affects training and performance,5 it is important for athletes to avoid URTl in order to maintain a good physical and mental condition.6 7

Several studies have shown that suppression of salivary immunoglobulin A (SIgA) level in elite athletes frequently results in URTl.8 9 SIgA, which is the predominant immunoglobulin in secretions of the mucosal immune system, penetrates the body through the mucosal tissues,10 11 neutralises various toxins and viruses, and inhibits the attachment and replication of pathogenic microorganisms.12 13

Acupuncture is used most commonly as an adjunctive treatment for the relief of chronic or acute pain. Acupuncture treatment is also a very common method for modulating the physical condition of various athletes.14 In athletics, acupuncture is most widely used to treat pain associated with musculoskeletal injuries, but it has also been used to treat physical conditions, such as relieving pain and tiredness, and promote regeneration of injured areas of the body. Furthermore, it has been reported that acupuncture treatment is effective for soothing tight muscles, improving peripheral blood circulation, increasing the pain threshold and adjusting the autonomic nervous system.15-17 Acupuncture also has been recognised to increase ‘natural healing power,’ including immunological responses that can improve the physical and mental condition and even can cure and prevent diseases.18

Although acupuncture treatment is considered to adjust the immune function and contribute to the stabilisation of human homeostasis,19 few studies have been conducted to determine the combined effects of acupuncture treatment and exercise on the human immune function. So far, our group has published only one study investigating the effect of acupuncture treatment on the immune function of athletes during competition.20 21 We reported that continuous acupuncture treatment counteracted intense training-induced decrease in SIgA and also had positive effects on physical and mental conditions. This study was the first to demonstrate that acupuncture treatment attenuated the reduction in the mucosal immune function of athletes during competition. Since this study was conducted during a period of sports competition, it could serve as a basic experiment for a new study, one that investigates the capability of acupuncture treatment to attenuate both the acute intense exercise-induced decrease in SIgA and the reduction in the mucosal immune function in athletes. The only conceptual change necessary in the experiment would be to omit the factors other than acupuncture treatment that affect SIgA levels.

The purpose of this study was to examine the effect of acupuncture treatment on the mucosal immune function after acute high-intensity exercise by measuring salivary SIgA levels. We hypothesised that acupuncture treatment would lessen the decrease of SIgA induced by the exercise.
VO2max. Subjects then underwent a submaximal cycle exercise (1) increased workload without corresponding increase in VO2; criteria for maximal effort included at least two of the following: increasing to 60, 80 and 100 W for 2 min. Subsequently, the workload was monitored on a cycle ergometer (AE280S; Minato Medical Science, Osaka, Japan). The protocol consisted of 2 min of unloaded pedalling and subsequent incremental exercise. The workload was increased at 60,80 and 100 W for 2 min. Subsequently, the workload increased by 30 W every 3 min until exhaustion. Objective criteria for maximal effort included at least two of the following: (1) increased workload without corresponding increase in VO2; (2) respiratory exchange quotient equal to or greater than 1.10; (3) pedal cadence less than 50 rpm in spite of maximal voluntary effort. The highest O2 uptake over a 30 s period was defined as VO2max. Subjects then underwent a submaximal cycle exercise test. The tests were taken within 1 week of the VO2max test, at least 4 d apart. In the submaximal cycle exercise test, subjects exercised on a cycle ergometer for 1 h at 75% of their VO2max.

**Exercise test**

Maximal oxygen uptake (VO2max) of the subjects was determined through the use of an incremental bicycle exercise test. The subjects rode the bicycle to exhaustion, and their breath-by-breath oxygen uptake and carbon dioxide production were monitored on a cycle ergometer (AE280S; Minato Medical Science, Osaka, Japan). The protocol consisted of 2 min of unloaded pedalling and subsequent incremental exercise. The workload was increased at 60,80 and 100 W for 2 min. Subsequently, the workload increased by 30 W every 3 min until exhaustion. Objective criteria for maximal effort included at least two of the following: (1) increased workload without corresponding increase in VO2; (2) respiratory exchange quotient equal to or greater than 1.10; (3) pedal cadence less than 50 rpm in spite of maximal voluntary effort. The highest O2 uptake over a 30 s period was defined as VO2max. Subjects then underwent a submaximal cycle exercise test. The tests were taken within 1 week of the VO2max test, at least 4 d apart. In the submaximal cycle exercise test, subjects exercised on a cycle ergometer for 1 h at 75% of their VO2max.

