Panax ginseng and salvia miltiorrhiza supplementation abolishes eccentric exercise-induced vascular stiffening: a double-blind randomized control trial

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Abstract

Background: Muscle damage induced by unaccustomed or eccentric exercise results in delayed onset vascular stiffening. We tested the hypothesis that a 7-day supplementation of panax ginseng and salvia miltiorrhiza prior to an acute eccentric exercise could attenuate arterial stiffening.

Methods: By using a double-blind study placebo-controlled randomized design, subjects were randomly assigned to either the Chinese herb (N = 12) or the placebo group (N = 11) and performed a downhill running (eccentric exercise) trial and a control (seated rest) trial.

Results: Muscle soreness increased 1–2 days after exercise similarly in both groups, whereas the herb group demonstrated a faster recovery on active range of motion. Plasma creatine kinase concentration increased significantly at 24 h in both groups but the magnitude of increase was attenuated in the herb group. Arterial stiffness as measured by carotid-femoral pulse wave velocity increased significantly at 24 h in the placebo group but such increase was absent in the herb group. Flow-mediated dilation did not change in either group. Plasma concentrations of CRP and IL-6 increased in the placebo group but no such increases were observed in the herb group. Changes in arterial stiffness induced by eccentric exercise were associated with the corresponding changes in IL-6 (r = 0.46, P < 0.05).

Conclusions: A short-term Chinese herb supplementation of panax ginseng and salvia miltiorrhiza ameliorated the delayed onset vascular stiffening induced by acute downhill running exercise.

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Keywords: muscle damage, inflammation, arterial stiffness

Background

Muscle damage induced by unaccustomed or eccentric exercise is associated with increases in oxidative stress, inflammatory response, and delayed onset muscle soreness [1]. The increases in circulating pro-inflammatory cytokine and C-reactive protein (CRP) are the characteristic responses induced by eccentric exercise [2, 3]. A growing body of evidence indicates that muscle damage may also exert adverse influences on vascular function a day or two days later on a similar time frame to delayed onset muscle soreness. An impairment in microcirculation [4, 5], an increase in vascular resistance [6], and a reduction in vascular reactivity [7, 8] have been observed after acute eccentric exercise. We [9] and others [10] have demonstrated that acute eccentric exercise induced significant increases in central arterial stiffness and arterial stiffening after eccentric exercise was associated with indicators of muscle damage [9].

In Chinese Medicine, Ginseng is one of the most commonly used herbs in over thousands of years [11]. Ginsenosides, the major compounds of ginseng, and its metabolites are considered to exert protective effects on the vasculature, acting as a free radical scavenger [12] and increasing nitric oxide production and antioxidant
Effects [13]. There have been a number of animal studies demonstrating that supplementation with either Asian ginseng (panax ginseng C. A. Meyer) or American ginseng (panax quinquefolium L) could protect against eccentric or strenuous exercise-induced muscle damage by attenuating CK release [14, 15] and inflammatory responses [16]. Danshen (salvia miltiorrhiza) is another widely used Chinese medicinal herb with diverse pharmacological properties to improve circulation and blood stasis [17], including dilating coronary arteries, increasing blood flow, and scavenging free radicals in ischemic diseases [18]. Indeed danshen has been prescribed to treat angina pectoris, hyperlipidemia, acute ischemic stroke [19], and coronary heart disease [20]. The major compounds of danshen, Tanshinone IIA and salvianolic acid B, have been shown to suppress vasoconstrictor endothelin-1 production [21] and reduce the expression of vascular adhesion molecules in vitro [22, 23]. Panax ginseng and danshen are often mixed in herb formulas in Chinese medicine, which is characterized by adapting several types of herbs or minerals as a combination of multiple components that could synergistically attack different pathological targets [24]. Panax ginseng and danshen has been practiced as a formula to treat cardiovascular disease in Chinese Medicine [24, 25]; however, scientific evidence to support its use is still lacking. In this study we tested the hypothesis that the supplementation with a combination of panax ginseng and danshen could exert protective effects on the vasculature following eccentric exercise.

