Renin–Aldosterone Paradox and Perturbed Blood Volume Regulation Underlying Postural Tachycardia Syndrome

Satish R. Raj, MD; Italo Biaggioni, MD; Paula C. Yamhure, RN; Bonnie K. Black, RN, NP; Sachin Y. Paranjape, BS; Daniel W. Byrne, MS; David Robertson, MD

Background—Patients with postural tachycardia syndrome (POTS) experience considerable disability, but in most, the pathophysiology remains obscure. Plasma volume disturbances have been implicated in some patients. We prospectively tested the hypothesis that patients with POTS are hypovolemic compared with healthy controls and explored the role of plasma renin activity and aldosterone in the regulation of plasma volume.

Methods and Results—Patients with POTS (n = 15) and healthy controls (n = 14) underwent investigation. Heart rate (HR), blood pressure (BP), plasma renin activity, and aldosterone were measured with patients both supine and upright. Blood volumes were measured with $^{131}$I-labeled albumin and hematocrit. Patients with POTS had a higher orthostatic increase in HR than controls (51 ± 18 versus 16 ± 10 bpm, P < 0.001). Patients with POTS had a greater deficit in plasma volume (334 ± 187 versus 10 ± 250 mL, P < 0.001), red blood cell volume (356 ± 128 versus 218 ± 140 mL, P = 0.010), and total blood volume (689 ± 270 versus 228 ± 353 mL, P < 0.001) than controls. Despite the lower plasma volume in patients with POTS, there was not a compensatory increase in plasma renin activity (0.79 ± 0.58 versus 0.79 ± 0.74 ng · mL$^{-1}$ · h$^{-1}$, P = 0.996). There was a paradoxically low level of aldosterone in the patients with POTS (190 ± 140 pmol/L versus 380 ± 230 pmol/L; P = 0.017).

Conclusions—Patients with POTS have paradoxically unchanged plasma renin activity and low aldosterone given their marked reduction in plasma volume. These patients also have a significant red blood cell volume deficit, which is regulated by the renal hormone erythropoietin. These abnormalities suggest that the kidney may play a key role in the pathophysiology of POTS. (Circulation. 2005;111:1574-1582.)

Key Words: tachycardia ■ renin ■ nervous system, autonomic ■ blood volume ■ aldosterone
We also explored the regulation of plasma volume by renin and aldosterone.

**Methods**

**Subjects**

Patients referred to the Vanderbilt University Autonomic Dysfunction Center with POTS between October 2002 and November 2003 were candidates for inclusion in this study. Patients met the current criteria for POTS. Briefly, patients developed symptoms of orthostatic intolerance accompanied by a heart rate rise ≥30 bpm (or a rate that exceeds 120 bpm) that occurred within the first 10 minutes of standing or head-up tilt, without any evidence of orthostatic hypotension (a fall in blood pressure of ≥20/10 mm Hg). Patients had at least a 6-month history of symptoms, in the absence of another chronic debilitating disorder or prolonged bed rest, and were at least 18 years of age. Healthy control subjects (who did not meet criteria for POTS, and were at least 18 years of age) underwent all of the same protocol elements. Patients and controls were free of medications that could impair autonomic tone and were not taking fludrocortisone for at least 5 days before testing. The Vanderbilt University Investigational Review Board approved this study, and written informed consent was obtained from each subject before the study began.

**Protocol**

Study investigations were performed at the Elliot V. Newman Clinical Research Center at Vanderbilt University. For 3 days before testing, subjects consumed a diet that contained 150 mEq of sodium per day and 70 mEq of potassium per day. The diet was free of caffeine-containing beverages.

