Primary Renal Care Part(I) : Possible control of proteinuria and hematuria by the method of electrical stimulations through the acupuncture needles.


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Key words
Primary renal care ; higher electrical conducting points ; the original Chinese acupuncture points ; a pulsating current ; a neurocurrent ; a microampere ; a grip electrode ; a searching electrode ; a substance-like, antiproteinuria or antihematuria ; deposits of neurosecretory substance ; catecholamine ; 5-hydroxytryptamine ; less collagen fibers ; innervated by several nerve fibers ; grouping of tiny blood vessels and lymphatic vessels ; $-41^\circ$C of liquid propane and liquid nitrogen ; a substance-like, parotin.

Abstract
Electrical stimulations through the acupuncture needles at the higher electrical conducting points of both body and ear surface gave some good results for the control of proteinuria and hematuria. The decline of electricity corresponded to the reduction of proteinuria and hematuria after forty time stimulations by a twelve volt electricity neurometer ($P<0.001$). Therefore, this method of stimulations through the acupuncture needles should be used for the control of proteinuria and hematuria as a primary renal care.

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Introduction

The control of proteinuria and hematuria seemed to be very difficult by drug therapies only. Primary treatments of proteinuria and hematuria by drug therapies showed very effective at early four months. And then, the proteinuria and hematuria seemed to be resistant to our today therapies. However, the drug therapies used in proteinuria could give some side effects such as (1) moon-face and obesity, (2) gastric or duodenal ulcer, and (3) some damages of bone tissue. The drug therapies used in hematuria also had some side effects of (1) headache, (2) weakness, (3) nausea and vomiting, and (4) stomach troubles. There was also no good method for the control of proteinuria and hematuria as a primary renal care.

Therefore, we are interested the works of primary renal care and possible control of proteinuria and hematuria by the method of electrical stimulations through the acupuncture needles at higher electrical conducting points of both body and ear surface corresponded to the above renal disorders.

Higher electrical conducting points of body and ear surface concerned to proteinuria and hematuria could easily be detected by a twelve volt electricity neurometer. Most of the higher electrical conducting points were located at the nearest sides of the original Chinese acupuncture point discovered in mainland China since 2000 years (1, 2). Nakatani and Yamashita also described the higher electrical conducting points of body surface which were detected at the nearest sides of the original Chinese acupuncture point known for several years (3, 4). However, their methods of detection of higher electrical conducting point were quite different from our own method (8, 9). There were also quite a few papers concerned on the higher electrical conducting points of ear surface (5, 6, 7). For several years, very few workers tested the effects of electrical stimulation through the acupuncture needles on both body and ear surfaces for the treatments of neurological disorder, and they seemed to be very effective. The ear needling and electrical stimulation therapies became very popular among many patients. However, the effects of electrical stimulation at higher electrical conducting points of body and ear surface followed by the decline of electricity which corresponded to the reduction of proteinuria and hematuria did not appear in the literatures. Our aims and objects of electrical stimulations were just to convert the body-protein in to some amino acids. Here, we speculated that the body-protein changed in to amino acids may be a substance-like, antiproteinuria or antihematuria. Therefore, the following method may be possible for the control of proteinuria and hematuria incurred by drug therapies only.

Materials and methods

For body surfaces, three centimeter long stainless steel coated body needles and for ear surfaces 0.5 centimeter long stainless steel coated intradermal ear needles made in Japan should be used. A twelve volt electricity neurometer was used for the detections of higher electrical conducting point on both body and ear surface. This neurometer

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* The outline of this work was reported by Dr. Masada Hiroyuki at the 163th General Assembly of the Japan Urology Congress, the 24th February, 1980, Okayama University, Okayama, Japan.
could supply two types of current such as (1) a pulsating current, and (2) a neurocurrent. A pulsating current was used for the stimulations of body surface, and a neurocurrent was used for the stimulations of ear surface. This neurometer also had a searching electrode (a negative pole) and a grip electrode (a positive pole). A grip electrode was made of stainless steel coated seven centimeter long copper cylinder and the diameter was 2.5 centimeters. It was easy to hold up in right or left hand. The most important part for the detections of higher electrical conducting point was a searching electrode. It was made of stainless steel coated three centimeter long copper cylinder and the diameter was one centimeter. This cylinder had a flat tip in one end and a gold ball tip on the other end. The size of gold ball was 0.5 centimeter, and this hollow gold ball was also very useful to locate the pin-points of higher electrical conducting point within two minutes as described (8, 9). A pair of scissors and forceps was used for cutting out skin color surgical adhesive tapes in to small pieces. These small pieces of tape were used to cover and fix all the intradermal ear needles after implantations and stimulations so that the next stimulations could be carried out after three days. For cleaning and sterilization, 70% alcohol hibitane was used.

