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Effect of continuous positive airway pressure therapy on inflammatory cytokines and atherosclerosis in patients with obstructive sleep apnea syndrome

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Abstract. Obstructive sleep apnea syndrome (OSAS) is known to be a risk factor for atherosclerosis (AS), derived from a series of chronic inflammatory reactions caused by hypoxia. However, the association between chronic inflammation and high blood pressure caused by hypoxia remains to be fully elucidated. The aim of the present study was to investigate the effect of continuous positive airway pressure (CPAP) therapy on inflammatory cytokines and AS. A total of 100 patients with OSAS and 50 healthy control subjects were enrolled. Fresh venous blood samples were collected prior to and following a 3-months period of CPAP treatment. The inflammatory factors, interleukin (IL)-18 and tumor necrosis factor (TNF)-α, C-reactive protein (CRP), intercellular cell adhesion molecule 1 (ICAM-1), vascular cell adhesion molecule 1 (VCAM-1), E-selectin and P-selectin, were detected using standard enzyme-linked immunosorbent assay kits. Intima-media thickness (IMT), brachial-ankle pulse wave velocity (Ba-PWV), apnea-hypopnea index (AHI) and transcutaneous oxygen saturation (SpO2) were also detected to compare differences prior to and following treatment. The results showed that, compared with the pre-treatment data, the expression levels of IL-18, TNF-α, CRP, ICAM-1, VCAM-1, E-selectin and P-selectin were significantly decreased following treatment (P<0.05). The AHI, IMT, blood pressure and Ba-PWV values were significantly decreased (P<0.05), and the SpO2 was increased (P<0.05). Taken together, by comparing the pre- and post-intervention data, it was confirmed that inflammatory factors were involved in the process of AS in patients with OSAS. Following CPAP treatment, blood pressure and primary indicators in the patients improved.

Introduction

Obstructive sleep apnea syndrome (OSAS), which affects 2-4% of the population worldwide, is a common sleep disorder and potential clinical risk factor. OSAS has been widely considered as a severe problem associated with a range of pathological consequences, including hypoxia (1), hypercapnia (2), imbalance of nerve regulation function (3), activation of the rennin angiotensin aldosterone system, catecholamine and endothelin secretion, endocrine dysfunction and hemodynamic changes (4). Of all the problems reported in patients with OSAS, the most important consequence is the risk of cardiovascular and cerebrovascular disease (5). OSAS has long been confirmed as one of the most important independent risk factors for atherosclerosis (AS) and hypertension (6). Tissue hypoxia followed by chronic inflammation damage and the involvement of a variety of inflammatory cytokines leads to AS (7).

Continuous positive airway pressure (CPAP) therapy is the most effective method for OSAS (8). Adequate CPAP treatment can increase pulmonary ventilation and ameliorate the inflammation of arteries in patients with OSAS (9). Patients show improvements in blood oxygen concentration on the first day following CPAP treatment, and further benefits are observed during an extended course of treatment (10). The molecular and immunological mechanisms underlying this type of therapy are a major concern when selecting CPAP in patients with OSAS.

According to clinical observations and previous investigations, the present study hypothesized that, in patients with OSAS without clinical interventions, the plasma levels of interleukin (IL)-18, tumor necrosis factor (TNF)-α (11), C-reactive protein...
(CRP), intercellular cell adhesion molecule 1 (ICAM-1), vascular cell adhesion molecule 1 (VCAM-1), E-selectin and P-selectin are significantly altered. The present study investigated these factors and compared changes following CPAP (12). On the basis of these observations, alterations in the clinical characteristics of OSAS following treatment were observed. In order to evaluate the effect of CPAP (5) on inflammatory factors and AS, the present study also examined the carotid intima-media thickness (IMT) and brachial-ankle pulse wave velocity (Ba-PWV) (13). The aim of the present study was to evaluate the effects of a longer (3-month) period of CPAP treatment on oxygen ventilation in patients with OSAS.