**Heart rates**

The heart rates of the subjects were measured at the same points of sample collections. The heart rate was measured by Polar S610 (Polar, Kempele, Finland).

**Subjective evaluations of physical and mental conditions**

Subjective evaluations of subject conditions were made during the experiment. The visual analogue scale for self-perceived tiredness was used at the same points as sample collections. This rating was assessed using a 100 mm continuous scale. In addition, a Japanese version of the profile of mood state (POMS) questionnaire was administered to assess the subjective mental states before the exercise test and after 24 h. Six mood states are used in POMS: T-A (tension), D (depression), A-H (anger), V (vigour), F (fatigue) and C (confusion). Subjects are given a score for each trait according to their responses to 65 statements. Each mood dimension was rated on a scale of 1 to 4 from ‘not at all’ to ‘extremely.’

**Acupuncture treatment**

After the submaximal cycle exercise test, the acupuncture treatment was performed by an experienced acupuncturist, and the same points were used as in our previous study. The following bilateral points were used as acupuncture points: LU6 in the forearms, LI4 in the hands, ST36 in the legs and ST6 in both sides of the face. After standard disinfecting of the insertion sites, disposable stainless needles (50 mm long, 0.20 mm in diameter, SEIRIN, Shizuoka, Japan) were inserted through the skin to a depth of 5–10 mm, and manipulated until the subjects felt a sensation from the needles. Electrodes were connected to three points (LU6, LI4 and ST36) and then attached to an electric stimulator (OhmPulsar LFP-4800; Zen Iryoki, Fukuoka, Japan). The points were electrically stimulated with a low frequency of 2 Hz for 30 min. At the same time, the fourth point (ST6) was manually stimulated for 30 min until needle sensation was reached every 5 min.

**Saliva collection**

Saliva samples were collected at rest before exercise (pre), immediately after exercise (P0), 1, 2, 3 and 4 h after exercise (P1, P2, P3, P4), and 24 h after exercise (P24), as described in our previous studies. The subjects were free to drink water during the experiment. Light meals were given to all the subjects at the sample collection, which was taken within 1 h after the exercise. The subjects rinsed their mouths for 30 sec 3 times with 100 ml of mineral water, and then rested for at least 5 min. Saliva production was stimulated by the chewing of sterilised cotton (Salivet; Saersted, Vumbrecht, Germany) for 2 min at a frequency of 1 chew/sec. Saliva was separated from the cotton swab by centrifugation at 3000 rpm for 15 min. The amount of saliva in grammes was converted to millilitres based on an assumed saliva density of 1 g/ml, as described in one of our previous studies. After the sample volume was measured, samples were frozen at −80°C and stored until analysis.

**SIgA measurement**

SIgA concentrations were measured by enzyme-linked immunosorbent assay, according to the procedures of our previous study. Rabbit antihuman secretory component IgG fraction (MBL, Nagoya, Japan) with coating buffer was added to each well of a microtitre plate (IMMULON-2, Dynex Technologies, Chantilly, VA) and kept at 4°C for more than 8 h. After 250 μl of phosphate-buffered saline (PBS) containing 1% bovine serum albumin (BSA, Sigma, St Louis, MO) was added, the wells were blocked for 2 h at normal room temperature. Saliva samples were thawed, centrifuged at 10 000 rpm for 5 min, diluted with 100 μl of PBS containing 1% BSA and incubated for 1 h. By using purified human SIgA (MP Biomedical, Inc, Santa Ana, CA), known concentrations of SIgA were also plated to establish standard values. After the plate was washed with PBS-Tween, goat Fab/anti-IgA conjugated with horse radish peroxide HRP (mose binding lectin) was added to the plate and incubated for 1 h. After washing, substrate solution was added, and the colour intensity produced after 10 min was measured by a microplate reader (Model550, Bio Rad, Hercules, CA) at 490 nm wavelength. For analysis of the SIgA levels, data were expressed as the SIgA secretion rate (μg/min). This rate was calculated by

![Table 1 Subjects’ characteristics](https://example.com/table1.jpg)

<table>
<thead>
<tr>
<th>Number</th>
<th>12</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (year)</td>
<td>23.6±0.3</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>171.4±1.6</td>
</tr>
<tr>
<td>Body mass (kg)</td>
<td>66.0±2.4</td>
</tr>
<tr>
<td>Body fat (%)</td>
<td>16.2±1.1</td>
</tr>
<tr>
<td>Maximal oxygen uptake (ml/kg/min)</td>
<td>41.6±1.6</td>
</tr>
</tbody>
</table>

**Date** were expressed mean±SE
multiplying the absolute SIgA concentration (μg/ml) by the saliva flow rate (ml/min), which was calculated by dividing the total volume of saliva obtained in each sample (ml) by the time taken to produce the sample (min).