Methods
Participants
A total of 24 apparently healthy young male adults surrounding by National Taiwan University community were recruited. Exclusions from the study participation were due to: (1) obesity (BMI >30 kg/m²); (2) smoking within past six months; (3) hypertension (high blood pressure >140/90 mmHg); (4) personal history of diabetes (fasting blood glucose >126 mg/dL), history of heart disease or other cardiovascular problems; (5) orthopedic injury that may prevent him or her from completing the exercise; or (6) the use of over-the-counter supplements or vitamins. Subjects must have been sedentary or recreational active, but not been participating in any type of resistance or endurance training. All subjects gave their inform written consent prior to study participation and procedures were reviewed and approved by Institutional Review Board of National Taiwan University Hospital. This study is listed in ClinicalTriall.gov (NCT02007304).

Experimental design
Subjects were randomly assigned into either the Chinese herb supplement or the placebo group after pre-screening and familiarization. In each group, subjects underwent two familiarization sessions followed by a pre-testing session that consists of the measurements of aerobic power and body composition. Eccentric exercise trials took place following a 7-day supplementation. Subjects were asked to keep their regular diet and sedentary lifestyle throughout the testing sessions.

Aerobic power
Individual peak aerobic power (VO₂peak) was determined using standard American College of Sports Medicine protocol. After a 5-min warm-up on the treadmill, subjects walked or ran while treadmill slope was increased 1 % every minute until the subjects could not continue the test. A mouthpiece and heart rate monitor were worn to collect expired air and assess heart rate throughout the test. VO₂peak was used to set the exercise intensity during the eccentric exercise.

Body composition
Percent body fat was measured noninvasively by using a bioimpedance analyzer Inbody 2.0 (Biospace Co. Ltd., Seoul, South Korea). To avoid the hydration effects, the test was performed in the morning when subjects were fasted.

Supplement administration
Following the pre-testing sessions, subjects were asked to take a total of 7 capsules of either Chinese herb or placebo per day for seven days. Herb supplement was prepared in capsules consisting of 250 mg of panax ginseng and 250 mg salvia miltiorrhiza via the water-extraction method, whereas placebo capsules contained microcrystalline cellulose. According to the pharmacopoeia of the People's Republic of China [26], the use of 1–3 g danshen extract per day is recommended [27]. A dosage of 1.75 g per day (~60 % of the maximal suggested dosage) was chosen for danshen in order to avoid unexpected adverse event and to take consideration that this was a combination herb therapy. To the best of our knowledge, a combination of panax ginseng and salvia miltiorrhiza together has not been investigated on humans in the literature [24]. Accordingly, we decided to adopt the 1:1 ratio (i.e., the same dosage) of panax ginseng and salvia miltiorrhiza as the supplementation. Similar to danshen, this particular dosage of panax ginseng has been shown to be effective. Hence, the total daily supplementation dosage was 3.5 g together.

Both Chinese herb and placebo capsules were identical in appearance and stored in identical bottles with labeled numbers generated by a study-independent researcher. All supplement products were prepared by the Brion Research Institute, Sun Ten Pharmaceutical Co. Analyses of ginsenosides of panax ginseng as well as Salvianolic acid B and
Tanshinone IIA in Radix *salvia miltiorrhiza* were performed by using high-performance liquid chromatography-electrospray mass (HPLC-MS) spectrometry method as previously described [26, 28]. These chromatographic quantification results of active compounds in herb supplement are shown in Table 1 and Fig. 1.

**Exercise protocol**

Subjects were instructed to fast at least 8 h and refrain from any strenuous exercise for at least 72 h before the test. Experimental trial consisted of baseline measurements, downhill running (eccentric exercise) or seated rest (control), and measurements during the recovery period. In order to eliminate diurnal variation of inflammatory response to eccentric exercise, participants were asked to perform both eccentric exercise and control trials at the same time of day. Subjects warmed up on treadmill on a level grade at the speed that could elicit 75 % of predetermined individual VO2 peak [29] for 5 min. Each subject performed downhill running exercise on treadmill with the same speed at −10° of slope for 30 min. Similar protocols have been successfully used elsewhere to induce delayed onset muscle soreness [10, 30].