**Supine and Upright Posture Study**

Heart rate, blood pressure, aldosterone, plasma renin activity, and plasma norepinephrine and epinephrine were assessed after overnight rest with subjects in the supine position and again after subjects had been standing for up to 30 minutes (as tolerated). The standing test was performed to assess the hemodynamic and biochemical responses to increased central hypovolemia (accentuated by the gravitational stress). For catecholamine measurements, blood was collected in plastic syringes and immediately transferred to chilled vacuum tubes with EGTA and reduced glutathione (Amersham International PLC) and immediately put on ice. The plasma was separated by refrigerated centrifugation at −4°C and stored at −70°C until the assay. Concentrations of norepinephrine and epinephrine were measured by batch alumina extraction, followed by high-performance liquid chromatography for separation with electrochemical detection and quantification. Plasma renin enzymatic activity was assayed by conversion of angiotensinogen to angiotensin I by a radioimmunoassay technique (antibodies from IgG Corporation) and reported in nanograms of angiotensin I per milliliter per hour. Blood for aldosterone was collected in chilled vacuum tubes without preservative, and the serum was extracted and sent to the laboratory on ice. Serum aldosterone was measured by radioimmunoassay (DPC Coat-a-Count, Diagnostic Products Corp). The aldosterone-plasma renin activity ratio was calculated with the conventional units for aldosterone (ng/dL; 1 ng/dL = 27.7 pmol/L) and plasma renin activity (ng · mL⁻¹ · h⁻¹) and reported without units.

**Blood Volume Assessment**

Plasma volume was determined by the indicator dye-dilution technique. In the morning after an overnight fast, patients were placed in a supine position for a minimum of 60 minutes before collection of the baseline sample. A 20-gauge intravenous catheter was placed in an antecubital vein, and blood samples could be obtained without stasis. A baseline venous sample of 5 mL was collected before injection of the tracer. With a prefilled 1-mL syringe, up to 25 μCi of 131I-labeled human serum albumin (Volumex, Iso-Tex Diagnostics Inc) was injected into the antecubital vein and flushed with 30 mL of normal saline. Starting at 12 minutes after injection, 5 mL of venous blood was collected at 6-minute intervals until 30 minutes after injection (5 samples, including baseline sample). Hematocrit was measured in duplicate from each sample after 10-minute centrifugation at 11 500 rpm on an International Equipment Co microcapillary centrifuge and read on an International Equipment Co microcapillary tube reader. Plasma radioactivity was measured in duplicate and averaged (for each sample and a reference standard) with an automated counter (BVA-100 Blood Volume Analyzer, DAXOR Corporation). A least-squares regression of the volume of distribution at each time point was automatically performed to determine the volume of distribution at the time of injection. Plasma volume was determined as the volume of distribution of albumin.

Total blood volume was calculated from measured plasma volume and microcapillary venous antecubital hematocrit corrected for the plasma-packing ratio (0.99), the ratio of mean body hematocrit to peripheral (measured) hematocrit (0.91), and the effects of heparin within the sampling syringe (0.97).

Red blood cell volume was calculated as the difference between total blood volume and plasma volume. This DAXOR method of red blood cell volume assessment was recently found to correlate well with the traditional 51Cr red blood cell-labeling method (Pearson correlation R = 0.96), with a mean difference between the techniques of 0.9% (personal communication with Dr Howard Dworkin, William Beaumont Hospital, Royal Oak, Mich).

Ideal plasma and total blood volume was determined on the basis of the height, weight, and gender of the individual subject. Individual “deficits” in plasma volume, red blood cell volume, and total blood volume were calculated as the ideal minus measured volume (in milliliters), or this difference divided by the ideal volume (percentage).

**Plasma Volume Shift With Upright Tilt**

All studies occurred between 10 AM and noon in a quiet, dimly lit room at a comfortable ambient temperature (21°C to 24°C). An antecubital venous catheter was inserted (if not already in situ and functioning) for blood sampling at least 15 minutes before the beginning of the test, with the patient supine. Subjects were tilted head-up to 60° for 30 minutes or until the subject experienced presyncope that required test termination.

Blood was drawn at baseline and then at 5, 10, 15, 20, and 30 minutes of tilt for measurement of hematocrit in quadruplicate (see Blood Volume Assessment section for details of hematocrit assessment). Relative changes in hematocrit from baseline were used to calculate the change in plasma volume with upright tilt. The percentage change in plasma volume (ΔPV%) = 100 × (HctBaseline − HctFinal)/HctBaseline, with the absolute change in plasma volume (ΔPV) = ΔPV% × measured plasma volume (where HctBaseline is hematocrit before tilt, and HctFinal is hematocrit at a given time after tilt). ΔPV_final and ΔPV%_final were defined as the ΔPV at the time of the last measured hematocrit before tilt termination. ΔPV_final was used in place of the individual late time points to minimize the confounding effect of late data dropout due to premature tilt termination. A negative value reflects a shift in plasma volume out of the vascular space.