Subjects and methods

Subjects and healthy donors. In the present study, a total of 100 patients were recruited from The Affiliated Jiangning Hospital of Nanjing Medical University (Nanjing, China) upon diagnosis on interpretation of clinical guidelines on the treatment of OSAS (American College of Physicians, 2013) (14). The patients included 82 men and 18 women, aged between 43 and 69 years old (mean age, 55.28±7.12 years). These patients were prospectively recruited on preparing to attend the Sleep Clinic for overnight polysomnography (PSG) between March 2014 and March 2016, who met the standard definition of an apnea-hypopnea index (AHI) ≥5. None of the patients were suffering from serious disease requiring treatment. As a healthy control group, 50 individuals undergoing physical examination at The Affiliated Jiangning Hospital of Nanjing Medical University were invited to join in this study sequence. There were no differences in age, gender or body mass index (BMI) between the healthy control group and the OSAS group.

The exclusion criteria included patients or healthy donors with coronary heart disease, valvular heart disease, cardiomyopathy, tumors, severe liver and kidney dysfunction, severe lung disease, hyperlipidemia, diabetes, hypertension, infection or trauma, or had undergone surgery in the previous 2 weeks. The present study was approved by the Ethics Committee of The Affiliated Jiangning Hospital of Nanjing Medical University. Written informed consent was obtained from each patient and healthy donor.

Specimen collection and detection of inflammatory factors. Blood samples were collected from the study subjects in the early morning (6:00 a.m.). Fasting blood was collected again from the patients with OSAS following CPAP treatment for 3 months (again at 6:00 a.m.). All blood samples were 10 ml in volume and obtained from the elbow vein. The samples underwent 300 x g centrifugation at 10°C for 10 min, and the upper plasma was stored at -80°C. IL-18 and TNF-α, CRP, ICAM-1, VCAM-1, E-selectin and P-selectin were detected using an enzyme-linked immunosorbent assay (ELISA). The concentrations of the inflammatory factors were measured using human ELISA kits (Amersham; GE Healthcare Life Sciences, Chalfont, UK).

CPAP treatment. The recording and comparing of sleep monitoring parameters during the 3-month treatment period were performed using a compumedics PSG monitoring system (ResMed, Sydney, Australia). The AHI and transcutaneous oxygen saturation (SpO2) were detected and recorded prior to and following the 3-month treatment period.

Ultrasound measurement of carotid IMT. The carotid artery IMT was detected using the Mylab 90 ultrasonic instrument (Esaote, Genoa, Italy) as the quality of IMT.

Detection of Ba-PWV. The Ba-PWV was detected using an automatic AS testing equipment (BP-203RPEII; Omron Colin, Tokyo, Japan).

The upper arm to the ankle propagation distance (L) and the pulse wave transit time (T) were automatically measured according to the height of the body. The formula Ba-PWV=L/T was calculated from Ba-Pwv on both sides, and the average value of both sides was recorded.

Statistical analysis. The statistical analyses performed included a matched t-test of measurement data and a rank-sum test using SPSS 16.0 software (SPSS, Inc., Chicago, IL, USA). All data are presented as the mean ± standard deviation. P<0.05 was considered to indicate a statistically significant difference.

Results

General characteristics of study subjects. All 100 patients and 50 healthy controls were included in the study. Table I lists the characteristics of the subjects. No differences in gender, age or BMI were found between the two groups (P>0.05). The baseline clinical data of the subjects are also listed in Table I.

Inflammatory factors in OSAS. Compared with the healthy control group, the expression levels of IL-8, TNF-α, CRP, ICAM-1, VCAM-1, E-selectin and P-selectin in the patients with OSAS were significantly increased. The respective statistical comparisons of expression in the healthy controls, vs. patients with OSAS were as follows: IL-8, 13.28±3.15, vs. 40.72±1.60 pg/ml (P=0.0013); TNF-α, 29.15±1.74, vs. 37.67±0.21 pg/ml (P=0.0037); CRP, 6.37±1.30, vs. 49.64±21.66 mg/l (P=0.0007); ICAM-1, 291.68±53.29, vs. 357.92±10.52 µg/l (P=0.0170); VCAM-1, 288.16±48.81, vs. 351.06±53.61 µg/l (P=0.0201); E-selectin, 42.57±10.4, vs. 50.65±8.29 µg/l (P=0.0233); P-selectin 30.26±6.80 and 54.79±3.34 µg/l (P=0.0149). The results are shown in Fig. 1.