**Measurements**

The measurements were made 5 times: 30 min pre, 90 min post, 24 h post, 48 h post, and 72 h post. Subjects were studied at the same time of day, during the morning hours to minimize the inconvenience of the 8-h fast and to avoid diurnal effects.

*Blood samples* were collected to determine metabolic risk factors, markers of muscle damage, inflammation, and redox state. Serum CK was used as an indicator of muscle membrane permeability or muscle damage [31]. Inflammatory markers (TNF-α, IL-6) as well as blood redox status marker, tiobarbituric acid-reactive substances (TBARS), were analyzed with the use of commercial ELISA kits. Due to the financial constraints, TNF-α, IL-6, and TBARS were measured only in the eccentric exercise condition. The inter- and intra-assay coefficients of variation were less than 10 % in all assays performed.

*Heart rate and blood pressure* were measured in the supine position. Heart rate was measured using an ECG, and blood pressure was measured using an automatic blood pressure monitor (Omron HEM907).

**Muscle soreness**

Subjects were asked to rate the perception of muscle soreness using a Visual Analog Scale of 0−10 with 0 describing no soreness and 10 describing unbearable soreness immediately after a downhill running [32]. In addition, active range of motion was measured while the subjects were placed on bed in prone position with full knee extension and then moved both legs gradually to the flexion point where pain in quadriceps muscle groups was experienced. A manual goniometer was used to measure the knee angle difference from full extension to flexion point with the initiation of pain. This test was repeated three times, and the average was used for statistical analysis.

**Arterial stiffness**

Arterial stiffness was measured using carotid-femoral pulse wave velocity (cfPWV), which was calculated from the traveling distance and foot-to-foot wave transit time between the two arterial recording sites in the supine position [33], and was the primary outcome measure of this study. Non-invasive pulse tonometer (SPT-301, Millar Inc. Houston TX) connected to a physiological signaling processing system (MP36, Biopac, Goleta CA) was used to detect pulse waves on the carotid and femoral arteries. The coefficient of variation for cfPWV in our laboratory were 6.2 %.

**Vascular reactivity**

Flow-mediated dilatation, the secondary outcome measure, was obtained noninvasively at the brachial artery using standardized procedure [34]. Brachial artery diameter was measured using an ultrasound machine (Sonosite Ultrasound System; Bothell, WA) equipped with a high-resolution linear array transducer. A blood pressure cuff was placed on the forearm 3−5 cm distal to the antecubital fossa, and longitudinal images of the brachial artery were acquired 5−10 cm proximal to the antecubital fossa. After the acquisition of baseline measurement, the probe position was clearly marked to ensure that the image was acquired from the same location throughout the test. The blood pressure cuff was inflated to 100 mmHg above resting systolic blood pressure for 5 min by using a customized rapid inflation system. After cuff deflation, ultrasound-derived measurements of artery diameters were taken for 3 min. FMD was calculated by the following equation: (maximum diameter − baseline diameter)/baseline diameter × 100. All ultrasound images were recorded and analyzed by the same investigator who was blinded to the groups and the conditions. Our coefficient

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**Table 1** Quantitative analyses of major compounds of the herb supplement used in the present study

| Compound          | Content (mg/g) |
|-------------------|----------------|
| *Panax ginseng*   |                |
| Rb1               | 2.24           |
| Re                | 1.13           |
| Rg1               | 1.47           |
| *Salvia miltiorrhiza* |            |
| Salvianola acid B | 28.2           |
| Tanshinone IIA    | 0.6            |
of variation of baseline diameter, maximal diameter, and FMD in our laboratory were 3.7, 4.0, and 14 %.

Randomization and blinding
Allocation to the herb or placebo supplementation was based on a computer-generated randomization list that was prepared by a study-independent researcher. Block-randomization with a block size of four was used; the group assignment was concealed in an envelope and revealed after all data analysis was performed. During study, researchers were blinded and unaware of subjects’ allocation. Subjects were instructed not to reveal any information regarding supplement and exercise treatment during intervention.

