The Difference of Lower Urinary Tract Symptoms Between Sympathetic Hyperactive and Hypoactive Men

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Purpose: Heart rate variability (HRV) is a tool used to measure autonomic nervous function; however, there is no evidence that it can be used to define sympathetic hyperactivity in men with lower urinary tract symptoms (LUTS). We suspected that LUTS would differ between sympathetic hyperactive and hypoactive patients. Therefore, we measured HRV and divided the LUTS patients into two groups, a sympathetic hyperactive group and a sympathetic hypoactive group according to the low frequency/high frequency (LF/HF) ratio and made clinical comparisons between the groups.

Methods: A total of 43 patients with symptomatic LUTS (International Prostate Symptom Score [IPSS] over 8) and 49 healthy volunteers were enrolled. No subjects had diseases that could affect the autonomic nervous system, such as diabetes or hypertension. Electrocardiographic signals were obtained from subjects in the resting state and HRV indexes were calculated with spectral analyses. We divided the LUTS patients into two groups by an LF/HF ratio of 1.9, which was the median value in the healthy volunteers, and compared the differences in clinical characteristics, IPSS, prostate-specific antigen (PSA), and transrectal ultrasound (TRUS) results. The parameters were compared by independent sample t-test by use of SPSS ver. 19.

Results: There were no significant differences in age, serum PSA, or volume of the prostate between the 2 LUTS groups. However, analyzing IPSS questionnaires between two groups showed that there were significant differences in mean of Q2 score (frequency) and storage symptom score ([Q2+Q4+Q7]/3) (P<0.05).

Conclusions: We suggest that an imbalance of autonomic nervous system activity may be a factor that evokes varieties of symptoms in men with LUTS. LUTS patients with hypoactive sympathetic tone may suffer from frequency and storage symptoms.

Keywords: Prostatic hyperplasia; Autonomic nervous system; Lower urinary tract symptoms; Heart rate variability
MATERIALS AND METHODS

We selected 43 patients who visited an outpatient clinic for LUTS and 49 healthy volunteers who visited the family medicine outpatient clinic for a general health examination. No subjects had diseases that could affect the ANS, such as diabetes, hypertension, coronary heart disease, or any malignancy. No subjects had a history of surgery, irradiation, or pelvic trauma. All subjects answered an International Prostate Symptom Score (IPSS) questionnaire and patients with an IPSS ≥ 8 were enrolled. We divided the LUTS patients into two groups (groups 1 and 2) by a low frequency/high frequency (LF/HF) ratio of 1.9, which was the median value in the healthy volunteers, and compared the clinical characteristics, IPSS, prostate-specific antigen (PSA), and transrectal ultrasound (TRUS) results between the two groups.

HRV Measurement

HRV was measured by using 2 methods: time domain and frequency domain analysis. The parameters of the time domain analysis were the mean heart rate in bpm, the standard deviation of the N-N interval (SDNN), and the square root of the mean squared difference of successive N-N intervals (RMSSD). SDNN reflects all of the cyclic components responsible for variability in the recording period and RMSSD represents parasympathetic activity. The parameters of the frequency domain analysis were total power (TP), very low frequency (VLF), LF, HF and LF/HF ratio. The LF peak of the spectrum (0.15 to 0.40 Hz) represents parasympathetic activity, whereas the LF peak (0.04 to 0.15 Hz) represents sympathetic activity. Thus, the LF/HF ratio represents the ANS balance (sympathetic to parasympathetic).

We used a system developed in-house for HRV acquisition and signal processing to measure ANS activity. We confirmed that bladders were filled with more than 100 mL of urine before testing by ultrasonography to represent the physiological state. All subjects were restricted in their consumption of tea and coffee, cigarette smoking, and medication use (e.g., beta-receptor agonists or antagonists). After 30 minutes of rest, each patient underwent electrocardiographic signal recording (SA-3000P, Medicore Co., Seoul, Korea) for 5 minutes while sitting. With the patients breathing normally, we calculated the mean heart rate, SDNN and RMSSD. We also determined the resting sympathetic and parasympathetic heartbeat rate modulation by using frequency domain methods, including TP, VLF, LF, HF, and LF/HF ratio. HRV parameters were compared by using the independent sample t-test.