**Gender Analysis**

POTS is a disorder that affects women more often than men. In addition to an overall analysis that included all subjects, a separate analysis was performed that included only female subjects. This was to ensure that the results were not skewed by the small number of men.

**Statistical Analysis and Sample Size Calculations**

Our primary end point was the plasma volume deficit (measured plasma volume minus ideal plasma volume). The null hypothesis was that the plasma volume deficit would not be statistically different between patients with POTS and control subjects. We calculated the size of our required sample after determining that a 7.5% deficit in plasma volume in patients with POTS would be
clinically significant. We did not expect the control subjects to have a plasma volume deficit. Assuming a pooled SD of 5% (giving an effect size of 1.5), a sample size of 13 subjects in each group would give 95% power to detect a statistically significant difference with a Student t test with a 2-sided significance level of 0.05. Differences between groups were analyzed with the Student t test. The Mann-Whitney U test was also used to confirm the results obtained from the Student t test, and the significance of the reported parameters was not different between the 2 tests. Categorical variables were analyzed with the Fisher exact test. Values are reported as means and SDs unless otherwise noted. Probability values of \( P < 0.05 \) were considered statistically significant, and all tests were 2 sided. Statistical analyses were performed with SPSS for Windows (version 12.0, SPSS). Sample size calculations were performed with nQuery Advisor (version 5.0). Prism for Windows 4 (version 4.02, GraphPad Software Inc.) was used for graphical presentation.

Results

Baseline Information

We enrolled 15 patients with POTS and 14 controls. The baseline characteristics are enumerated in Table 1. There was no significant difference in baseline characteristics between patients with POTS and control subjects for the overall group or when just the female subjects were considered.

Supine and Upright Posture Study

As seen in Figure 1A, patients with POTS had a higher heart rate than control subjects both when supine (77 ± 12 versus 64 ± 12 bpm, \( P = 0.008 \)) and when standing upright for up to 30 minutes (128 ± 18 versus 80 ± 11 bpm, \( P < 0.001 \)). As would be expected on the basis of the diagnostic criteria for POTS, patients with POTS had a significantly greater increase in heart rate (51 ± 18 bpm) on assuming the upright position than did the control subjects (16 ± 10 bpm, \( P < 0.001 \)). The supine systolic blood pressure was similar between the 2 groups (POTS versus control, 111 ± 14 versus 114 ± 13 mm Hg; \( P = 0.525 \); Figure 1B). Both groups experienced a small increase in systolic blood pressure that was not statistically significant on standing, with no difference between groups (POTS versus control, 123 ± 20 versus 128 ± 18 mm Hg; \( P = 0.195 \)).

TABLE 1. Baseline Information

<table>
<thead>
<tr>
<th>Overall Group</th>
<th>Female Subjects Only</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>POTS (n=15)</td>
</tr>
<tr>
<td>Age, y</td>
<td>36±11</td>
</tr>
<tr>
<td>Height, cm</td>
<td>166±8</td>
</tr>
<tr>
<td>Mass, kg</td>
<td>66.3±19.3</td>
</tr>
<tr>
<td>Body mass index, kg/m²</td>
<td>24.2±6.8</td>
</tr>
<tr>
<td>Na⁺, mEq/L</td>
<td>138±2</td>
</tr>
<tr>
<td>K⁺, mEq/L</td>
<td>4.1±0.3</td>
</tr>
<tr>
<td>Calculated osmolality, mEq/L</td>
<td>286±4</td>
</tr>
<tr>
<td>Hematocrit, %</td>
<td>37.8±2.4</td>
</tr>
</tbody>
</table>

All values are presented as mean±SD. \( P \) value for gender distribution from a Fisher exact test, other \( P \) values from Student t test.