Statistical analyses
Descriptive statistics were used for the analyses of subject characteristics using SPSS statistical package (version 16.0; Chicago, IL). Dependent variables were analyzed within each treatment to determine the time effect (pre, 90 min, 24 h, 48 h, and 72 h post exercise) using repeated measures ANOVA. A 2-way mixed model ANOVA was used for analyses of time and treatment effects. Bonferroni post-hoc analysis was performed when significance was achieved. Associations were determined by Spearman rank correlations. To detect difference in cfPWV of 5 % at a SD of 4.5 % unit change with 80 % power (α was set at 0.05) and accounting for a 10 % attrition rate, a total of 24 subjects were to be recruited and tested.

Results
After screening, twenty-four subjects were eligible and entered into the study (Fig. 2); one subject withdrew from the study during intervention due to time commitment unrelated to the study. Selected subject characteristics are presented in Table 2. All the subjects were non-obese, normolipidemic, and normotensive. There were no significant differences in body composition, lipid profile, and baseline hemodynamic parameters between the placebo and the Chinese herb groups.

An acute bout of downhill running exercise increased muscle soreness significantly at 90 min, 24 and 48 h post exercise (Fig. 3). Active range of motion decreased significantly at 24 and 48 h post exercise in both groups, but remained significant at 72 h in the placebo group (Fig. 3). As shown in Fig. 4, the magnitude of the increase in plasma CK concentration was significantly greater in the placebo group than in the herb group.

An acute bout of eccentric exercise increased cfPWV at 24 h post exercise in the placebo group (Fig. 5). No such significant increase in cfPWV was observed in the herb group. There were no changes in blood pressure in either group (Table 3). As shown in Fig. 6, there were no significant changes in FMD.
Plasma CRP concentration increased significantly at 24 h post eccentric exercise in both groups (Table 4). Plasma TBARs and TNF-α concentrations did not change in either group. Plasma IL-6 concentration increased significantly at 90 min after eccentric exercise in the placebo group.

The associations between changes in cfPWV and changes in selected biomarkers in combined groups are shown in Table 5. Changes in IL-6 were associated with changes in cfPWV at 48 h post exercise ($r = 0.46$, $P < 0.05$). In addition, changes in TNF-α were associated with changes in cfPWV at 24 to 48 h after exercise ($r = 0.57 - 0.60$, $P < 0.05$).

**Discussion**
The major findings of this study are as follows. Seven days of herb supplementation of *panax ginseng* and *salvia miltiorrhiza* prior to downhill running exercise did not affect muscle soreness, but prevented the significant and transient increase in arterial stiffness, and facilitated the recovery of active range of motion induced by muscle damage. This ‘destiffening’ effect was independent of blood
pressure changes as arterial pressure did not change in either group. A lack of changes in arterial stiffness with the herb supplementation was in part associated with the attenuated increases in inflammatory markers. These results suggest that the Chinese herb supplementation may be an effective strategy to minimize the delayed onset vascular stiffening induced by eccentric exercise.

The presence of elevated plasma CK has been recognized a marker of increased sarcolemma permeability or muscle damage resulted from unaccustomed exercise or eccentric muscle contractions [31, 35, 36]. In the present study, an acute bout of downhill running exercise increased plasma CK concentration significantly following eccentric exercise. The increase in plasma CK concentration was greater in the placebo group at 24 h post exercise than in the herb group. Previous studies in animal models reported that North American ginseng, panax ginseng or panax quinque-folius, decreased plasma CK levels after eccentric exercise [15, 37] and preserved mitochondria integrity [14] and decreased macrophage infiltration [16] in skeletal muscle. One recent human study [38] also found that American ginseng decreased CK concentration at 72 h post exercise when compared with the placebo control. Collectively, these results suggest that Chinese herb supplementation may have reduced the amount of muscle damage induced by eccentric exercise.

