POSTURAL TACHYCARDIA SYNDROME

Supine Parasympathetic Withdrawal and Upright Sympathetic Activation Underly Abnormalities of the Baroreflex in Postural Tachycardia Syndrome

Effects of Pyridostigmine and Digoxin

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ABSTRACT: Upright postural tachycardia syndrome (POTS) resembles hemorrhage with reduced central blood volume, parasympathetic withdrawal, and sympathetic activation. Baroreflex dysfunction causes low heart rate variability, enhanced blood pressure variability, and decreased maximum baroreflex gain (G_{max}) putatively measured by spontaneous fluctuation of blood pressure and heart rate. We investigated whether/how cardiovagal baroreflex in POTS differ from control, supine, and upright by comparing indices of spontaneous baroreflex function to that measured using the reference standard modified Oxford method. This uses sodium nitroprusside and phenylephrine to generate the sigmoidal cardiovagal baroreflex curve. Baroreflex in POTS was evaluated supine and upright untreated and then treated to determine whether pyridostigmine or digoxin (a vagotonic agent) corrects baroreflex deficits. Supine, G_{max} was reduced by 25% in POTS compared with controls, and descriptors of this sigmoidal relationship showed a reduction, downward shift, and left shift of the response to the pharmacological decrease and increase in blood pressure. Digoxin normalized supine cardiovagal baroreflex while pyridostigmine resulted in partial normalization as G_{max}, and other descriptors of these relationships were similar to control. Upright, cardiovagal curves were distorted and displaced in untreated POTS, while digoxin and pyridostigmine left shifted the cardiovagal curves due to sympathetic activity. Cardiovagal baroreflex deficits in POTS relate to parasympathetic withdrawal supine, remediated completely by digoxin, and sympathetic activation upright through alteration of baroreflex responsivity. Since these baroreflex effects resemble those measured following microgravity/chronic bedrest, vagotonic/sympatholytic treatment combined with aerobic exercise might normalize the cardiovagal baroreflex and provide therapeutic benefit in patients with POTS. (Hypertension. 2021;77:1234-1244. DOI: 10.1161/HYPERTENSIONAHA.120.16113.)

Data Supplement

Key Words: baroreflex ▪ digoxin ▪ orthostasis ▪ modified oxford method ▪ postural tachycardia syndrome ▪ pyridostigmine

Standing upright (orthostasis) produces central hypovolemia by translocating blood to the dependent body. Orthostatic intolerance is defined by signs and symptoms, such as lightheadedness, exercise intolerance, tachycardia, hypotension, hypertension, headache, fatigue, cognitive deficits, and nausea, while upright relieved by recumbence.1,2

Postural tachycardia syndrome (POTS) is defined in adults by orthostatic intolerance symptoms plus an excessive increase in heart rate exceeding 30 beats/min within 10 minutes of standing or upright tilt.3
Systolic arterial pressure (SAP) is maintained and may increase. Larger heart rate increments by 40 beats/min or more are required to diagnose POTS in younger patients <19 years old. More than 85% are women aged 14 to 50 years.

POTS resembles hypovolemia with reduced systemic venous return and reduced cardiac output (CO). In some instances, absolute hypovolemia is found; in others, a redistributive central hypovolemia occurs when upright. The latter is designated neuropathic POTS and results from decreased adrenergic vasoconstriction in the legs, or splanchnic vasculature, causing baroreflex unloading, decreased reflex cardiovagal stimulation, increased reflex sympathetic excitation, and reflex tachycardia. An alternative form of POTS in which sympathetic adrenergic function is primarily increased is denoted hyperadrenergic POTS.

In both cases, baroreflex regulation is affected. Changes in blood pressure (BP) during orthostasis are primarily buffered by arterial baroreflexes, which are negative feedback inhibitory stretch reflexes. An increase in arterial pressure causes decreased heart rate mediated primarily by parasympathetic (cardiovagal) efferent activity, decreased peripheral vasoconstriction mediated primarily by reduced vascular synaptic sympathetic activity, and suppression of adrenal medullary epinephrine secretion. POTS patients have normal muscle sympathetic nerve activity while supine, but increased muscle sympathetic nerve activity when upright compared with healthy controls suggesting a reflex response with intact arterial baroreflexes afferents.

