Neurology® Clinical Practice

Hydration status substantially affects chronic cerebrospinal venous insufficiency assessments

Claudiu I. Diaconu, BS
Robert J. Fox, MD
Alia Grattan, RVT
Alexander Rae-Grant, MD
Mei Lu, MD, PhD
Heather L. Gornik, MD, MPH, RPVI
Esther Soo H. Kim, MD, MPH, RPVI

Summary
We sought to determine the effect of hydration on the criteria for chronic cerebrospinal venous insufficiency (CCSVI), a proposed hypothesis for the etiology of multiple sclerosis (MS). Sixteen subjects (11 MS and 5 controls) were asked to fast overnight. The following morning, 2 CCSVI ultrasound examinations were performed: 1 in the mildly dehydrated state, and another 30–45 minutes after rehydrating with 1.5 L of Gatorade. Seven subjects fulfilled CCSVI criteria in the dehydrated state. Of these, 5 (71%) no longer fulfilled CCSVI criteria after rehydration. One additional subject met CCSVI criteria only after rehydration. Hydration status has a substantial effect on CCSVI criteria, suggesting that the sonographic findings of CCSVI may represent a physiologic rather than pathologic state.

The chronic cerebrospinal venous insufficiency (CCSVI) hypothesis posits that abnormalities in the extracranial venous outflow contribute to the pathogenesis of multiple sclerosis (MS), although results from different groups vary widely. The chronic cerebrospinal venous insufficiency (CCSVI) hypothesis posits that abnormalities in the extracranial venous outflow contribute to the pathogenesis of multiple sclerosis (MS), although results from different groups vary widely.

Cleveland Clinic Lerner College of Medicine of Case Western Reserve University (CID), Mellen Center for Multiple Sclerosis Treatment and Research (RJF, AR-G), Heart and Vascular Institute, Cardiovascular Medicine, Non-Invasive Vascular Laboratory (AG, HLG, ESHK), and Neurological Institute, Cerebrovascular Center (ML), Cleveland Clinic, Cleveland, OH.

Funding information and disclosures are provided at the end of the article. Full disclosure form information provided by the authors is available with the full text of this article at Neurology.org/cp.

Correspondence to: kims@ccf.org

© 2013 American Academy of Neurology

© 2013 American Academy of Neurology. Unauthorized reproduction of this article is prohibited.
A key method used for assessment of CCSVI involves ultrasound of the internal jugular veins (IJVs), vertebral veins (VVs), and deep cerebral veins (DCVs); catheter venography is not as widely used in CCSVI assessments due to its invasive technique. Two or more of 5 sub-criteria must be met to fulfill the criteria for CCSVI: 1) reflux in IJVs or VVs, 2) reflux in the DCVs, 3) stenosis of IJV, 4) no flow detected in IJVs or VVs, and 5) reverted postural flow. To date, no studies have investigated the relation between volume status and CCSVI. There is anecdotal evidence, both from patients and from experience of clinicians, that people with MS tend to be more dehydrated than the general population. Drinking less fluid can help patients self-manage neurogenic bladder symptoms. By measuring urine osmolality, one study demonstrated that 42% of subjects with MS were not adequately hydrated. Our goal was to evaluate the effect of hydration status on CCSVI ultrasound results.

**METHODS**

**Standard protocol approvals, registrations, and patient consents**

The study was reviewed and approved by the Cleveland Clinic Institutional Review Board. All participants provided written informed consent to participate.

**Patients**

Sixteen subjects (11 subjects with MS and 5 healthy controls) were recruited from an ongoing CCSVI study. Participants were required to have met 0–4 CCSVI subcriteria on their previous CCSVI ultrasound.

**Study design**

CCSVI ultrasound assessment was performed by a single trained sonographer and over-read by a vascular ultrasound physician familiar with CCSVI assessment, both of whom were blinded to disease status and number of CCSVI subcriteria previously met during the separate CCSVI study from which patients were recruited.