Changes of inflammatory cytokines following treatment. PSG was performed on patients treated with CPAP using an automatically programmed PSG system following 3 months of treatment. Compared with the pre-therapy data, the expression levels of IL-8, TNF-α, CRP, ICAM-1, VCAM-1, E-selectin and P-selectin were significantly decreased following treatment. The statistical comparisons of post-treatment, vs. pre-treatment data were as follows: IL-8, 35.79±1.63 and 40.72±1.60 pg/ml (P=0.0392); TNF-α, 34.97±3.18 vs. 37.67±0.21 pg/ml (P=0.0412); CRP, 27.41±8.86 vs. 49.64±21.66 mg/l (P=0.0138); ICAM-1, 338.29±43.03 vs. 357.92±10.52 µg/l (P=0.0219); VCAM-1, 322.36±38.25 vs. 357.92±10.52 µg/l (P=0.0019); E-selectin, 47.46±8.58 vs. 50.65±8.29 µg/l (P=0.0143); P-selectin, 44.05±6.97 vs. 54.79±3.34 µg/l (P=0.0019). The results are shown in Fig. 2.
AHI and Ba-Pwv. Following CPAP treatment, the AHI and Ba-Pwv improved significantly. As shown in Fig. 3A, the average value of AHI decreased significantly (37.80±6.70 vs. 26.73±4.34; P=0.0019), as did that of Ba-Pwv (1,418.86±199.58 vs. 1,265.31±219.36; P=0.0239; Fig. 3).

IMT and SPO$_2$. Compared with pre-treatment, the IMT following treatment was significantly reduced (1.35±0.55 vs. 1.12±0.52, respectively; P=0.0381), whereas SPO$_2$ was significantly increased (89.18±7.23 vs. 89.18±5.19, respectively; P=0.0283), as shown in Fig. 4A and B.

SBP and DBP. In the patients whose blood pressure remained high despite medication (n=78), the blood pressure was improved following treatment. As is shown in Fig. 5A and B, the decreases in SBP (168.84±32.57 vs. 144.29±17.85; P=0.0013) and DBP (84.21±11.85 vs. 79.47±8.98; P=0.0021) were statistically significant.

### Discussion

In previous years, OSAS has been widely accepted as a mechanism for hypertension, and specific treatment for this change has also been considered. In the present study, CAPA was used for the treatment of patients with OSAS and hypertension, and the efficacy was observed in order to examine the novel treatment approach for patients with OSAS and hypertension.

The pro-inflammatory cytokine IL-18 is important in the occurrence and development of atherosclerotic plaque rupture (15). An epidemiological follow-up study revealed that IL-18 is an independent risk factor for coronary events, and is also a predictor of life-threatening cardiac events in patients with acute coronary syndrome (10). TNF-α is critical in the

<table>
<thead>
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<th>Variable</th>
<th>OSAS</th>
<th>Healthy control</th>
<th>P-value</th>
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<tbody>
<tr>
<td>Number</td>
<td>100</td>
<td>50</td>
<td>ND</td>
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<tr>
<td>Male/female</td>
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<td>Age (years)</td>
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<td>BMI (kg/m$^2$)</td>
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<td>SBP (mmHg)</td>
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<td>DBP (mmHg)</td>
<td>95.32±5.515</td>
<td>83.42±4.216</td>
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<tr>
<td>TC (mmol/l)</td>
<td>6.71±2.998</td>
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<td>AHI (/h)</td>
<td>38.01±8.040</td>
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<td>SPO$_2$ (%)</td>
<td>89.18±7.234</td>
<td>95.27±2.490</td>
<td>0.018</td>
</tr>
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</table>

OSAS, obstructive sleep apnea syndrome; BMI, body mass index; SBP, systolic blood pressure; DBP, diastolic blood pressure; TC, total cholesterol; AHI, apnea hypopnea index; SPO$_2$, pulse oxygen saturation; ND, no data.
inflammatory cascade reaction of AS. Although CRP cannot predict disease specificity, it is an important inflammatory response factor in the majority of types of chronic inflammation, particularly in AS. In addition, inflammatory factors, including ICAM-1, VCAM-1 (16), E-selectin and P-selectin, have been found to be important in predicting the occurrence and development of AS (17). The pathological basis of AS involves chronic inflammation of the vascular wall (18), however, whether inflammation is also involved in the pathogenesis of cardiovascular disease in patients with OSAS remains to be fully elucidated. The present study hypothesized that hypoxia causes chronic inflammation in patients with OSAS, and chronic inflammation leads to AS and hypertension, whereas mechanical ventilation improves the condition of hypoxia and the inflammatory response in patients.