In the present study, arterial stiffening effects were attenuated when the subjects were supplemented with Chinese herbs for 7 days before the unaccustomed exercise was performed. These effects were not related to changes in blood pressure in either group. To our knowledge, this is the first study to demonstrate that Chinese herb supplementation effectively prevented the adverse effects on vascular stiffening induced by eccentric exercise. Specifically, downhill running increased arterial stiffness significantly as early as 24 h after the eccentric exercise in the placebo group. In the previous study [10] including ours [9], the increase in arterial stiffness was somewhat delayed showing up at 48 h after exercise. The exact reasons for the slight differences in the time course are not known. But compared with our previous study [9] that used localized resistance exercise as an eccentric stimulus, in the present study, we used more systemic downhill running exercise that elicit greater and more robust responses [39]. Moreover, compared with the previous study that utilized moderately-trained young adults [10], the subjects in the current study were mainly sedentary adults. Current literature indicates that people who repeatedly perform eccentric muscle contractions demonstrate so-called “repeated bout effect” or adaptation in skeletal muscle that inflammatory and muscle biomarker responses were substantially lowered in the

Table 2

|                     | Placebo | Herb  |
|---------------------|---------|-------|
| (n = 11)            | (n = 12) |
| Age, yr             | 24 ± 1  | 26 ± 5 |
| Height, cm          | 173 ± 1 | 174 ± 3 |
| Body mass, kg       | 68 ± 2  | 68 ± 3 |
| BMI, kg/m²          | 23 ± 1  | 22 ± 1 |
| Body fat percentage, % | 19 ± 1  | 18 ± 2 |
| Waist-hip ratio     | 0.85 ± 0.01 | 0.85 ± 0.01 |
| Heart rate at rest, bpm | 67 ± 3  | 58 ± 3 |
| Systolic BP, mmHg    | 118 ± 2 | 112 ± 3 |
| Diastolic BP, mmHg   | 66 ± 2  | 62 ± 2 |
| VO2peak, ml/kg/min  | 47 ± 2  | 47 ± 2 |
| HDL cholesterol, mg/dL | 53 ± 2  | 53 ± 3 |
| LDL cholesterol, mg/dL | 97 ± 9  | 86 ± 5 |
| Total cholesterol, mg/dL | 180 ± 9 | 192 ± 8 |
| Triglyceride, mg/dL  | 78 ± 9  | 58 ± 8 |
| HbA1C, %            | 5.4 ± 0.1 | 5.4 ± 0.1 |

Values are means ± SEM

BMI: body mass index, BP: blood pressure, VO2peak: peak oxygen consumption, HbA1c: glycosylated hemoglobin A1c

![Fig. 3](image-url) Delayed onset muscle soreness (a) and active range of motion (AROM) (b) following downhill running exercise. *P < 0.05 vs. Pre in the same condition.
following challenge [1], suggesting that exercise training status plays a role in determining the response to this exercise challenge.

The underlying mechanisms by which muscle damage induced by eccentric exercise results in increased arterial stiffness remain unclear. However, arterial stiffening following eccentric exercise has been associated with increases in subjective muscle soreness [10], a marker of muscle damage (i.e., plasma CK) [9]. In the present study, plasma IL-6 levels increased significantly after eccentric exercise in the placebo group. Previous studies have demonstrated that the acute inflammation induced by a vaccination increased not only IL-6 and CRP concentrations [40, 41] but also arterial stiffness [42]. Collectively, these results suggest that arterial stiffening induced by muscle damage is associated with markers of muscle damage and/or systemic inflammation.

Consistent with this concept, we found that Chinese herb supplementation minimized the increases in IL-6 levels. Additionally, changes in arterial stiffness were