Figure 1. Hemodynamic parameters and catecholamines: supine and upright. Supine and upright values for heart rate (A), systolic blood pressure (BP; B), venous plasma norepinephrine (C), and venous plasma epinephrine (D) for patients with POTS and control subjects. Error bars represent SE. Probability values are from between-group comparison with Student t test.
had a significantly higher heart rate (P=0.001), a significantly higher plasma norepinephrine level (P=0.025), and a significantly lower aldosterone level (P=0.038) than control subjects. There were no differences between the groups while standing for blood pressure, plasma epinephrine, or plasma renin activity. In the setting of the smaller sample size in this subgroup analysis, none of the supine hemodynamic and biochemical parameters were significantly different between groups.

### Blood Volumes

As shown in Table 2, plasma volume was significantly lower in patients with POTS (2348±438 mL) than in control subjects (2823±480 mL, P=0.010), whereas ideal plasma
by guest on April 30, 2017 http://circ.ahajournals.org/ Downloaded from

with POTS had a plasma volume deficit of 334 \pm 187 mL, which represented 12.8 \pm 7.6\% of their ideal plasma volume, whereas control subjects had no deficit (10 \pm 250 mL [0.8 \pm 8.8\%]). Figure 3A illustrates both this highly significant difference and the variability in plasma volume deficits within the 2 groups. The plasma volume of 1 patient with POTS actually exceeded expectations. Conversely, 3 patients with POTS had a plasma volume deficit of \geq 20\%, with a plasma volume deficit as high as 27\% in 1 patient with POTS.

The measured red blood cell volume was also lower by a mean of \geq 300 mL in patients with POTS than in control subjects (Table 2). Control subjects experienced a modest deficit in red blood cell volume. In contrast, patients with POTS had a mean deficit in red blood cell volume of \geq 350 mL, which represented a 22.7\% deficit from the expected red blood cell volume (Figure 3B). The difference between the 2 groups was highly significant (P=0.003). Supine hematocrit values were not different between patients with POTS and control subjects (37.8 \pm 2.4\% versus 38.0 \pm 3.0\%, P=0.812), which reflects the parallel decrease in both plasma volume and red blood cell volume in patients with POTS.

The total blood volume followed the same pattern as the plasma volume and red blood cell volume components. The measured total blood volume was significantly lower in patients with POTS than in control subjects (P=0.010; Table 2). Even after we corrected for individual differences in ideal total blood volume, patients with POTS still had a significantly larger relative deficit in total blood volume (16.5 \pm 6.8\% versus 5.6 \pm 7.8\%, P<0.001; Figure 3C). This works out to a mean absolute total blood volume deficit of 460 mL compared with the control subjects.

Blood volumes for the female subjects are shown in Figures 3D through 3F and in Table 2. Concordant with the overall group, each of the 3 measured blood volume deficits was greater for patients with POTS than for control groups.

**Plasma Volume Shifts With Upright Tilt**

Plasma volume shifts with upright tilt were calculated both early during the tilt (5 minutes [\Delta PV_%5 minutes]) and near the end of tilt (\Delta PV_%End) as a percentage of the baseline plasma volume. Neither the mean \Delta PV_%5 minutes (–11.7 \pm 3.2\% versus –10.4 \pm 3.5\%, P=0.298) nor the mean \Delta PV_%End (–16.6 \pm 4.7\% versus –15.3 \pm 4.6\%, P=0.463) was different in patients with POTS compared with control subjects. As can be seen in Figure 4, there was significant heterogeneity in the maximal plasma volume shift with upright tilt in both groups.

**Discussion**

This study sought to assess both blood volume and the role of the renin–angiotensin–aldosterone system in the regulation of blood volume in POTS. The main findings from this prospective study were that compared with control subjects, patients with POTS (1) have a significant deficit of plasma volume, (2) have a significantly lower level of serum aldosterone, (3) have an inappropriately low level of plasma renin activity given the degree of hypovolemia that they exhibit, and (4) have a significant deficit of red blood cell volume in the setting of an elevated standing heart rate and plasma norepinephrine.