Since POTS is characterized by tachycardia, investigations have frequently focused on cardiovagal baroreflex impairments. Methods, such as transfer function analysis, are often used that measure small spontaneous fluctuations of HR (or RR interval) and SAP to obtain a linear frequency domain relationship between RR variability and SAP variability near 0.1 Hz, low frequency in which the slope, ∆RR/∆SAP, is thought to represent the sensitivity or gain characterizing the cardiovagal baroreflex. Low-frequency SAP oscillations are thought to reflect sympathetic activity, while low-frequency RR interval oscillations result from cardiovagal baroreflex transduction of SAP oscillations. The slope of ∆RR/∆SAP is typically reduced in POTS compared with controls supine and even more when upright.

Since spontaneous fluctuations generate information over a very narrow range of arterial pressures their ability to accurately represent baroreflex function, particularly during orthostatic stress, is questionable. This is because the cardiovagal baroreflex is often characterized by its slope (sensitivity, gain) at a single point referred to as the operating point, the particular SAP and its corresponding RR interval defining the state of the system at which the slope is measured. But cardiovagal baroreflex cannot be defined at a single point. Rather, it is defined

Nonstandard Abbreviations and Acronyms

| Abbreviation | Meaning       |
|--------------|---------------|
| BP           | blood pressure|
| CO           | cardiac output|
| HUT          | head-up tilt  |
| POTS         | postural tachycardia syndrome |
| SAP          | systolic arterial pressure |
| SNP          | sodium nitroprusside |

Summary

Cardiovascular baroreflex deficits in POTS relate to parasympathetic withdrawal while supine, remediated completely by digoxin, and to sympathetic activation while upright distorting the baroreflex response range. Since these baroreflex effects resemble those measured following microgravity/chronic bedrest, vagotonic/sympatholytic treatment combined with aerobic exercise might normalize the cardiovagal baroreflex and provide therapeutic benefit in patients with POTS.

Novelty and Significance

What Is New?

- We showed differences in characteristics of the cardiovagal baroreflex in patients with postural tachycardia syndrome (POTS) compared with control both supine and while upright during an orthostatic challenge.
- Parasympathetic stimulation with digoxin remediated the baroreflex deficit in POTS while supine.
- Parasympathetic stimulation with both pyridostigmine and digoxin partially corrected the baroreflex deficit in POTS while upright.

What Is Relevant?

- Patients with POTS have baroreflex deficits while upright shown using the nonlinear modified Oxford method.
- Parasympathetic stimulation with pyridostigmine and digoxin partially corrected this deficit.

- Since many patients with POTS are bed rested and physically deconditioned, a combination of exercise reconditioning which increases parasympathetic activity and reduces sympathoexcitation, plus pyridostigmine/digoxin may prove beneficial for treatment of POTS.
by the nonlinear, functional relationship between the RR interval and the SAP, closely fit by a sigmoidal (logistic) function over a wide range of arterial pressures.\textsuperscript{23}

Slopes obtained at operating points cannot define the cardiovagal baroreflex, but rather represent a changeable aspect of the current state of the system, corresponding to the variation of RR interval with variation of SAP derived from spontaneous fluctuations of RR interval and SAP. Results are only applicable near that particular operating point\textsuperscript{24} and not elsewhere; the assumption that the slope at the operating point estimates the maximum cardiovagal baroreflex gain, G\textsubscript{max}, may be erroneous. As shown below in Figure 1, if the operating point is near the centering point where the curve is nearly linear (ie, operating point 1), its slope will closely approximate G\textsubscript{max}; otherwise, the slope is less than G\textsubscript{max} and can neither accurately reflect G\textsubscript{max} nor assist in constructing the baroreflex relationship.\textsuperscript{24} Thus, for example, if the SAP corresponds to operating point 2, the slope at that point is much smaller than G\textsubscript{max}.

We have shown previously in healthy volunteers that orthostasis shifts or resets the centering point, saturation, and threshold en bloc, while G\textsubscript{max} and response range are unchanged.\textsuperscript{25} BP can be modestly affected resulting in different operating points. Resetting is attributable to the modulating effects of cardiopulmonary baroreflexes on arterial baroreflexes with decreased cardiac filling in the upright position.\textsuperscript{26} Often, operating points are almost directly displaced downwards, and the local slope at OP-upright may not accurately estimate G\textsubscript{max}.

The purpose of this investigation was, therefore, to examine whether and how cardiovagal baroreflex relationships in POTS differ from healthy controls in both the supine and upright positions. We used the so-called modified Oxford method to construct the sigmoidal baroreflex curve and tested whether parasympathetic stimulation with pyridostigmine (Mestinon)\textsuperscript{26} or digoxin\textsuperscript{27} can correct baroreflex deficits.

