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## Electroacupuncture for Treating Prostatic Hyperplasia: A Randomized Controlled Trial

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**Abstract:** Objective: To assess the clinical effect of electroacupuncture for treating prostatic hyperplasia. Methods: Sixty patients with prostatic hyperplasia were randomized into a treatment group and a control group, 30 in each. The treatment group was treated by electroacupuncture, and the control was prescribed with orally taken *Jing Zhu Qian Lie Long Bi Tong* Capsule. After treatment, the two groups were compared with each other in terms of treatment result, international prostate symptom score (I-PSS), quality of life index (QLI), maximum flow rate (Q<sub>max</sub>), and residual urine volume (RUV). Results: The total effective rate was 96.7% in the treatment group versus 70.0% in the control, and the difference was significant ( $P < 0.05$ ). Both groups had marked improvement in I-PSS, QLI score, Q<sub>max</sub> and RUV ( $P < 0.05$ ). Conclusion: By improving I-PSS, QLI, Q<sub>max</sub> and RUV, electroacupuncture is an effective way for treating prostatic hyperplasia.

**Key Words:** Electroacupuncture; Acupuncture Therapy; Prostatic Hyperplasia; Difficulty in Urination; Randomize Controlled Trial

Benign prostatic hyperplasia is a common disease in elderly males. According to the survey conducted by Gu[1] in 20 hospitals, 13.6% of the inpatients in urinary department were prostatic hyperplasia. We treated this disease by electroacupuncture and compared the effect with medication. The report is now given as follows

### Materials and methods.

#### 1 Clinical Data

##### 1.1 Diagnosis criteria[2]

① Urinary symptoms such as urinary hesitancy, obstructed urine, urination frequency worse at night, urine leakage afterwards, distending pain in the lower belly, but absence of irritating pain in the urinary tract; ② digital rectal examination finds prostate enlargement, and dullness to percussion in lower belly; ③ diagnosis confirmed by ultrasonic scanning; ④ urinary infection, urinary tract stone and prostate cancer should be excluded.

##### 1.2 Inclusion criteria

① Conformed to the diagnosis criteria; ② willing to participate in this randomized controlled clinical study and sign the informed consent form.

##### 1.3 Exclusion criteria

① Allergic to the medication used in the present study; ② accompanied by mental health problem, liver or kidney dysfunction, hepatopathy, or infectious diseases; ③ patients who adopted other treatment methods in recent 2 weeks.

##### 1.4 Termination criteria

① Patients who cannot finish the study; ②patients who fail to comply with the treatment protocol; ③ severe adverse effect event occurs; ④when a complication occurs or the disease condition is aggravated. For terminated cases, doctor has to record the termination time and analyze the reason.

### 1.5 General data

Sixty eligible subjects were recruited from the Geriatric Department of the Second Hospital Affiliated to Heilongjiang University of Traditional Chinese Medicine, and then randomized into a treatment group and a control group, 30 cases in each. The subjects were aged 50-75 years. There were no significant differences in comparison of age, disease duration, international prostate symptom score (I-PSS), quality of life index (QLI), maximum flow rate (Qmax), and residual urine volume (RUV) (table 1-3).

Table 1. Comparison of age distribution (Cases)

Age (year)	Treatment group (n=30)	Control group (n=30)
50-55	4	3
56-60	5	6
61-65	7	6
66-70	9	11
71-75	5	4

Table 2. Comparison of disease duration (Cases)

Disease duration (year)	Treatment group (n=30)	Control group (n=30)
0-1	4	4
1-5	8	9
5-10	12	10
10-6	6	7

Table 3. Comparison of I-PSS, QLI, Qmax, RUV before

Indexes	Treatment group (n=30)	Control group (n=30)	t
I-PSS	17.30±4.74	18.70±4.81	1.14
QLI	4.33±1.24	4.97±1.22	1.20
Qmax	10.47±2.49	9.30±2.70	1.74
RUV	75.33±16.13	78.50±16.14	0.76

## 2 Treatment Methods

2.1 Treatment group Acupoints: Foot Motor-Sensory Region (scalp acupuncture area); Qugu (CV 2), Zhongji (CV 3), Guanyuan (CV 4), Henggu (KI 11), Sanyinjiao (SP 6).

Operation: After sterilization, the selected acupoints were inserted with the needles of 0.30 mm in diameter and 40 mm in length. The Foot Motor-Sensory Region was punctured subcutaneously by depth of 30mm, until it reached the subgaleal and producing a gasping feeling. Before Qugu (CV 2), Zhongji (CV 3), Guanyuan (CV 4), Henggu (KI 11) were punctured, the patient was asked to empty the urine bladder first and then to lie on his back. These points were punctured perpendicularly by depth of 35-38 mm, with needling sensation radiated to the urinary tract, perineum and the upper inner thigh. Sanyinjiao (SP 6) was also needled perpendicularly by depth of 35 mm, better to produce a distending needling sensation. After qi arrival, the bilateral Foot Motor-Sensory Region, Qugu (CV 2) and Guanyuan (CV 4) were connected to the acupuncture apparatus (Yingdi KWD-808) respectively, with sparse-dense wave and tolerable strength of current

for 30 min in total. The treatment was given once every day, 10 times as a treatment course, with an interval of 2-3 d between each two courses, and the treatment outcomes were evaluated after 3 treatment courses.