**Plasma Volume**

Patients with POTS had a lower basal plasma volume (Table 2) than the control subjects. Accurate plasma volume assessments can be affected by several environmental, dietary, and patient-related factors. These variables may in part explain

---

**TABLE 2. Blood Volumes**

<table>
<thead>
<tr>
<th></th>
<th>Overall Group</th>
<th>Female Subjects Only</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>POTS (n=15)</td>
<td>Control (n=14)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>P</td>
</tr>
<tr>
<td></td>
<td></td>
<td>P</td>
</tr>
<tr>
<td><strong>Plasma volume</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Measured, mL</td>
<td>2348 \pm 438</td>
<td>2823 \pm 480</td>
</tr>
<tr>
<td>Ideal, mL</td>
<td>2682 \pm 379</td>
<td>2833 \pm 287</td>
</tr>
<tr>
<td>Deficit, mL</td>
<td>334 \pm 187</td>
<td>10 \pm 250</td>
</tr>
<tr>
<td>Deficit, %</td>
<td>12.8 \pm 7.6</td>
<td>0.8 \pm 8.8</td>
</tr>
<tr>
<td><strong>Red cell volume</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Measured, mL</td>
<td>1201 \pm 221</td>
<td>1510 \pm 405</td>
</tr>
<tr>
<td>Ideal, mL</td>
<td>1557 \pm 268</td>
<td>1729 \pm 330</td>
</tr>
<tr>
<td>Deficit, mL</td>
<td>356 \pm 128</td>
<td>218 \pm 140</td>
</tr>
<tr>
<td>Deficit, %</td>
<td>22.7 \pm 7.2</td>
<td>13.5 \pm 8.2</td>
</tr>
<tr>
<td><strong>Total blood volume</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Measured, mL</td>
<td>3550 \pm 642</td>
<td>4334 \pm 868</td>
</tr>
<tr>
<td>Ideal, mL</td>
<td>4239 \pm 631</td>
<td>4562 \pm 608</td>
</tr>
<tr>
<td>Deficit, mL</td>
<td>689 \pm 270</td>
<td>228 \pm 353</td>
</tr>
<tr>
<td>Deficit, %</td>
<td>16.5 \pm 6.8</td>
<td>5.6 \pm 7.8</td>
</tr>
</tbody>
</table>

All values are presented as mean \pm SD. P values from Student t test. Blood volume deficits are calculated as the ideal volume minus the measured volume.
the mixed results obtained by other investigators who have tried to assess plasma volume in patients with POTS. Several measures were undertaken in the present study to ensure that the plasma volume was measured accurately. Study subjects were withdrawn from all medications that might alter the plasma volume (such as fludrocortisone) for at least 5 days before the study. They consumed a controlled sodium diet for at least 3 days before assessment, because sodium intake can alter activation of the renin–angiotensin–aldosterone axis and subsequently alter plasma volume. Finally, all plasma volume assessments were performed in the same temperature-controlled room on the research unit after the subjects had been supine for at least 1 hour. Other factors that can physiologically alter plasma volume include the subject’s size and gender. To correct for individual variations in subject size, we calculated the expected plasma volume for each subject based on his or her size and gender and subtracted the measured plasma volume to arrive at the individual’s actual plasma volume deficit, as shown in Figure 3A. Patients with POTS had a mean plasma volume deficit of almost 350 mL, whereas control subjects had no such deficit. The deficit of 13% in patients with POTS constitutes a moderate to severe hypovolemia. This reduction in effective circulating volume could trigger a cascade of perturbations associated with POTS. In the supine position, this hypovolemia may cause only modest or nonsignificant changes in heart rate and plasma norepinephrine. In the upright position, in the setting of gravitational blood pooling, the additional reduced volume could decrease the cardiac output and cause a reflex increase in sympathetic nerve activity. The result would be an increase in the upright plasma norepinephrine.

Figure 3. Blood volumes. Individual and mean values are presented for plasma volume (PV) deficit (A) as percentage (individual deficit = [ideal plasma volume – measured plasma volume]/ideal plasma volume) for control subjects and patients with POTS. Negative value for deficit implies excess in plasma volume. Probability values are from between-group comparison with Student t test. Similar figures are presented for red blood cell volume deficit (B) and total blood volume (TBV) deficit (C). D through F, Data when only female subjects were included.