**METHODS**

The data that support the findings of this study can be made available from the corresponding author upon reasonable request.

**Subjects**

We enrolled 36 female patients with POTS aged 15 to 30 (mean=21.7±7.7 SD) and 20 healthy female volunteer subjects aged 18 to 24 years (median=21.3±5.2 SD years) of similar weight, height, and body mass index. POTS subjects fulfilled POTS criteria defined previously. Symptoms of orthostatic intolerance were present on a daily basis for >6 months and were relieved once supine. Symptoms were replicated during a prior 10 minutes, 70° upright tilt showing excessive tachycardia without hypotension or syncope. A similar upright tilt test on control subjects showed the absence of orthostatic intolerance. Only female subjects were employed because we could not sufficiently power a study of male patients with POTS as the majority of patients with POTS are female.\textsuperscript{8} All subjects were free of systemic or infectious disease. Subjects were either therapeutically naive or weaned off drugs for at least 2 weeks. Only nonpharmaceutical birth control was employed by subjects during the study. Subjects did not use nicotine. Subjects refrained from caffeine for at least 72 hours before testing. Subjects fasted for a minimum of 4 hours before testing. There were no trained athletes or bedridden subjects. Studies were performed in the midluteal phase when autonomic tone is maximized.\textsuperscript{28} Informed consent was obtained from all participants. All protocols were approved by the Committee for the Protection of Human Subjects of New York Medical College.

**Protocol**

Testing began at 9:30 am. Subjects were instrumented for ECG, beat-to-beat AP using a Finometer (FMS, Amsterdam) calibrated against an oscillometric BP cuff. CO was estimated using the Finometer which models the circulation as an adaptive Windkessel. Systemic vascular resistance was computed using the formula systemic vascular resistance = mean arterial pressure/CO (mm Hg/L per minute). This was initially calibrated against an inert gas rebreathing (Innocor, Innovision, Denmark) while supine before experiments began. An intravenous catheter was placed in the left antecubital vein.

Supine and upright cardiovascular baroreflex relationships were obtained using the modified Oxford method described below, and estimate of the slope ΔRR/ΔSAP (spontaneous baroreflex sensitivity) at the operating point were calculated by Fourier based transfer function analysis, when squared coherence >0.5, and by autoregressive modeling.\textsuperscript{19}

1. Following a 30-minute rest period, data were recorded throughout a 10-minute supine baseline period. During this and all subsequent study periods, HR and AP data were used for the calculation of RR interval variability,
SAP variability, and baseline spontaneous baroreflex sensitivity.

2. Following baseline measurements, the modified Oxford method was performed in the supine position. For this and all subsequent uses, performance of the modified Oxford method consisted of an intravenous bolus injection of 100 µg sodium nitroprusside (SNP) was administered followed 1 minute later by an intravenous bolus injection of 150 µg phenylephrine. Subjects were then allowed to recover to resting levels over a 30-minute period, and the method was repeated.

3. Next was a 2-hour period for drug absorption (no drug given in control or POTS on day 1=untreated).

4. A second baseline data collection for spontaneous fluctuation variabilities was obtained after 2 hours and a second supine modified Oxford method was performed.

5. After a 30 minutes recovery, subjects were tilted to 40° upright for 10 minutes. Data from minutes 2 to 7 were used for the calculation of upright spontaneous baroreflex sensitivity. The modified Oxford method was repeated once at 8 minutes to avoid syncope. After upright HR and AP returned to near pre-Oxford levels, subjects were returned to the supine position. All subjects were able to tolerate a 40° tilt + modified Oxford.

Control subjects underwent 1 day of testing without medication, while patients with POTS repeated the protocol during 3 additional days during which they received oral placebo, oral pyridostigmine 60 mg, or oral digoxin 500 mg. One of the 3 drugs was given just before step 3. Digoxin was given last because of its long (≈36 hours) half-life. The intervening 2 days were used to administer placebo and pyridostigmine in random order. The dose of pyridostigmine corresponds to a short-acting therapeutic dose. The dose of digoxin corresponds to a half loading dose for an adult and plasma levels reach the therapeutic range in 2 hours.

A flow sheet of the course of study is shown in Figure 2.

**Data Analysis**

Approximately 5 minutes of baseline data were analyzed preceding the modified Oxford method in the supine position. Upright HR and AP data were obtained during minutes 2 to 7 of head-up tilt (HUT) to avoid the hemodynamic equilibration phase that occurs during the first minutes of tilt.