|                         | Pre  | 90 min | 24 h  | 48 h  | 72 h  |
|-------------------------|------|--------|-------|-------|-------|
| **cfPWV, cm/s**         |      |        |       |       |       |
| Placebo Control         | 530 ± 19 | 535 ± 17 | 546 ± 16 | 547 ± 15 | 513 ± 13 |
| Exercise                | 531 ± 12 | 548 ± 19 | 577 ± 20* | 570 ± 24 | 552 ± 22 |
| Herb Control            | 482 ± 14 | 491 ± 17 | 490 ± 14 | 487 ± 13 | 505 ± 19 |
| Exercise                | 523 ± 20 | 497 ± 17 | 503 ± 16 | 500 ± 13 | 505 ± 23 |
| **Heart rate, bpm**     |      |        |       |       |       |
| Placebo Control         | 63 ± 4  | 58 ± 3  | 62 ± 3  | 66 ± 3  | 65 ± 3  |
| Exercise                | 62 ± 3  | 69 ± 4* | 66 ± 3  | 63 ± 3  | 59 ± 5  |
| Herb Control            | 57 ± 3  | 54 ± 1  | 56 ± 3  | 59 ± 4  | 55 ± 3  |
| Exercise                | 56 ± 2  | 65 ± 4* | 56 ± 3  | 55 ± 2  | 53 ± 3  |
| **Systolic BP, mmHg**   |      |        |       |       |       |
| Placebo Control         | 119 ± 2 | 116 ± 2 | 118 ± 2 | 119 ± 2 | 119 ± 2 |
| Exercise                | 119 ± 2 | 114 ± 2 | 120 ± 2 | 120 ± 3 | 117 ± 2 |
| Herb Control            | 114 ± 2 | 111 ± 2 | 114 ± 2 | 115 ± 3 | 115 ± 2 |
| Exercise                | 116 ± 3 | 114 ± 3 | 115 ± 2 | 112 ± 2 | 111 ± 3 |
| **Diastolic BP, mmHg**  |      |        |       |       |       |
| Placebo Control         | 66 ± 2  | 67 ± 1  | 67 ± 2  | 64 ± 2  | 67 ± 2  |
| Exercise                | 66 ± 3  | 65 ± 2  | 65 ± 3  | 66 ± 2  | 64 ± 2  |
| Herb Control            | 63 ± 3  | 63 ± 1  | 62 ± 2  | 62 ± 1  | 62 ± 1  |
| Exercise                | 63 ± 3  | 61 ± 2  | 62 ± 1  | 63 ± 3  | 60 ± 1  |
| **Pulse pressure, mmHg**|      |        |       |       |       |
| Placebo Control         | 52 ± 2  | 49 ± 1  | 51 ± 2  | 55 ± 2  | 53 ± 2  |
| Exercise                | 53 ± 2  | 48 ± 2  | 55 ± 2  | 53 ± 2  | 53 ± 1  |
| Herb Control            | 51 ± 3  | 48 ± 3  | 52 ± 3  | 54 ± 3  | 53 ± 3  |
| Exercise                | 53 ± 3  | 52 ± 3  | 53 ± 2  | 49 ± 3  | 51 ± 3  |

Values are means ± SEM

*P<0.05 vs. Pre in the same condition

**Fig. 4** Relative changes in serum creatine kinase (CK) concentration in response to eccentric exercise sessions. *P<0.05 vs. Pre in the same condition. †P<0.05 vs. Placebo at the same time point.

**Table 3** Hemodynamic responses in control (seated rest) and eccentric exercise sessions.

**Fig. 5** Effects of Chinese herb supplementation on carotid-femoral pulse wave velocity (cfPWV). *P<0.05 vs. Pre. †P<0.05 vs. Herb supplementation.

**Fig. 6** Changes in flow-mediated vasodilatation in response to the eccentric exercise.
Changes in muscle damage markers, inflammatory and oxidative stress markers in response to downhill running exercise in the placebo and herb group