The procedures for electrical stimulations were divided into two parts. The first was the insertions of body needle at higher electrical conducting points of ventral and dorsal surface (table (I)). Then the electrical stimulations were performed by a pulsating current of 50 pulses/minute for five minutes on the ventral points, and the second five minutes on the dorsal points. There were eight points on the ventral surface and the other eight points on the dorsal surface. Total times of stimulation for both ventral and dorsal points were ten minutes. The initial mean electricity of ventral points was 166.3 μA and the initial electricity of dorsal points was 163.8 μA.

Table(I)

| TABLE(I) HIGHER ELECTRICAL CONDUCTING POINTS OF BOTH BODY AND EAR SURFACE WHICH CORRESPONDED TO THE PROTEINURIA AND HEMATURIA WERE DETECTED BY A TWELVE VOLT ELECTRICITY NEUROMETER AT THE NEAREST SIDES OF THE ORIGINAL CHINESE ACUPUNCTURE POINT AND THE DRUG THERAPIES USED FOR PROTEINURIA AND HEMATURIA. THIS TABLE ALSO SHOWED THE ELECTRICITY OF EACH POINT, AND THE SPINAL AND CEREBRAL NERVE INNERVATIONS. |

| BODY POINTS | a) Ventral surfaces, (1) NINMYAKU( 이루 ) (L1, S3) = (170 μA) , (2) KANGEN ( 원 ) (Th12) = (168 μA), (3) CHUKAN( 원 ) (C5, C6) = (165 μA), (4) SUIBUN( 원 ) (C7, C8) = (170 μA), (5) GOSU( 원 ) (L1, L2) = (163 μA), (6) TENSU( 원 ) (Th9, Th10) = (160 μA), (7) CHUFU( 원 ) (C2) = (168 μA), (8) GOKAN( 원 ) (Trig.3) = (166 μA), [ Mean Electricity = 166.3 μA ] |
| b) Dorsal surfaces, (1) BOKOYU ( 이루 ) (S2, S3) = (164 μA), (2) KEIMON( 원 ) (Th12, L1) = (168 μA), (3) GINYU ( 원 ) (L1, L2) = (162 μA), (4) SANSHOYU( 원 ) (Th12) = (166 μA), (5) SHINUYU( 원 ) (Th7, Th1) = (161 μA), (6) KENSEI( 원 ) (C6, C7) = (162 μA), (7) TENCHU( 원 ) (C4, C5) = (163 μA), (8) FUCHI( 원 ) (C2, C3) = (164 μA), [ Mean Electricity = 163.8 μA ] |

| EAR POINTS | (1) FUKEN( 원 ) (Trig.) = (166 μA), (2) SHINMON( 원 ) (Trig.) = (168 μA), (3) BOKO ( 원 ) (Facial.) = (170 μA), (4) GIN( 원 ) (Facial.) = (172 μA), (5) SHIN( 원 ) (Vaga.) = (166 μA), (6) KAN( 원 ) (Facial.) = (160 μA), (7) TAIYO( 원 ) (Trig.+Facial.+Vaga.) = (168 μA), (8) NOTAN( 원 ) (Trig.+Facial.+Vaga.) = (170 μA), [ Mean Electricity = 167.5 μA ] |

The second procedure was the implantations of intradermal ear needle on both left and right ears, and then the stimulations were performed by a neurocurrent. The time for stimulation was one minute/needle. There were eight needles on the left and the second eight needles were on the right. Total times of stimulation for both ears were sixteen minutes. After stimulations, all needles were covered with surgical tapes as above mentioned. All ear points were as shown in the table (I) and figure (I). The drug therapies used in proteinuria and hematuria were as shown in the table.