Statistical Analysis

Statistics were performed by using the IBM SPSS ver. 19.0 (IBM Co., Armonk, NY, USA). The data were presented as means ± standard deviations. Comparisons between the two groups were performed with the independent t-test. P-values less than 0.05 were considered to be statistically significant.

RESULTS

The statistical analysis showed no significant differences between the control (healthy volunteers) and LUTS group in age, body mass index, serum PSA, or volume of the prostate (Table 1). Sympathetic hypoactive LUTS patients who LF/HF ratio below 1.9 were group 1, and sympathetic hyperactive LUTS patients who LF/HF ratio over 1.9 were group 2. The results of the analysis of the parameters of the IPSS questionnaire in LUTS patients (groups 1 and 2) are shown in Table 2. There were significant differences in mean of IPSS Q2 score (frequent urination) and storage symptom score ([Q2+Q4+Q7]/3) of IPSS according to ANS balance (P < 0.05). Sympathetic hypoactive patients suffer from more severe storage symptoms especially frequency than sympathetic hyperactive patients.

DISCUSSION

The extended underlying pathophysiology leading to LUTS remains unclear, although it is increasingly recognized that the causes of LUTS extend well beyond prostate enlargement and

Table 1. Clinical characteristics between control and LUTS group

|                         | Control group (n = 49) | LUTS group (n = 43) | P-value |
|-------------------------|------------------------|---------------------|---------|
| Age (yr)                | 61.4 ± 10.4            | 58 ± 7.8            | 0.40    |
| Body mass index (kg/m²) | 24.5 ± 3.4             | 24.3 ± 3.0          | 0.87    |
| Total IPSS              | 23 ± 8.1               | 17.3 ± 6.9          | 0.09    |
| PSA (ng/mL)             | 2.67 ± 2.0             | 1.45 ± 1.5          | 0.15    |
| TRUS (mL)               | 31.41 ± 9.1            | 32.7 ± 13.8         | 0.82    |

Values are presented as mean ± standard deviation. LUTS, lower urinary tract symptoms; IPSS, International Prostate Symptom Score; PSA, prostate-specific antigen; TRUS, transrectal ultrasonography.
Table 2. The results of characteristics and IPSS parameters in two groups of LUTS patients by sympathetic activity

| Variable            | Group 1  | Group 2  | P-value |
|---------------------|----------|----------|---------|
|                     | (n = 24) | (n = 19) |         |
| Age (yr)            | 59.6 ± 9.3 | 58.8 ± 7.8 | 0.25    |
| PSA (ng/mL)         | 1.77 ± 1.68 | 1.46 ± 1.73 | 0.67    |
| TRUS (mL)           | 29.4 ± 9.1 | 32.2 ± 12.0 | 0.21    |
| IPSS parameters     |          |          |         |
| Q1                  | 3.46 ± 1.77 | 2.58 ± 1.84 | 0.27    |
| Q2                  | 3.67 ± 1.63 | 2.79 ± 1.62 | 0.04a   |
| Q3                  | 3.08 ± 1.86 | 2.95 ± 1.81 | 0.87    |
| Q4                  | 2.83 ± 1.99 | 2.32 ± 1.89 | 0.74    |
| Q5                  | 4.08 ± 1.28 | 3.79 ± 1.32 | 0.54    |
| Q6                  | 3.13 ± 1.85 | 2.53 ± 1.35 | 0.41    |
| Q7                  | 2.83 ± 1.49 | 2.34 ± 1.46 | 0.35    |
| Total IPSS score    | 23.08 ± 7.48 | 19.26 ± 7.37 | 0.08    |
| Quality of life     | 4.58 ± 0.97 | 4.16 ± 0.90 | 0.15    |
| Voiding Sx score    | 3.29 ± 1.35 | 2.90 ± 1.36 | 0.39    |
| Storage Sx score    | 3.02 ± 1.44 | 2.11 ± 1.26 | 0.03a   |

Values are presented as mean ± standard deviation.