Figure 4. Plasma volume shift with upright tilt. Individual and mean shifts in plasma volume (PV Shift), as percentage of baseline plasma volume, in response to upright tilt are presented for control subjects and patients with POTS. Negative value represents shift of plasma volume from intravascular to extravascular space. Mean differences were not statistically significant between groups. Max indicates maximum.
Renin–Aldosterone Paradox

The renin–angiotensin–aldosterone system plays an important role in the regulation of plasma volume. Hypovolemia, acting via reduced renal blood flow and possible cardiorenal mechanisms, would be expected to increase plasma renin activity, which should augment levels of angiotensin II and aldosterone. Angiotensin II promotes renal sodium retention both directly, through receptors in the renal proximal tubule, and indirectly by stimulating the secretion of the mineralocorticoid aldosterone. Through this augmented sodium retention, the renin–angiotensin–aldosterone axis should restore extracellular fluid volume (Figure 5A).

Plasma renin activity was similar in patients with POTS and control subjects, whereas supine and upright levels of aldosterone were significantly lower in patients with POTS (Figure 2). Given their degree of hypovolemia, however, one would expect both plasma renin activity and aldosterone levels to be significantly higher in the POTS group than in controls. Both the plasma renin activity and, to a greater extent, aldosterone levels were inappropriately low given the hypovolemic status of the patients with POTS. We have termed this dysregulation of plasma renin activity and aldosterone in POTS the “renin–aldosterone paradox” (Figure 5B).

Alfaldosterone secretion is controlled at many levels: it is stimulated by angiotensin II, potassium, and hypoxemia, and acutely by the adrenocorticotroic hormone; it is inhibited by dopamine and atrial natriuretic factor (ANF). Electrolyte abnormalities are not likely to explain the low aldosterone as the sodium and potassium levels were similar in the POTS group and controls (Table 1). Although we cannot exclude the possibility that there are abnormalities in ANF or increases in adrenal dopamine concentrations that could contribute to the low aldosterone state, the most likely explanation for the renin–aldosterone paradox is an inappropriately low level of angiotensin II.

The cause of the inappropriately low levels of plasma renin activity and aldosterone in POTS is not clear. Low-flow states across the juxtaglomerular apparatus (as is seen in renal artery stenosis) are known to increase plasma renin activity. It is possible that the opposite effect somehow occurs in POTS, presumably by impaired vascular function. Other possibilities include problems with the sensor mechanisms at the level of the macula densa, or in the transmission of this signal to the juxtaglomerular apparatus, or in the response of the juxtaglomerular apparatus. It is also possible that patients with POTS have a low blood vessel “capacitance” that in turn limits the blood volume. Such a phenomenon has been proposed to exist in patients with pheochromocytoma, who have low blood volume. Plasma volume and total blood volume increase in response to treatment with an α-adrenergic antagonist.

Red Blood Cell Volume

In addition to deficits in plasma volume patients with POTS also had a significant reduction in red blood cell volume (Figure 3B and Table 2). This finding would not have been apparent on cursory assessment. Hematocrit levels were not different between the 2 groups (Table 1). Because there was a parallel reduction in the 2 largest components of blood volume (plasma volume and red blood cell volume), the percentage of the blood column that was due to the red blood cells appeared to be normal. Thus, we required a formal radioisotope dilution assessment of blood volumes to document the red blood cell volume deficit.

This red blood cell volume deficit has been observed previously in patients with POTS. The pathogenesis of this deficit, however, is not known. The renal hormone erythropoietin is the primary agonist for red blood cell production in the bone marrow. It is possible that a deficit in erythropoietin production might play a pathophysiological role in POTS, although this is not yet clear.

There are several pieces of evidence that point to an important role for angiotensin II and the renin–angiotensin–aldosterone axis in the regulation of erythropoietin production. First, in healthy subjects, infusions of angiotensin II caused serum erythropoietin levels to increase significantly, but this stimulation of erythropoietin was blocked when the subjects were premedicated with losartan, an angiotensin receptor blocker. Second, plasma renin activity is higher among hemodialysis patients who do not require exogenous erythropoietin to maintain a hematocrit of 30% (which suggests adequate endogenous erythropoietin) than in those patients who require exogenous erythropoietin. Third, plasma ultrafiltration induced a doubling of plasma renin activity, which was accompanied by a 69% rise in serum erythropoietin over 4 hours. This increase in erythropoietin was abolished with the use of an ACE inhibitor. Finally, some patients develop a persistently elevated hematocrit after renal...
transplant.\textsuperscript{38} Inactivation of the renin–angiotensin–aldosterone system by an ACE inhibitor or an angiotensin receptor blocker can correct this polycthemia,\textsuperscript{39,40} and conversely, withdrawal of the ACE inhibitor has been associated with “rebound” polycthemia.\textsuperscript{38} Taken together, these pieces of evidence suggest that the paradoxically low renin activity seen in patients with POTS could be the cause of the low red blood cell volume through a direct hormonal effect.