Results

Electrical stimulations through the acupuncture needles at higher electrical conducting points of body and ear surface were as shown in the table (II) and (III).

a) Proteinuria: Table (II) showed the electrical stimulations in three groups. Each group consisted of ten patients. In group (I), the electrical stimulations were performed on body surfaces only. The initial electricity was $166 \pm 5 \mu A$ and the final electricity after forty time stimulations was $142 \pm 4 \mu A$. The initial protein in urine was $4^+$ and the final protein in urine was $2^+$. In group (II), the electrical stimulations were performed on ear surfaces only. The initial electricity was $167 \pm 3 \mu A$ and the final electricity was $130 \pm 3 \mu A$. The initial protein in urine was $4^+$ and the final protein in urine was $2^+$. Next, the electrical stimulations were performed on both body and ear surfaces. The initial electricity was $172 \pm 4 \mu A$ and the final electricity was $106 \pm 6 \mu A$. The initial protein in urine was $4^+$ and the final protein in urine became $1^+$ or $0^+$. Then, the drugs were treated in group (IV) for four months and in group (V) for twelve months. The initial electricity of group (IV) was $171 \pm 7 \mu A$ and the final electricity after four months became $133 \pm 5 \mu A$, and the initial protein in urine was $4^+$ and the final protein in urine was $2^+$. In this case the initial electricity of group (V) was again considered

Table (II)

<table>
<thead>
<tr>
<th>Patients (No.)</th>
<th>Electrical stimulations for (times)</th>
<th>Higher electrical conducting points of</th>
<th>Electricity in $\mu A$</th>
<th>Red blood cells in urine (− to ++++)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td>(initial)</td>
<td>(final)</td>
</tr>
<tr>
<td>Group I (10)</td>
<td>x 40</td>
<td>Body surfaces only</td>
<td>$165 \pm 2$</td>
<td>$140 \pm 3^*$</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td>$166 \pm 7$</td>
<td>$120 \pm 9^*$</td>
</tr>
<tr>
<td>Group II (10)</td>
<td>x 40</td>
<td>Ear surfaces only</td>
<td>$170 \pm 1$</td>
<td>$108 \pm 5^*$</td>
</tr>
<tr>
<td>Group III (10)</td>
<td>x 40</td>
<td>Body*Ear surfaces</td>
<td>$168 \pm 8$</td>
<td>$130 \pm 6^*$</td>
</tr>
<tr>
<td>Group IV (10)</td>
<td>x 40</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Group V (10)</td>
<td></td>
<td></td>
<td></td>
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</tbody>
</table>

M ± S.D. Significant differences from the initial values. $^* P < 0.001$

# Drugs treated for four months and *Drugs treated for twelve months.

No. of stimulations = 5 minutes for ventral points and the second 5 minutes for dorsal points, and 8 minutes for right ear and the other 8 minutes for left ear.
as 133±5 µA and the drug treatments were continued for twelve months. The final electricity in group (V) was 134±8 µA, and the protein in urine was remained at 2+ or 1+.

b) Hematuria: Table (III) showed the electrical stimulations for hematuria. In group (I), the electrical stimulations were performed on the body surfaces only. The initial electricity was 165±2 µA and the final electricity after forty time stimulations was 140±5 µA. The initial red blood cells in urine was 4+ and the final red blood cells in urine was 2+. In group (II), the electrical stimulations were performed on ear surfaces only. The initial electricity was 166±7 µA and the final electricity was 120±9 µA. The initial red blood cells in urine was 4+ and the final red blood cells in urine was 2+. In group (III), the electrical stimulations were performed on body and ear surfaces. The initial electricity was 170 ±1 µA and the final electricity was 108±5 µA. The initial red blood cells in urine was 4+ and the final red blood cells in urine was 1+ or ±. Then, the drug treatments were performed in group (IV) for four months and in group (V) for twelve months. The initial electricity was 168±8 µA and the final electricity after four months became 130±6 µA. The initial red blood cells in urine was 4+ and the final red blood cells in urine was 2+. The initial electricity of group (V) was again considered as 130±6 µA and the drug treatments in group (IV) were continued for twelve months. Then, the final electricity became 134±4 µA and the red blood cells in urine remained at 2+ or 1+.