**Blood collection and measurement of plasma catecholamine concentrations**

Blood samples were collected at the same times as saliva collections. Blood was sampled from the antecubital vein in the arm. Plasma was generated by centrifuging blood at 3000 rpm for 15 min at 4°C; it was frozen at −80°C until the analysis of catecholamine concentrations. Plasma concentrations of epinephrine and norepinephrine were measured by an automatic catecholamine analyser (HLC-725CA II, Tosoh, Tokyo, Japan).

**Statistical analysis**

All data were represented as mean±SE. The effect of acupuncture treatment was determined by using two-way analyses of variance with repeated measures. A Tukey-Kramer adjustment was applied to the posthoc test. The significance level of all statistical analyses was set at p<0.05.

**RESULTS**

**Salivary SIgA levels**

As shown in figure 1, the SIgA secretion rate significantly decreased from P0 to P3 in the control group (p<0.05) but did not decrease significantly in the acupuncture group. In contrast, the SIgA of the acupuncture group significantly increased at P24. The SIgA secretion rate was significantly higher in the acupuncture group than in the control group at P1 and P24 (p<0.05). The concentration of SIgA significantly decreased from P0 to P3 in the control group (p<0.05) and decreased significantly only at P2 in the acupuncture group (figure 2).

**Saliva flow rates**

Figure 3 shows that the saliva flow rate did not significantly change in the control group. In contrast, the saliva flow rate of the acupuncture group significantly increased at P1 and P24 (p<0.05). The saliva flow rate was significantly higher in the acupuncture group than in the control group at P1 and P24 (p<0.05).

**Plasma catecholamine concentrations**

As shown in table 2, the epinephrine concentration increased at P0 and P1 in the control group. In contrast, the epinephrine concentration of the acupuncture group increased at P0. The norepi-
Acupuncture 0.03±0.01

Epinephrine (ng/ml)

<table>
<thead>
<tr>
<th></th>
<th>Pre</th>
<th>Post</th>
<th>Post 1 h</th>
<th>Post 2 h</th>
<th>Post 3 h</th>
<th>Post 4 h</th>
<th>Post 24 h</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acupuncture</td>
<td>0.03±0.01</td>
<td>0.09±0.04*</td>
<td>0.04±0.02</td>
<td>0.03±0.01</td>
<td>0.03±0.01</td>
<td>0.04±0.02</td>
<td>0.04±0.02</td>
</tr>
<tr>
<td>Control</td>
<td>0.03±0.01</td>
<td>0.11±0.04*</td>
<td>0.05±0.03*</td>
<td>0.03±0.01</td>
<td>0.03±0.01</td>
<td>0.04±0.02</td>
<td>0.03±0.02</td>
</tr>
<tr>
<td>Norepinephrine (ng/ml)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Acupuncture</td>
<td>0.22±0.08</td>
<td>0.69±0.12*</td>
<td>0.30±0.09</td>
<td>0.32±0.10*</td>
<td>0.32±0.09</td>
<td>0.30±0.07</td>
<td>0.29±0.07</td>
</tr>
<tr>
<td>Control</td>
<td>0.25±0.04</td>
<td>0.69±0.24*</td>
<td>0.31±0.08</td>
<td>0.37±0.14*</td>
<td>0.33±0.08*</td>
<td>0.30±0.05</td>
<td>0.29±0.06</td>
</tr>
</tbody>
</table>

Date were expressed as mean±SE.
*p<0.05 (vs Pre)