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**Figure 2.** It shows the flow sheet and design of the described studies.

The experimental protocol is shown on top and the administration of drugs is shown on the bottom.
were collected continuously during the hypotensive and hypertensive phases of the modified Oxford method in both supine and upright positions and included ECG, CO, systolic (SAP), diastolic AP, and mean arterial pressure calculated using the Finometer calibrated against oscillometric cuff pressure. RR interval and HR were calculated from the ECG. Data were sampled at 200 Hz using custom signal processing software and analyzed offline.

Cardiovagal Baroreflex Determined by the Modified Oxford Method

RR intervals were plotted as a function of SAP to generate the cardiovagal baroreflex response during the modified Oxford method, and the methods for calculation of the cardiovagal baroreflex is described in the Data Supplement using established methods.\textsuperscript{23,24}

Spontaneous Cardiovagal Baroreflex Indices

To measure spontaneous cardiovagal baroreflex function, the slope corresponding to spontaneous fluctuations in SAP and RR intervals were calculated during supine baseline, supine 2 hours after drug administration, and during 40° upright tilt before the modified Oxford method, as described in the Data Supplement using established methods.\textsuperscript{23,24,32}

Statistics

A repeated-measures analytic approach was used. We used a multilevel approach using linear mixed models to account for the within-subject repeated measurements,\textsuperscript{35,36} as described in the Data Supplement.

RESULTS

The following is a summary of the hemodynamic data in patients with POTS while supine before (supine premed), and after (supine postmed) treatment with placebo, pyridostigmine or digoxin, or controls, and during upright tilt (tilt) and are shown in Table S1 of the Data Supplement. Data averaged over 15 seconds is reported for pre-SNP. Data at the time of minimum SAP are reported for SNP and at the time of maximum SAP is reported for phenyl.

Hemodynamics Before and During the Modified Oxford Method

HR Sodium Nitroprusside

SNP-premed increased HR in all groups but most in patients with POTS. A proposed nitrergic nitric oxide deficiency in POTS may produce a potentiated response to exogenous nitric oxide.\textsuperscript{38} The addition of pyridostigmine blunted the SNP-mediated HR increase in both POTS-pyridostigmine postmed and POTS-pyridostigmine tilt. This may indicate ganglionic potentiation of nitrergic parasympathetic nerves.

HR Phenyl

There was a relative reduced bradycardic response to phenylephrine in POTS-tilt groups except for POTS-pyridostigmine.

SAP Presodium Nitroprusside

In contrast, SAP was reduced when supine in POTS unmedicated, increasing with the addition of pyridostigmine or digoxin to control levels while remaining supine. With tilt, SAP was reduced in all POTS compared with control regardless of treatment. This corresponds to a left shift of the operating point of baroreflex curves for POTS compared with control independent of medication.

SAP Sodium Nitroprusside

SNP produced similar reductions of systolic BP when supine, independent of medication. Systolic BP increased during tilt in control but remained unchanged in POTS treatment groups.

SAP Phenyl

SAP response to phenylephrine was similar for all POTS compared with control except for POTS-pyridostigmine which blunted systolic BP increase.

Diastolic AP Pre-SNP

This was reduced in all POTS compared with control pre-SNP when tilted.

CO and Systemic Vascular Resistance

CO during tilt in POTS was similar to control pre-SNP; CO during tilt was lower in POTS than control during SNP, except for POTS-digoxin, which was similar to control, and higher than control during phenyl in POTS except for POTS-digoxin. CO was enhanced by phenylephrine when upright. Systemic vascular resistance was decreased in POTS supine and upright compared with control.

Sigmoidal Baroreflex Results From the Modified Oxford Method

In addition to the fitted cardiovagal baroreflex parameters ($G_{max}$, SAP, and RR interval at center point, threshold, saturation, and response range), we tabulated the SAP and RR interval at operating points (Table).

Supine and Supine Postmed

As shown in the Table, $G_{max}$ supine in POTS and control subjects was smaller in POTS compared with control.
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G\text{max} was unchanged in untreated or placebo-treated POTS but increased, becoming similar to control after either pyridostigmine or digoxin were administered.