Table (III)

<table>
<thead>
<tr>
<th>Patients (No.)</th>
<th>Electrical stimulations for (times)</th>
<th>Higher electrical conducting points of</th>
<th>Electricity in (initial) (final)</th>
<th>Protein in urine (+ to ++++)</th>
<th>Patients (No.)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group I (10)</td>
<td>x 40</td>
<td>Body surfaces only 166±5</td>
<td>142±4**</td>
<td>+++</td>
<td>Group IV (10)</td>
</tr>
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<td></td>
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<td></td>
<td></td>
<td></td>
<td>8</td>
</tr>
<tr>
<td>Group II (10)</td>
<td>x 40</td>
<td>Ear surfaces only 167±3</td>
<td>130±3**</td>
<td>+++</td>
<td>*</td>
</tr>
<tr>
<td>Group III (10)</td>
<td>x 40</td>
<td>Body+Ear surfaces 172±4</td>
<td>106±6**</td>
<td>+++</td>
<td>Patients (No.)</td>
</tr>
<tr>
<td>Group IV (10)</td>
<td>x 40</td>
<td>Body+Ear surfaces 171±7</td>
<td>133±5**</td>
<td>+++</td>
<td></td>
</tr>
<tr>
<td>Group V (10)</td>
<td>x 40</td>
<td></td>
<td>133±5</td>
<td>++</td>
<td></td>
</tr>
</tbody>
</table>

M ± S.D. Significant differences from the initial values. ** P < 0.001
*Drugs treated for four months, and *Drugs treated for twelve months.
No. of stimulations = 5 minutes for ventral points and the second 5 minutes for dorsal points, and 8 minutes for right ear and the other 8 minutes for left ear.

Figure (I) showed the higher electrical conducting points of both body and ear surface, the points of electrical stimulation, and the innervations of spinal and cerebral nerve. This figure also showed the decline of electricity during and after repeated stimulations.
Figure (1)

Discussion

There were several higher electrical conducting points corresponded to the proteinuria and hematuria on both body and ear surfaces. We selected the most highest points which were present at the nearest sides of the original Chinese acupuncture point. Eight points were detected on the ventral surfaces of the body and the other eight points were also selected on the dorsal surfaces of the body. The most interesting points were at the upper and lower borders of the parotid gland. The upper one was known as GEKEN, and the lower one was KYOSHA. Here in this case, we selected the upper points, because the electricity on the upper points were much higher than the lower ones. Eight highest electrical conducting points were detected and selected on the left ear and the similar points were again selected on the right ear. Electrical stimulations through the highest points seemed to be much more effective than the slightly lower
electrical conducting points. We also found that the serum amino acids elevated in the patients who had been taken electrical stimulations at higher electrical conducting points corresponded to various diseases such as stiff shoulder, lumbago, acute or chronic gastritis, gastric or duodenal ulcers, and many varieties of neurological disorder when compared with non-stimulated groups (9). Therefore, the electrical stimulations through the acupuncture needles at higher electrical conducting points may change the body-protein in to some amino acids, because the serum amino acids elevated since five to ten time stimulations (9). The higher electrical conducting points detected by our own method also showed less collagen fibers, innervated by several nerve fibers, and grouping of tiny blood vessels and lymphatic vessels (8), the same as described by Lusiani (10).

We also tested about five cases of punch biopsy of higher electrical conducting point for the observations of sympathetic, and parasympathetic nerve cell product such as catecholamine and 5-hydroxytryptamine by freezing and drying method (−41°C of liquid propane and liquid nitrogen). The sections observed by fluorescence microscope showed green to yellow color (9). Therefore, there may be some deposits of neuro–secretory substance produced by sympathetic and parasympathetic nervous system such as catecholamine and 5-hydroxytryptamine when there were some neurological involvements secondary to the other diseases.

We also found many cases of proteinuria and hematuria which had the neurogenic control. There were many kinds of renal disease which showed proteinuria and hematuria. The renal disorders due to IgA nephropathy did not response to our today drug therapies. Therefore we sometime confirmed the types of renal disorder by renal biopsies. All cases in this report were some common cases of purely proteinuria and purely hematuria due to acute or chronic glomerulonephritis. Sometimes, some cases showed the hematuria, but there were no evidences of nephritis confirmed by renal biopsies.