Another possible explanation for the low red blood cell volume seen in POTS is that it is a direct result of the low plasma volume. The kidney may function as the key organ involved in the regulation of hematocrit, because it controls both plasma volume (through salt regulation) and red blood cell volume (through erythropoietin).\textsuperscript{41} To maintain an appropriate hematocrit (the hematocrit was similar between patients with POTS and controls), the red blood cell volume may be adjusted downward through a physiologically reduced level of erythropoietin to match the deficit in plasma volume.

Erythropoietin replacement, by itself, is not likely to restore normal physiological function in patients with POTS. Hoeldtke et al\textsuperscript{12} studied 8 patients with POTS and found 6 of those patients to have a low red blood cell volume. Treatment with open-label erythropoietin improved the red blood cell volume but did not increase the plasma volume. The orthostatic tachycardia was corrected in only 1 of their patients, although 3 patients subjectively reported feeling better.

**Study Limitations**

We have found that plasma renin activity and aldosterone are not appropriately regulated in patients with POTS. The low aldosterone-renin ratio seen in POTS suggests a mismatch between these 2 hormones. One limitation of the present study was that levels of angiotensin II, a biochemical link between renin and aldosterone, were not measured directly. Other potential regulators of aldosterone secretion (dopamine and ANF) and salt and water regulation (such as antidiuretic hormone, serum osmolality, and B-type natriuretic peptide) may also provide useful insights into the renin–aldosterone paradox seen in POTS. Future studies will include an assessment of these markers.

**Conclusions**

In summary, we have found that patients with POTS have a reduction in plasma volume. They have inappropriately low levels of renin and low levels of aldosterone, 2 hormones that promote sodium retention and increase plasma volume and are regulated by the kidneys. These patients also have a significantly low volume of red blood cells. Red blood cell production is primarily stimulated by erythropoietin, a hormone that is released by the kidney. Taken together, these findings suggest that abnormalities in the kidney might be critical in the pathophysiology of POTS.

**Acknowledgments**

This study was supported in part by National Institutes of Health grants 2P01 HL56993 and M01 RR00995 (General Clinical Research Center) from the National Institutes of Health. Dr Raj is a Vanderbilt Clinical Research Scholar, supported by a K12 grant from the National Institutes of Health. The DAXCOR Corporation (New York, NY) kindly donated the equipment and supplies needed for the blood volume assessment.

**References**


Renin-Aldosterone Paradox and Perturbed Blood Volume Regulation Underlying Postural Tachycardia Syndrome
Satish R. Raj, Italo Biaggioni, Paula C. Yamhure, Bonnie K. Black, Sachin Y. Paranjape, Daniel W. Byrne and David Robertson

_Circulation._ 2005;111:1574-1582; originally published online March 21, 2005;
doi: 10.1161/01.CIR.0000160356.97313.5D
_Circulation_ is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
Copyright © 2005 American Heart Association, Inc. All rights reserved.
Print ISSN: 0009-7322. Online ISSN: 1524-4539

The online version of this article, along with updated information and services, is located on the World Wide Web at:
http://circ.ahajournals.org/content/111/13/1574

Data Supplement (unedited) at:
http://circ.ahajournals.org/content/suppl/2005/03/21/01.CIR.0000160356.97313.5D.DC1

Permissions: Requests for permissions to reproduce figures, tables, or portions of articles originally published in _Circulation_ can be obtained via RightsLink, a service of the Copyright Clearance Center, not the Editorial Office. Once the online version of the published article for which permission is being requested is located, click Request Permissions in the middle column of the Web page under Services. Further information about this process is available in the Permissions and Rights Question and Answer document.

Reprints: Information about reprints can be found online at:
http://www.lww.com/reprints

Subscriptions: Information about subscribing to _Circulation_ is online at:
http://circ.ahajournals.org//subscriptions/