**Centering Point, Response Range**
Based on the results shown in the Table and depicted in Figures 3 through 5, G\text{max}, which by definition occurs at the centering point, was displaced downwards (smaller RR interval) and to the left (lower SAP) in POTS compared with control. This is not a simple resetting because the shape of the sigmoidal function is not preserved; rather, G\text{max} and the response range are both reduced while supine in POTS. Administration of either pyridostigmine or digoxin repaired the BP location of the centering point as well as G\text{max}, but only digoxin corrected its displacement along the RR interval axis and the response range. Thus, digoxin repairs the supine cardiovagal baroreflex in POTS.

**Operating Point and Operating Point Slope**
The operating point response is expressed as a percentage of the response range from the threshold. Relative to control, untreated or placebo-treated POTS shifted the operating point towards lower percentile, that is, towards the knee of the threshold where the sigmoidal nonlinear and flat. Control remained near the center. RR interval was reduced in POTS. The operating point slope was much smaller in premed POTS than in control. This corresponds to a reduction in the alpha index estimate of operating point gain (see below), and, therefore, a marked reduction of heart rate variation for given degree of BP variation (the sigmoid curve is flat) and a general

### Table. Baroreflex Results From Sigmoidal Curves

| Parameter | Condition | Control | POTS-untreated | POTS-placebo | POTS-pyridostigmine | POTS-digoxin |
|-----------|-----------|---------|---------------|--------------|--------------------|--------------|
| G\text{max} (max gain, slope), ms/mm Hg | Supine premed | 28.8±3.6 | 20.2±3.5* | 20.7±3.4* | 18.9±3.9* | 19.5±5.0* |
| | Supine postmed | 24.2±3.0 | 19.7±2.5* | 18.4±3.0* | 25.8±3.0 | 27.3±4.0 |
| | Tilt | 24.9±1.5 | 10.9±2.7* | 13.4±1.8* | 25.0±2.8* | 19.0±4.2* |
| SAP at G\text{max}, mm Hg, center point | Supine premed | 117±3 | 112±6 | 113±4 | 111±3 | 114±3 |
| | Supine postmed | 119±2 | 113±3* | 115±3* | 118±5 | 122±2* |
| | Tilt | 121±6 | 116±4 | 118±5 | 119±4* | 121±8* |
| RR at G\text{max}, ms, center point | Supine premed | 890±50 | 709±45* | 710±45* | 799±31* | 856±38* |
| | Supine postmed | 900±50 | 709±45* | 710±45* | 799±31* | 856±38* |
| | Tilt | 771±31* | 616±25*† | 632±47* | 680±32 | 688±45 |
| Threshold, ms | Supine premed | 666±21 | 572±26* | 579±26* | 547±35* | 561±15* |
| | Supine postmed | 644±21 | 548±23* | 545±28* | 616±28* | 573±24 |
| | Tilt | 579±21* | 489±21*† | 498±17*† | 564±24* | 526±30* |
| Saturation, ms | Supine premed | 1094±91 | 938±92 | 922±83 | 879±101 | 940±59 |
| | Supine postmed | 1157±92 | 870±27* | 874±27* | 946±42 | 1138±76* |
| | Tilt | 843±106* | 744±45* | 806±97 | 796±27* | 796±27* |
| Response range, ms | Supine premed | 427±27 | 366±16* | 342±42* | 359±38* | 338±48* |
| | Supine postmed | 522±44 | 348±23* | 329±35* | 330±52* | 564±52* |
| | Tilt | 385±42 | 255±50*† | 316±53 | 232±28*† | 251±41*† |
| Slope (gain) at O.P., ms/mm Hg | Supine premed | 18.9±3.1 | 14.1±3.4* | 13.7±3.2* | 11.6±4.4* |
| | Supine postmed | 17.9±3.5 | 12.8±2.2* | 11.4±2.4* | 22.1±3.5* | 16.0±3.9* |
| | Tilt | 19.9±3.5 | 6.5±2.7*† | 6.8±1.6*† | 6.3±3.3*† | 8.2±2.7* |
| Operating point (% response) | Supine premed | 59±11 | 36±9* | 29±11* | 34±8* | 37±11 |
| | Supine postmed | 62±10 | 40±10* | 28±6* | 66±9 | 47±13 |
| | Tilt | 54±11 | 24±11* | 27±12* | 18±10* | 26±5* |
| SAP at O.P., mm Hg | Supine premed | 118±2 | 117±4 | 112±4 | 112±3 | 113±4 |
| | Supine postmed | 120±3 | 113±4* | 109±5* | 117±7 | 118±4 |
| | Tilt | 124±6 | 111±5* | 112±3* | 110±6* | 115±5* |
| RR at O.P., ms | Supine premed | 893±37 | 736±54* | 653±42* | 687±52* | 716±55* |
| | Supine postmed | 927±48 | 700±44* | 634±41* | 854±39* | 746±42* |
| | Tilt | 834±32* | 585±45*† | 571±27*† | 643±39* | 605±44* |

O.P. indicates operating point; POTS, postural tachycardia syndrome; and SAP, systolic arterial pressure.