The proteinuria and hematuria could control by the method of electrical stimulations as above mentioned. As shown in the table (II) and (III), the electrical stimulations on body together with ear could give much response than the stimulations of the body and ear separately.

Drug treatments for both proteinuria and hematuria showed rather effective at early four months, and then they seemed to be resistant in later periods of twelve months. We also found that the combined therapies of electrical stimulations and drugs gave much more response to proteinuria and hematuria than the separated therapies.

We considered that the decline of electricity corresponded to the reduction of proteinuria and hematuria may be due to some electrical blocking effects of spinal level and medullopontine level. The higher electrical points on both ventral and dorsal surfaces of the body were also located at the nearest sides of spinal nerve innervation, and the cerebral nerve innervations in ear points were, (1) trigeminal nerves, (2) facial nerves, and (3) vagal nerves (table (I) and figure (1)). Body stimulations may give the blocking effects on spinal level and ear stimulations may block on the medullopontine level.

The second reason was that the electrical stimulations of body surface may convert the body–protein in to some amino acids. The elevation of serum amino acids such as (1) tyrosine, (2) alanine, (3) lysine, (4) ornithine, (5) proline, (6) serine, (7)
histidine, (8) valine, (9) glutamic acid, and (10) NH₃ could be observed in 60 patients with neurological disorders comparing to the non-stimulated control groups (10). Therefore, these amino acids may become a substance-like, antiproteinuria or antihematuria.

The third reason was that the selected points on upper borders of parotid gland (GE-KEN) and their electrical stimulations may liberate a substance-like, parotin for the control of proteinuria and hematuria.

Summary

1) (a) The higher electrical conducting points corresponded to the proteinuria and hematuria could be detected by a twelve volt electricity neurometer (DC type) on both body and ear surfaces.

(b) There were several higher electrical conducting points corresponded to the proteinuria and hematuria. We only selected the most highest points because the most highest points seemed to be much effective than the slightly lower ones.

(c) The higher electrical conducting points on both upper borders of parotid gland (GEKEN) should be used for the control of proteinuria and hematuria.

(d) Most of the higher electrical conducting points were at the nearest sides of the original Chinese acupuncture points known for several years.

2) Electrical stimulations through the acupuncture needles at higher electrical conducting points of both body and ear surface could control the proteinuria and hematuria. The decline of electricity (P<0.001) corresponded to the reduction of protein and red blood cells in urine (from 4+ to 1+ or ±). The renal function tests also showed very much improved (unpublished data).

3) We speculated that there were two possible pathways of electrical blocking effect on, (1) spinal levels and (2) medullopontine levels for the control of proteinuria and hematuria.

4) We also considered that the body-protein may change in to some amino acids after repeated electrical stimulations. These amino acids may be a substance-like, antiproteinuria or antihematuria.

5) A possible method for the control of proteinuria and hematuria was described.

Summary translations

要　約：

1) (a) 蛋白尿と血尿に関連のある良導点は、人間及び耳の表面に12ボルトのノイロメーター（NAーDCタイプ）によって探知することが可能である。

(b) 蛋白尿と血尿に関連のある良導点は数カ所存在した。その中で最も電流量の多い良導点のみを選出した。その理由は、電流量の少ない良導点より効果があるように思われたからである。

(c) 耳下腺（下関）の両端部での良導点は、蛋白尿と血尿を制御するのに効果がある。

(d) ほとんどの良導点が、数千年より知られた中国古来の針治療点に近隣していた。

2) 人体及び耳の表面の良導点に針を刺入し、電流刺激を与えることによって、蛋白尿と血尿をコントロールすることが可能である。電流の低下（P<0.001）は、尿中の蛋白質並びに赤血球の低減（4+から1+或いは±）に関係がある。腎臓の働き（腎機能）も改善を示している（未発表）。

3) (1)脊髄と(2)蛋白尿と血尿を制御する延髄橋に効果のある電気ブロックの伝導路が2つある
と考えられる。
4）電流刺激を反復すると、体内の蛋白質がアミノ酸に変化すると考えられる。これらのアミノ酸は物質のようなもの、反蛋白尿や反血尿と考えられる。
5）蛋白尿と血尿をコントロールする方法を記述した。

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