Table 3 Temporal change in heart rate after the exercise

<table>
<thead>
<tr>
<th></th>
<th>Pre</th>
<th>Post</th>
<th>Post 1 h</th>
<th>Post 2 h</th>
<th>Post 3 h</th>
<th>Post 4 h</th>
<th>Post 24 h</th>
</tr>
</thead>
<tbody>
<tr>
<td>Heart rate (beats/min)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Acupuncture</td>
<td>57.0±2.1</td>
<td>170.0±3.5*</td>
<td>72.2±2.7*</td>
<td>74.8±2.2*</td>
<td>70.6±2.5*</td>
<td>65.8±2.8*</td>
<td>59.3±2.0</td>
</tr>
<tr>
<td>Control</td>
<td>61.0±3.3</td>
<td>169.8±2.7*</td>
<td>75.8±3.0*</td>
<td>78.5±2.8*</td>
<td>74.7±2.4*</td>
<td>68.3±3.0</td>
<td>60.3±1.9</td>
</tr>
</tbody>
</table>

Date were expressed as mean±SE.
*p<0.05 (vs pre)

neuropehrine concentration increased at P0 and P2 in the control group. In contrast, the norepinephrine concentration of the acupuncture group increased at P0, P2 and P3.

Heart rates

Table 3 shows that the heart rate increased from P0 to P3 in the control group. In contrast, the heart rate of the acupuncture group increased from P0 to P4.

Subjective evaluations of physical and mental conditions

The visual analogue scale score on fatigue increased for both groups at P0 but returned to baseline after P0. All the mood states except ‘vigour’ at P24 in both groups were lower than they were at pre. However, there were no significant differences between the groups on any items.

DISCUSSION

In the present study, we provided evidence that acupuncture treatment positively affects SIgA levels and saliva flow rates in healthy sedentary male subjects after a single bout of intense exercise. This result suggests that acupuncture treatment works on the autonomic nervous system and may improve mucosal immunity protection.

SIgA has the important function of blocking entrance of pathogenic organisms and preventing infections. It has been reported that reduction in SIgA levels might increase the risk of URTI. Moreover, it has been reported that the autonomic nervous system reaction relate to central nerve system which stimulated by acupuncture. This means that stimulating the sympathetic, the parasympathetic nerve, or both nerves by acupuncture treatment could promote SlgA secretion. In this study, besides saliva, epinephrine, norepinephrine and heart rate, which are related to the autonomic nervous system, were measured. In the acupuncture group, the saliva flow rate increased at P1 and P24 but did not change in the control group throughout the study. On the other hand, the acupuncture treatment did not affect epinephrine, norepinephrine and heart rate whereas the intensive exercise did affect those factors. These results might suggest that the acupuncture treatment did not affect the whole autonomic nervous system, but only affected a part of it after intensive exercise. Therefore, it is possible that acupuncture treatment might improve SIgA secretion by stimulating the autonomic nervous system.

In this study, acute high-intensity exercise decreased SIgA levels, but then acupuncture treatment increased SIgA levels. However, the acupuncture treatment had no effect on the self-perceived tiredness resulting from the acute high-intensity exercise. There also were no effects on the POMS score in the acupuncture group. On the other hand, the acupuncture treatment did not affect epinephrine, norepinephrine and heart rate whereas the intensive exercise did affect those factors. Therefore, it is possible that acupuncture treatment might improve SlgA secretion by stimulating the autonomic nervous system.

It has been shown that acupuncture treatment restores the balance of the body’s function. In addition, in actual acupuncture treatment, specific acupoints are selected based on each patient’s individual conditions. In our study, the same acupuncture points were selected and the same treatments were given to
all subjects, in order to keep all experimental conditions equal. Furthermore, the acupuncture treatment given in our study was physical stimulation by a needle, which might have a little different effect from that of a typical acupuncture treatment in clinics.

There is a need in the future to validate how effective acupuncture treatment is for decreasing URTI risks, by examining the relation between the influence of acupuncture treatment on SlgA levels and the incidence of URTI during a period of sports competition. There also is a need to define the mechanism by which acupuncture treatment promotes SlgA secretion. In addition, further research is needed to determine whether the effect of acupuncture was point specific in this study and whether electrical stimulation is necessary in the future. Once all the tasks mentioned above have been completed successfully, it is conceivable that acupuncture could be widely recognised as an effective treatment to keep athletes from getting URTI by adjusting their immune systems and even help athletes with their self-conditioning.

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Competing interests None.

Provenance and peer review Not commissioned; externally peer reviewed.

Ethics approval This study was conducted with the approval of the Ethical Committees of the Institute of Health and Sport Sciences and the Institute of Clinical Medicine.

Patient consent Obtained.

REFERENCES