*P<0.05 compared with control.
†P<0.05 compared with supine premed.
‡P<0.05 compared with POTS-untreated.
loss in the buffering effects of HR on BP. Treatment with either pyridostigmine or digoxin brought the operating point closer to the centering point and the operating point slope increased.

**Tilt**

$G_{max}$

With tilt, $G_{max}$ remained similar to control when treated with pyridostigmine or digoxin but fell for untreated or placebo-treated POTS.

**Centering Point, Response Range**

The centering point for control was shifted downwards (higher HR when upright) and slightly to the right. The response range remained unchanged and the operating point moved closer to the centering point optimizing BP buffering. Operating point slope (sensitivity) thus more closely approximates $G_{max}$ for control. Centering points for POTS-untreated or POTS-placebo moved downwards (higher HR) and to the left (higher SAP) but response range remained reduced compared with control. The operating point moved sharply to the left and thus the operating point slope was greatly reduced. Pyridostigmine or digoxin shifted the centering point to lower RR interval but response range and operating point moved sharply to the left such that the slope at the operating point was comparable to POTS-untreated and much lower than control.

The sigmoidal baroreflex results from the modified Oxford Method were used to generate Figures 3 through 5 using mean values for sigmoidal parameters averaged over all subjects, from data in the Table. In all determinations, POTS-placebo results were similar to untreated and are not depicted graphically.

Figure 3 shows POTS-untreated both supine and following HUT. $G_{max}$ and the response range remained unchanged with tilt in control, is reduced in POTS, and further reduced in POTS when tilted. Additional key features of POTS include left lateral and downwards shifts of centering points and operating points and decreased response range with tilt. Placebo-treated results were not different from untreated POTS.

Figure 4 compares control with POTS-pyridostigmine. $G_{max}$ is restored to near control values supine and following HUT. Supine SAP at centering and operating points are similar to control, but threshold is displaced downward and response range is curtailed. Centering point SAP and $G_{max}$ were sustained but response range and operating point were greatly reduced during HUT and contributed to the left shift of the operating point for pyridostigmine with consequent reduced slope. It appears, therefore, that a reduction in response range with unchanged centering point and $G_{max}$ may account for left-shifted operating points.

Figure 5 compares control with POTS-digoxin supine and following HUT. Digoxin sustains the centering point and $G_{max}$ at close to control values. Digoxin, alone, increases response range supine because of a reduced threshold. Tilted, threshold was further reduced. SAP at centering point and $G_{max}$ were sustained but response range fell precipitously with tilt and contributed to the left shift of the operating point for digoxin and consequently reduced slope.

**Spontaneous Fluctuations—Comparison With Modified Oxford Method Results**

Table S2 shows total RR interval and SAP variability, and variability measures over low-frequency and high-frequency bands. Baroreflex gain (local slope) is presented for the sequence method and for the $\alpha$-index method obtained from transfer function analysis.

**Baroreflex Sensitivity (Gain)**

Alpha index and sequence methods yielded similar results. These indices were larger in control than in
POTS, increased by pyridostigmine and digoxin comparable to control when supine postmed, and decreased once tilted. Indices in control decreased to a lesser extent. Pyridostigmine ameliorated the decrease with tilt. Slopes from sigmoidal baroreflex at the operating point (Table) were similar to spontaneous fluctuation alpha indices.

**DISCUSSION**

Our data are the first to show deficits in both supine and upright cardiovagal baroreflex in POTS, measured using the reference standard modified Oxford method. While supine, deficits were primarily reflective of severely reduced vagal activity. Although G_max was reduced that did not account for the majority of vagal loss. Rather, the response range was reduced, and centering points shifted to lower SAP and RR. This combination resulted in movement of operating points to the flat-curvilinear portion of the baroreflex function with consequent profound reductions in operating point slope. Comparing the responses shown in Figures 3 through 5, in patients with POTS, pyridostigmine ameliorated and digoxin corrected these deficits while supine: G_max was similar to control, centering point SAP was similar to control although the RR interval at the centering point SAP was shifted towards lower RR intervals. The response range exceeded control for digoxin.