CCSVI subcriteria 1, 3, 4, and 5 were assessed. DCV reflux was not assessed in this hydration study since transcranial Doppler examination in the main study has been consistently negative for reflux. After a 12-hour fast, including abstention from fluids, a baseline “dehydrated” CCSVI ultrasound was performed. Patients were then asked to drink 1.5 L of Gatorade over 30–45 minutes, and a second “rehydrated” ultrasound was subsequently performed. Inferior vena cava (IVC) cross-sectional area (CSA) was measured during both studies, and change in IVC CSA was used as a quantitative marker for change in hydration status.

**Data analysis**

Changes in subcriteria 1, 3, 4, and 5 between the dehydrated and rehydrated states were recorded. The maximal change in CSA was recorded, regardless of whether it occurred in the right or left IJV. Two-tailed paired t tests were used to compare changes between the 2 hydration states. Because of the small sample size, outlier values were not used in the statistical calculations.

**RESULTS**

Sixteen subjects were recruited: 11 subjects with MS (9 relapsing-remitting MS, 2 primary progressive MS) and 5 healthy controls. Overall, there were 5 men and 11 women, with an average age of 54.9 years.
Seven subjects fulfilled the CCSVI criteria in the dehydrated state. Of these, 5 (71%) no longer fulfilled CCSVI criteria after rehydration (table 1). One additional subject met CCSVI criteria only after rehydration. The most common change in criteria was losing the stenosis subcriterion, followed by a change in reflux time. Rehydration decreased reflux time in 8 subjects and increased it in 7; the average change in reflux over all subjects was not significant (table 2). One subject had reverted postural control that normalized with hydration, whereas another had reverted postural control induced by hydration. In 3 subjects, flow became detectable in one vertebral vein in the supine position with rehydration. In 2 of these subjects, there was no detectable flow in one vertebral vein in the upright position despite rehydration, thus not changing subcriterion 4. After rehydration, IVC CSA increased on average by 14% ($p = 5.6 \times 10^{-6}$) and IJV CSA increased on average by 32% ($p = 0.0018$) (table 2).

**DISCUSSION**

Studies of CCSVI have yielded conflicting results, although a meta-analysis of 8 studies suggested a correlation between CCSVI and MS. Because the venous system is a low pressure
We found that 71% of subjects who met CCSVI criteria while dehydrated no longer met CCSVI criteria after rehydration.

and high compliance system, it is particularly affected by hydration status, and so we hypothesized that hydration may play a substantial role in CCSVI ultrasound assessments. We found that 71% of subjects who met CCSVI criteria while dehydrated no longer met CCSVI criteria after rehydration. Since patients with MS often prefer to drink less fluid because of bladder dysfunction, our results suggest that at least some of the differences between patients with MS and controls without MS in previous CCSVI studies may be explained by differential hydration between patients with MS and controls without MS. We could not draw definite conclusions from our sample size about the effect of hydration on CCSVI within MS subgroups. To our knowledge, no other study has investigated the effect of hydration on MS subgroups or on CCSVI.

Our observations suggest that the sonographic findings of CCSVI may not represent a pathologic state but instead may represent a physiologic state in which hydration status plays a substantial role. When interpreting CCSVI ultrasound findings, care must be taken to recognize the potential effect from patient hydration status.