The results of the present study showed that IL-18, TNF-α, high-sensitivity CRP, and serum levels of ICAM-1, VCAM-1, E-selectin and P-selectin in the OSAS group were significantly increased, compared with those in the healthy control group. It was indirectly confirmed that these inflammatory factors may be involved in hypoxia and in vascular sclerosis. In order to directly confirm this mechanism, CPAP treatment and follow-up investigations were performed on patients with OSAS. Following CPAP treatment, significant differences
were found in these inflammatory markers, compared with those at the pre-treatment stage.

In patients with OSAS with significant characteristics of upper airway collapse (19) and effects on the circulatory system, the primary mechanism involves the excitement of sympathetic nerves (3), increase of endothelin levels (20), vascular endothelial function, abnormalities in vascular active substances (21), vascular tension (22) and excessive renin secretion causing sustained hypertension (23). In previous studies, plasma levels of CRP and other inflammatory cytokines in patients with OSAS caused by inflammation have been reported to promote cardiovascular and cerebrovascular diseases (24), including AS. The overexpression of IL-18, TNF-α, E-selectin and P-selectin in OSAS is an important factor in the occurrence and development of hypertension (25).

It has been reported that OSAS is closely associated with the occurrence and development of hypertension (22). Under the condition of low oxygen, substantial cellular metabolic waste accumulation leads to the generation of oxygen free radicals (26). Free radicals react with the unsaturated fatty acids in cells (27), in a process called lipid peroxidation, generating cytotoxic effects of peroxide (28). Oxyradicals and subsequent lipid peroxidation lead to various types of damage in the body. In the electron transport chain of the redox reaction, co-enzyme Q in complex III can produce high activity free radical intermediates in the process of reduction (29). The reactive oxygen species generated in these cells include hypochlorite, hydrogen peroxide, and free radicals, including superoxide anions and hydroxyl radicals (30). The chemical properties of hydroxyl radical are unstable and can be divided into specific biological molecules. These biological macromolecules are predominantly produced by the catalytic reduction of hydrogen peroxide by metal enzymes. The oxidant can trigger a chain reaction, for example, lipid peroxidation or oxidation of DNA and protein, resulting in cell damage. Damaged DNA can cause mutations and induce several diseases if not repaired. The damage causes the degradation of protein and inhibits the activity of the enzyme. CPAP can improve patient ventilation status, and subsequently improve in the long-term chronic hypoxia status, with reductions in the redox reaction and generation of free radicals (31).

In the present study, in addition to the changes of inflammatory factors and clinical indices of the patients, AHI, IMT, Ba-Pwv (32) and blood pressure were detected. Patients with OSAS and hypertension exhibited significantly higher IMT and Ba-Pwv, compared with the normal population. Following treatment of the patients with OSAS, blood pressure (SBP and DBP) and Pso2 was decreased. This showed that CAPA alleviated the changes in OSAS patients with hypertension.

In conclusion, CPAP increases oxygen supply, improves chronic inflammation in OSAS, and inhibits the expression of inflammatory factors and other factors promoting cardiovascular and cerebrovascular diseases, in addition significantly improving to pulmonary function in patients (33). In the present study, it was confirmed that CPAP inhibited the inflammatory response of patients with OSAS and hypertension, and inhibited the pathological basis of OSAS. Therefore, the patients with OSAS exhibited hypertension and hypoxia, leading to inflammation. CPAP indirectly improved AS and high blood pressure in the patients. However, due to the complexity of clinical investigation and confounding factors, there are still some deficiencies in the present study. More cases and longer follow-up are needed to investigate the association between OSAS, chronic inflammation and hypertension.

References