Upright posture in POTS elicited more marked baroreflex impairment compared with supine. For POTS-untreated G_max decreased, response range decreased while centering points were conserved (being shifted to the left and downwards as when supine). Operating points are shown by blue and black circles. G_max is restored to near control values supine and upright. Supine systolic arterial pressure (SAP) at centering and operating points are similar to control, but threshold is displaced downwards and response range is curtailed. During tilt centering point, SAP and G_max were sustained but response range was greatly reduced during tilt contributing to the left shift of the operating point.

**Figure 4.** It compares patients with postural tachycardia syndrome (POTS) treated with pyridostigmine (blue lines) with control (black lines) supine and during head-up tilt (HUT). Operating points are shown by blue and black circles. G_max is restored to near control values supine and upright. Supine systolic arterial pressure (SAP) at centering and operating points are similar to control, but threshold is displaced downwards and response range is curtailed. During tilt centering point, SAP and G_max were sustained but response range was greatly reduced during tilt contributing to the left shift of the operating point.

**Figure 5.** It compares patients with postural tachycardia syndrome (POTS) treated with digoxin (teal lines) with control (black lines) supine and during head-up tilt (HUT). Operating points are shown by cyan and black circles. Digoxin returns the centering point and G_max of patients with POTS to similar to control values. The response range supine postdigoxin is greater in treated POTS than control. After tilt, threshold was more in POTS while systolic arterial pressure (SAP) at centering point and G_max were sustained. However, response range decreased markedly with tilt in POTS and contributed to the left shift of the operating point for digoxin and consequently reduced operating point slope.
slopes were on the flat portion of the curve, and operating point slope was very low. Both digoxin and pyridostigmine shifted the operating point to the vicinity of the knee of the sigmoid curve where the second derivative is maximum. Because of these shifts, an increase in SAP can give rise to a large increase in RR interval while a small decrease in SAP fails to appreciably lower RR interval. The system then responds asymmetrically and buffers increases in SAP with a decrease in HR but with little buffering capacity when SAP decreases.

Impaired upright cardiovagal baroreflex is not corrected by pyridostigmine or digoxin, although $G_{\text{max}}$ and centering point are reset to a lower RR interval while SAP is reasonably well sustained. Also, SAP at the operating points is only slightly diminished compared with supine. However, although SAP only changes slightly, the operating points for both pyridostigmine and digoxin are shifted to the left onto the flat portion of the curve where slope is quite small. This is the result of reduced response range. Thus, for example, if the response range is small while $G_{\text{max}}$ and centering points remain unchanged, the linear portion of the sigmoidal curve occurs between narrow SAP limits. Then, even a small left shift in SAP can shift the operating point to the flat part of the sigmoidal curve where response to change is reduced.

Sympathetic activation reduces the response range to baroreflex stimulation, and POTS has a much larger increase in sympathetic activity when upright compared with control, and SAPV-low frequency is an indicator of sympathetic activation which increase in SAPV-low frequency in POTS when upright. The relative reduction of the bradycardic response to phenylephrine during tilt in POTS except for POTS-pyridostigmine is consistent with anti-$\alpha$, adrenergic antibodies recently reported.

Cardiovagal deficits are described by changes in the functional sigmoidal relationship between SAP and RR interval. A resetting of the baroreflex (centering point translation) accommodates changes in posture, BP, or activity by changing the location of the sigmoidal curve without changing its shape. Apart from resetting, $G_{\text{max}}$ and response range suffice to define the sigmoidal curve. The position along the curve (the operating point) offers an additional degree of freedom and can be defined by determining the extant SAP or RR interval. The cardiovagal baroreflex cannot be defined by the slope at a single point but can only be usefully defined as a functional relationship between SAP and RR interval over a wide range of BPs. The cardiovagal baroreflex helps to buffer beat-to-beat changes in BP by changing HR in the opposite direction. If the operating point is uniformly placed on a flat portion of the sigmoidal curve, then HR cannot vary much with changing SAP, and changes in HR inadequately compensate for either increasing or decreasing BP. As a consequence, BP variability is enhanced at low frequency with tilt. We have previously shown that this interferes with cerebral autoregulation and cognitive capability.