### Table 2 Change in CCSVI parameters with hydration, as measured by ultrasound

<table>
<thead>
<tr>
<th>Subject no.</th>
<th>CSA: Dehydrated</th>
<th>CSA: Hydrated</th>
<th>% CSA Δ</th>
<th>Reflux: Dehydrated</th>
<th>Reflux: Hydrated</th>
<th>% Reflux Δ</th>
<th>% IVC CSA Δ</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0.38</td>
<td>0.39</td>
<td>3</td>
<td>1.125</td>
<td>1.300</td>
<td>16</td>
<td>84*</td>
</tr>
<tr>
<td>2</td>
<td>0.61</td>
<td>1.38</td>
<td>126</td>
<td>1.080</td>
<td>900</td>
<td>−17</td>
<td>16</td>
</tr>
<tr>
<td>3</td>
<td>0.21</td>
<td>0.88</td>
<td>319*</td>
<td>1.080</td>
<td>1,296</td>
<td>20</td>
<td>7</td>
</tr>
<tr>
<td>4</td>
<td>2.22</td>
<td>2.43</td>
<td>9</td>
<td>600</td>
<td>539</td>
<td>−10</td>
<td>22</td>
</tr>
<tr>
<td>5</td>
<td>0.70</td>
<td>0.86</td>
<td>23</td>
<td>351</td>
<td>364</td>
<td>4</td>
<td>2</td>
</tr>
<tr>
<td>6</td>
<td>1.33</td>
<td>1.44</td>
<td>8</td>
<td>560</td>
<td>1,066</td>
<td>90</td>
<td>33</td>
</tr>
<tr>
<td>7</td>
<td>0.29</td>
<td>0.36</td>
<td>24</td>
<td>820</td>
<td>701</td>
<td>−15</td>
<td>30</td>
</tr>
<tr>
<td>8</td>
<td>0.93</td>
<td>1.26</td>
<td>35</td>
<td>1,198</td>
<td>656</td>
<td>−45</td>
<td>16</td>
</tr>
<tr>
<td>9</td>
<td>1.79</td>
<td>1.91</td>
<td>7</td>
<td>830</td>
<td>842</td>
<td>1</td>
<td>14</td>
</tr>
<tr>
<td>10</td>
<td>0.48</td>
<td>0.83</td>
<td>73</td>
<td>938</td>
<td>700</td>
<td>−25</td>
<td>13</td>
</tr>
<tr>
<td>11</td>
<td>1.08</td>
<td>1.44</td>
<td>33</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>12</td>
<td>0.10</td>
<td>0.13</td>
<td>30</td>
<td>300</td>
<td>1,883</td>
<td>529*</td>
<td>14</td>
</tr>
<tr>
<td>13</td>
<td>0.75</td>
<td>1.11</td>
<td>48</td>
<td>891</td>
<td>737</td>
<td>−17</td>
<td>13</td>
</tr>
<tr>
<td>14</td>
<td>0.44</td>
<td>0.51</td>
<td>16</td>
<td>831</td>
<td>762</td>
<td>−8</td>
<td>4</td>
</tr>
<tr>
<td>15</td>
<td>0.63</td>
<td>0.58</td>
<td>−8</td>
<td>0</td>
<td>539</td>
<td>N/A</td>
<td>10</td>
</tr>
<tr>
<td>16</td>
<td>0.34</td>
<td>0.51</td>
<td>50</td>
<td>364</td>
<td>195</td>
<td>−46</td>
<td>2</td>
</tr>
<tr>
<td>Average</td>
<td>32</td>
<td></td>
<td></td>
<td>(0.0018)</td>
<td>(0.9492)</td>
<td>(5.6E-06)</td>
<td></td>
</tr>
</tbody>
</table>

Abbreviations: CCSVI = chronic cerebrospinal venous insufficiency; CSA = cross-sectional area; N/A = not applicable.

Values are shown for internal jugular vein (IJV) CSA, IJV reflux time, and inferior vena cava CSA in the dehydrated and hydrated states. Areas were measured in centimeters and reflux in milliseconds.

*Outliers and thus not included in the statistical calculations.
REFERENCES


STUDY FUNDING

Supported by a grant from the Cleveland Clinic Research Program Committee (RPC grant 2010-1074) and a Research Grant (RC 1004-A-5) from the National MS Society (USA).