Group 1, patients with LF/HF below 1.9; Group 2, patients with LF/HF 1.9 and over 1.9; IPSS, International Prostate Symptom Score; PSA, prostate-specific antigen; TRUS, transrectal ultrasonography; Sx, symptom; Q, question number.

*P < 0.05.

bladder outlet obstruction [9]. The multiple pathways leading to the onset and progression of LUTS not only complicate diagnosis, but also limit the overall effectiveness and satisfaction of targeted symptom management strategies [10]. The lower urinary tract is generally under the control of the ANS, is innervated by 3 sets of peripheral nerves (parasympathetic, sympathetic, and somatic nervous systems), and contains afferent and efferent motor axons [11].

HRV depends on the influence of sympathetic and vagal activity on the sinus node, and variability reflects spontaneous changes in autonomic activity [12]. HRV is a simple and important tool for studying the autonomic control of the heart and autonomic dysfunction, and it represents one of the most promising of such markers [13,14]. As a development or progression, we suggest the possibility of ANS imbalance and use HRV as a method to evaluate ANS tone.

Previous epidemiological studies in men with BPH have suggested that a relationship between may exist between sympa-

thetic overactivity and LUTS. Meigs et al. [15] noted that men with symptomatic BPH were also more likely to have characteristics known to be associated with increased sympathetic activation, specifically heart disease, β-blocker use, and a sedentary lifestyle. Hammarsten and Hogstedt [16] also found a relationship between more rapidly developing BPH and diseases including obesity, increased insulin, increased risk of type 2 diabetes and hypertension.

Choi et al. [17] reported ANS activity in men with voiding symptom-predominant LUTS and storage symptom-predominant LUTS. They showed that patients with LUTS exhibited decreased HF than healthy controls, which indicated that they may had some disease or imbalance in the ANS, that may distinguish LUTS patients from healthy men. However, the LF/HF ratio, which represents ANS balance, was not significantly different between LUTS patients and healthy controls. Thus, in the present study, we divided the LUTS patients into two groups by the median LF/HF value of the healthy controls. Because most investigators believe that LF and HF represent sympathetic and parasympathetic nervous system activity, respectively. Previous studies LUTS patients had an increased sympathetic tone, and voiding symptom predominant patients had more increased sympathetic tone. But our results suggest that LUTS patients could have relatively hypoactive sympathetic tone than healthy men, and they suffered from storage symptom more than those with hyperactive sympathetic tone, especially frequency.

Unfortunately, no specific modalities can be used to improve autonomic dysfunction. In some studies, HRV was improved by exercise or cessation of smoking. Accordingly, it may be beneficial for physicians to advise their patients to perform more exercise and to quit smoking [18,19].

This study was limited in that LUTS are not specific and can be changed by factors unrelated to prostatic conditions, including diet, fluid intake, alcohol intake, anticholinergic effects of commonly used over-the-counter medications, smoking and even mood or physical activity [20-22]. We attempted to reduce such effects by asking patients to not consume anything that could affect the ANS before the HRV recording. Second, the sample size of both groups was small: thus, additional study with large numbers will be needed. Last, electrocardiography recording was measured only for 5 minutes in a sitting position and not for 24 hours to simplify the measurement and to reduce the discomfort of the subjects.

Many patients suffer from LUTS that do not respond to current medications or surgical treatments, and we suggest that an
imbalance in ANS activity may be a factor that cause or progression of diseases in men with LUTS. In particular, LUTS patients with hypoactive sympathetic tone may suffer from frequency and storage symptom by ANS dysfunction. We believe it is time to shift our concern to general health and ANS dysfunction to find the other etiologies of LUTS.

CONFLICT OF INTEREST

No potential conflict of interest relevant to this article was reported.

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