Local slope at the operating point may characterize the local change in RR interval with BP and may, therefore, be considered to represent local baroreflex function but it cannot predict the sigmoid baroreflex relationship. Rather, centering point location which is reset with orthostasis and the distance of the operating point from the centering point, determined by the relations of $G_{\text{max}}$ with threshold and response range yield operating point locations. These may be on the linear portion of the curve close to the centering point, as in control, in which case the local slope approximates $G_{\text{max}}$ or on the flat portion of the curve distant from the centering point, as in untreated POTS, in which case the local slope is far smaller than $G_{\text{max}}$.

Pyridostigmine and digoxin each have vagotonic actions and repair the supine cardiovagal baroreflex in POTS by normalizing $G_{\text{max}}$, correcting the centering point, and reexpanding the response range. The resulting operating point falls on the linear portion of the curve and slopes become good estimates of $G_{\text{max}}$. Digoxin is also sympatholytic while pyridostigmine may also exert sympathotonic effects.

When tilted upright, SAPV is elevated, specifically in the low-frequency range. This is consistent with increased sympathetic activity. Downward resetting of the centering point and reduced response range conforms to increased sympathetic activity which may explain their abnormalcy in POTS in which sympathetic activation is excessive when upright. Neither pyridostigmine nor digoxin maintain the baroreflex once upright because increased cardiac sympathetic activity reduces the response range which left shifts the operating point without important changes in SAP. Potential causes of excessive sympathetic activation in POTS include absolute or positional central hypovolemia, intrinsically increased sympathetic CNS outflow as proposed in hyperadrenergic POTS, or increased central command in the face of postural muscle atrophy. Sympatholytic agents, in addition to digoxin or pyridostigmine, should increase the response range and normalize baroreflex control.

Administration of either pyridostigmine or digoxin repaired the SAP of the centering point as well as $G_{\text{max}}$. However, digoxin corrected its displacement along the RR interval axis and the response range. Pyridostigmine enhances acetylcholine and nitric parasympathetics, resulting in increased nitric nitric oxide which acts at prejunctional and postjunctional sites to reduce adrenergic sympathetic transduction. Reduction of $G_{\text{max}}$, reduced response range, leftwards and downwards translation of the centering point are also evident in microgravity deconditioning, which can be simulated by bed rest deconditioning and to a lesser degree by cardiovascular deconditioning.
Many of our patients with POTS are bed rested and cardiovascular deconditioned. Our findings are, therefore, highly consistent with studies indicating that exercise reconditioning may be the ideal treatment for POTS because it increases parasympathetic and decreases sympathetic cardiac activity. In addition, since digoxin was able to repair the SAP of the centering point and $G_{\text{max}}$ and correct its displacement along the RR interval axis and response range, the use of this drug in combination with directed exercise/reconditioning may be a beneficial treatment modality for patients with POTS.

**LIMITATIONS**

Least squared fits of threshold, saturation, centering point, and response range are model-dependent and obtained after nonlinear fitting of data to the sigmoid curve. We always captured the threshold and most linear portion of the curve (the center) and some portion of the saturation in every subject.

The mean age of our study subjects was relatively young, and results may not generalize to older age groups. Data were collected only from female subjects since POTS is predominantly a female illness from menarche to menopause. Male and female physiology may differ; there was no way we could sufficiently power a study of POTS in males as well as females.

Medication was not delivered in a completely random manner and, therefore, its sequential delivery, combined with the training effect of repeated tilt testing may have introduced bias. Also, both pyridostigmine and digoxin have additional effects other than increasing vagal tone. Administration of drugs that elicit multiple drug effects can be problematic when studying systemic outcomes, however, we used these because both have well known vagotonic effects.

**PERSPECTIVES**

Orthostasis produces central hypovolemia by translocating blood to the dependent body. POTS is defined by orthostatic intolerance and increased in heart rate and maintenance or increase in arterial pressure. Orthostatic changes in BP are buffered by arterial baroreflexes. The cardiovasal baroreflex regulation is deficient in POTS by reducing the local slope at the operating point estimated by spontaneous fluctuations of heart rate and BP over a very narrow range of arterial pressures; local slopes may not represent true baroreflex function which can be assessed by reference standard modified Oxford method to generate the nonlinear sigmoidal relationship between the RR interval and the SAP over a wide range of arterial pressures. We describe the determinants of abnormal cardiovascular baroreflex response in POTS, supine and upright, compared with control. Supine baroreflex abnormalities were corrected by digoxin and ameliorated by Mestinon, although sympathetic stimulation of orthostasis produced further baroreflex deficits.

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