## 2.2 Control group

The control group was given Jing Zhu Qian Lie Long Bi Tong Capsules for orally taken (Z20025304), 4 capsules each time (0.5 g/capsule), and 3 times a day. The treatment course was the same as the treatment group.

## 3 Therapeutic Effects

3.1 Criteria of therapeutic effect [3, 4] Marked effect: I-PSS  $\leq 7$ ; QLI  $\leq 1$ ; Qmax  $\geq 18$  mL/s. The marked effect can be confirmed by achieving two of the above levels. Effect: I-PSS  $\leq 13$ ; QLI score drops from 4-6 to 2-3 after treatment; RUV is reduced by over 50%; Qmax  $\geq 12$  mL/s. Effect can be determined by achieving one of the above levels.

Invalid: None of the level mentioned for determining an effect is achieved.

## 3.2 Statistical analysis

The values were expressed by ( $\bar{x} \pm s$ ); the measurement data were analyzed by t-test, and the numerical data were by Chi-square test.

## 3.3 Treatment results

3.3.1 Comparison of treatment effect The comparison of total effective rate found that the treatment group had a significantly better result than the control ( $P < 0.01$ ) (table 4).

### 3.3.2 Comparison of I-PSS

It found that both groups had marked improvement in I-PSS after treatment ( $P < 0.05$ ), and the inter-group comparison also showed significant difference after treatment ( $P < 0.05$ ) (table 4)

Table 4. Comparison of clinical effect (Cases)

Groups	<i>n</i>	ME	Effect	Invalid	TER (%)
Treatment	30	11	18	1	96.71
Control	30	6	15	9	70.0

Note: ME=Marked effect; TER=Total effective rate; compared with the control group, 1)  $P < 0.01$

Table 5. Comparison of I-PSS ( $\bar{x} \pm s$ , Score)

Groups *n* Before treatment After treatment

Treatment 30 17.30 $\pm$ 4.74 7.73 $\pm$ 3.53 (1)2)

Control 30 18.70 $\pm$ 4.81 12.17 $\pm$ 4.02 (1)

Note: Intra-group comparison between before and after treatment,

1)  $P < 0.05$ ; compared with after treatment in the control group, 2)

$P < 0.05$

### 3.3.3 Comparison of QLI

It found that both groups had marked improvement in QLI after treatment ( $P < 0.05$ ), and the inter-group comparison also showed significant difference after treatment ( $P < 0.05$ )

### 3.3.4 Comparison of Qmax

It found that both groups had marked improvement in Qmax after treatment ( $P < 0.05$ ), and the inter-group comparison also showed significant difference after treatment ( $P < 0.05$ ).

### 3.3.5 Comparison of RUV

It found that both groups had marked improvement in RUV after treatment ( $P < 0.05$ ), and the inter-group comparison also showed significant difference after treatment ( $P < 0.05$ ) (table 5).

Table 5. Comparison of RUV ( $\bar{x} \pm s$ , mL)

Groups	<i>n</i>	Pre-treatment	Post-treatment
Treatment	30	75.33 $\pm$ 16.13	28.50 $\pm$ 10.35
Control	30	78.50 $\pm$ 16.14	43.37 $\pm$ 19.81

### 3.3.6 Adverse effects

None of the participants ever disobeyed the treatment schedule, or had a severe adverse effect

during the whole procedure, and there was no terminated case.

### **Result and Discussion**

Prostatic hyperplasia belongs to the scopes of "Long Bi (uroschesis)" or "Lin Zheng(stranguria)" according to traditional Chinese medicine (TCM) theory. The pathogenesis can be summarized to be deficiency of lung, spleen and kidney as the root cause, with retained stasis and dampness as the symptom [5,6]. The Foot Motor-Sensory Region is located at the paracentral lobule, and is also the projection area of the advanced micturition center. When this region is punctured, it can activate the function of micturition center, restore the modulation of subcortical micturition center, inhibit the hyperexcitation of detrusor muscle, and reduce the frequency and urgency of urination [7]. Qugu (CV 2), Zhongji (CV 3), and Guanyuan (CV 4) are all from the Conception Vessel; Qugu (CV 2) is the crossing point of the Conception Vessel and the Liver Meridian of Foot Jueyin; Zhongji (CV 3) and Guanyuan (CV 4) both are crossing points of the Conception Vessel and the Three Foot Yin Meridians. Therefore, the combined use of the three acupoints can regulate qi and blood in the Conception Vessel. Henggu (KI 11) is the crossing point of the Thoroughfare Vessel and the Kidney Meridian of Foot Shaoyin. Acupuncture at this point works to adjust the function of the bladder, for achieving proper bladder tonicity [8]. Qugu (CV 2), Zhongji (CV 3), Guanyuan (CV 4), and Henggu (KI 11) are all located on bladder and prostate. Hence, needling these points can improve the local symptoms [9,10]. Meanwhile, in association with needling the Foot Motor-Sensory Region and Sanyinjiao (SP 6), as well as electric stimulation, a satisfactory result was finally obtained.

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