DISCLOSURES

C.I. Diaconu reports no disclosures. R.J. Fox serves on scientific advisory boards for Biogen Idec and Novartis; serves on the editorial boards of Neurology® and Multiple Sclerosis; serves as a consultant for Allozyne, Avanir, Biogen Idec, Novartis, Questcor, EMD Serono, and Teva Neurosciences; and receives research support from Novartis and the National MS Society. A. Grattan reports no disclosures. A. Rae-Grant has received speaker honoraria from Novartis, Avanir, Biogen Idec, and Teva Neurosciences for speaking; serves on the Evidence Review Team for Neurology; receives publishing royalties for Handbook of Multiple Sclerosis (Springer Healthcare, 2010), Comprehensive Review of Clinical Neurology (Wolters Kluwer, 2012), and 5 Minute Consult in Neurology (Wolters Kluwer, 2012); serves on the speakers’ bureau for Biogen Idec; and receives research support from the NIH and the National MS Society. M. Lu has received research support from the National MS Society. H. Gornik serves on an internal DSMB for Roy Greenberg IDE; serves on the editorial board of Vascular Medicine and Current Treatment Options in Cardiovascular Medicine; is author on patents re: Non-invasive diagnosis of lower extremity peripheral artery disease by oscillometric means (license fee payments received) and Remote monitoring of anticoagulation (royalty payments received); serves as Medical Director, Non-Invasive Vascular Laboratory, Cleveland Clinic; and receives research support from Theravasc, Astra-Zeneca, and the NIH/NHLBI, E.S.H. Kim received consulting fees from Philips; received grant support from the American College of Cardiology supported by GE, as well as grant support directly from GE; reviews vascular ultrasound (15% clinical effort) at the Cleveland Clinic Non-Invasive Vascular Laboratory; and receives research support from the Multiple Sclerosis Society. Full disclosure form information provided by the authors is available with the full text of this article at Neurology.org/cp.
# Related articles from other AAN physician and patient resources

<table>
<thead>
<tr>
<th>Source</th>
<th>Title</th>
<th>Date and Volume</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Neurology</strong>®</td>
<td>Progressive multiple sclerosis is not associated with chronic cerebrospinal venous insufficiency</td>
<td>August 30, 2011;77:844-850.</td>
</tr>
<tr>
<td></td>
<td>Prevalence, sensitivity, and specificity of chronic cerebrospinal venous insufficiency in MS</td>
<td>July 12, 2011;77:138-144.</td>
</tr>
<tr>
<td><strong>Neurology Now</strong>®</td>
<td>This Way In: CCSVI and Multiple Sclerosis</td>
<td>July/August 2010;6:9-11.</td>
</tr>
<tr>
<td><strong>Neurology Today</strong>®</td>
<td>Two New Papers Question MS-CCSVI Theory</td>
<td>October 7, 2010;10:40-41.</td>
</tr>
</tbody>
</table>
Hydration status substantially affects chronic cerebrospinal venous insufficiency assessments
Claudiu I. Diaconu, Robert J. Fox, Alia Grattan, et al.
Neurol Clin Pract 2013;3;386-391 Published Online before print October 3, 2013
DOI 10.1212/CPJ.0b013e3182a78f15

This information is current as of October 3, 2013

Updated Information & Services
including high resolution figures, can be found at:
http://cp.neurology.org/content/3/5/386.full.html

References
This article cites 7 articles, 2 of which you can access for free at:
http://cp.neurology.org/content/3/5/386.full.html##ref-list-1

Citations
This article has been cited by 2 HighWire-hosted articles:
http://cp.neurology.org/content/3/5/386.full.html##otherarticles

Subspecialty Collections
This article, along with others on similar topics, appears in the following collection(s):
Autoimmune diseases
http://cp.neurology.org//cgi/collection/autoimmune_diseases
Multiple sclerosis
http://cp.neurology.org//cgi/collection/multiple_sclerosis
Ultrasound
http://cp.neurology.org//cgi/collection/ultrasound

Permissions & Licensing
Information about reproducing this article in parts (figures, tables) or in its entirety can be found online at:
http://cp.neurology.org/misc/about.xhtml#permissions

Reprints
Information about ordering reprints can be found online:
http://cp.neurology.org/misc/addir.xhtml#reprintsus

Neurol Clin Pract is an official journal of the American Academy of Neurology. Published continuously since 2011, it is now a bimonthly with 6 issues per year. Copyright © 2013 American Academy of Neurology. All rights reserved. Print ISSN: 2163-0402. Online ISSN: 2163-0933.