|          | Placebo | Pre | 90 min | 24 h | 48 h | 72 h |
|----------|---------|-----|--------|------|------|------|
| **CRP, mg/dL** |         |     |        |      |      |      |
| Control   | 0.07 ± 0.04 | 0.07 ± 0.04 | 0.07 ± 0.03 | 0.06 ± 0.02 | 0.05 ± 0.02 | |
| Exercise  | 0.08 ± 0.02 | 0.11 ± 0.05 | 0.15 ± 0.05* | 0.11 ± 0.05 | 0.10 ± 0.03 | |
| **CK, U/L**  |         |     |        |      |      |      |
| Control   | 94 ± 10 | 98 ± 8 | 89 ± 8 | 87 ± 9 | 90 ± 11 | |
| Exercise  | 93 ± 8  | 126 ± 13 | 396 ± 72** | 257 ± 48** | 192 ± 32* | |
| **Herb**  |         |     |        |      |      |      |
| CRP, mg/dL |         |     |        |      |      |      |
| Control   | 0.13 ± 0.05 | 0.14 ± 0.05 | 0.11 ± 0.04 | 0.11 ± 0.04 | 0.11 ± 0.03 | |
| Exercise  | 0.10 ± 0.03 | 0.09 ± 0.03 | 0.15 ± 0.05* | 0.10 ± 0.02 | 0.10 ± 0.02 | |
| CK, U/L   |         |     |        |      |      |      |
| Control   | 110 ± 12 | 108 ± 9 | 119 ± 17 | 109 ± 11 | 113 ± 16 | |
| Exercise  | 106 ± 9 | 195 ± 51* | 291 ± 35** | 204 ± 26** | 151 ± 20 | |
| **TBARs, µM** |         |     |        |      |      |      |
| Placebo   | 7.1 ± 1.1 | 8.1 ± 1.5 | 7.9 ± 1.0 | 6.5 ± 1.0 | -     | |
| Herb      | 7.0 ± 1.0 | 6.6 ± 0.8 | 8.1 ± 0.9 | 5.7 ± 0.6 | -     | |
| **IL-6, pg/ml** |         |     |        |      |      |      |
| Placebo   | 0.44 ± 0.1 | 0.69 ± 0.1* | 0.32 ± 0.1 | 0.46 ± 0.1 | -     | |
| Herb      | 0.50 ± 0.2 | 0.45 ± 0.1 | 0.23 ± 0.1* | 0.29 ± 0.1 | -     | |
| **TNF-α, pg/ml** |         |     |        |      |      |      |
| Placebo   | 0.45 ± 0.14 | 0.28 ± 0.06 | 0.30 ± 0.06 | 0.30 ± 0.06 | -     | |
| Herb      | 0.44 ± 0.06 | 0.34 ± 0.06 | 0.38 ± 0.06 | 0.34 ± 0.06 | -     | |

Values are means ± SEM. TBARs, IL-6, and TNF-α were measured only during the eccentric exercise session. CRP C-reactive protein, CK creatine kinase, TBARs thiobarbituric acid reactive substances, IL-6 interleukin-6, TNF-α tumor necrosis factor-α

*P<0.05 vs. Pre in the same condition. **P<0.05 vs. Control or Placebo at the same time point

Table 5 - Associations between relative changes (%) in arterial stiffness and selected biomarkers

|          | ΔcfPWV 24 h | ΔcfPWV 48 h |
|----------|-------------|-------------|
| ΔCK      | 0.05        | 0.08        |
| CRP      | 0.17        | 0.21        |
| ΔIL-6    | 0.11        | 0.46*       |
| ΔTNF-α   | 0.60*       | 0.57*       |
| ΔTBARs   | 0.09        | 0.36        |

ΔcfPWV carotid-femoral pulse wave velocity, CK creatine kinase, CRP C-reactive protein, IL-6 interleukin-6, TNF-α tumor necrosis factor-α, TBARs thiobarbituric acid reactive substances

*P<0.05

Conclusions

A short-term Chinese herb supplementation incorporating panax ginseng and salvia miltiorrhiza was effective in ameliorating the delayed onset vascular stiffening.
induced by acute eccentric exercise, possibly via the reductions in oxidative stress and systemic inflammation.

**Abbreviations**

BMI, body mass index; BP, blood pressure; cFPWV, carotid-femoral pulse wave velocity; CK, creatine kinase; CRP, C-reactive protein; HbA1c, glycated hemoglobin; IL-6, interleukin-6; TNF-a, tumor necrosis factor-a; VO2peak, peak oxygen consumption

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**Availability of data and materials**

The datasets supporting the conclusions of this article are presented in this paper.

**Authors’ contributions**

Study concept and design: HL, HT, CL, JL; data collection: HL, TK, CC; data analysis: HL, TK, CC, CL; data interpretation: HL, TK, JL, HT; manuscript drafting: HL, HT, revising manuscript critically for important intellectual content: HL, TK, CC, CL, JL, HT. All authors have read, edited, approved the final manuscript, and have agreed to be held accountable for all aspects of manuscript in ensuring that questions related to the accuracy or integrity of any part of manuscript are appropriately investigated and resolved.

**Competing interests**

The authors declare that they have no competing interests.

**Consent for publication**

Not applicable.

**Ethics approval and consent to participate**

All subjects gave their written informed consent prior to study participation and procedures were reviewed and approved by the Institutional Review Board of National Taiwan University Hospital.

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