

Validity of the diagnostic criteria for chronic cerebrospinal venous insufficiency and association with multiple sclerosis

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ABSTRACT

Background: The chronic cerebrospinal venous insufficiency theory proposes that altered cerebral venous hemodynamics play a role in the pathophysiology of multiple sclerosis. We aimed to explore the validity of this hypothesis by assessing the diagnostic criteria for chronic cerebrospinal venous insufficiency in persons with and without multiple sclerosis.

Methods: We compared the proportion of venous outflow abnormalities between patients with multiple sclerosis and healthy controls using extracranial Doppler ultrasonography and gadolinium-enhanced magnetic resonance venography. Interpreting radiologists were blinded to the clinical status of participants.

Results: We enrolled 120 patients with multiple sclerosis and 60 healthy controls. High proportions of both patients (67/115 [58%]) and con-

trols (38/60 [63%]) met 1 or more of the proposed ultrasound criteria for diagnosis of chronic cerebrospinal venous insufficiency ($p = 0.6$). A minority of patients (23/115 [20%]) and controls (6/60 [10%]) fulfilled 2 or more of the proposed criteria ($p = 0.1$). There were no differences between patients and controls in the prevalence of each individual ultrasound criterion. Similarly, there were no differences in intracranial or extracranial venous patency between groups, as measured by magnetic resonance venography.

Interpretation: We detected no differences in the proportion of venous outflow abnormalities between patients with multiple sclerosis and healthy controls. Moreover, our study revealed significant methodologic concerns regarding the proposed diagnostic criteria for chronic cerebrospinal venous insufficiency that challenge their validity.

Competing interests: See end of article.

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Multiple sclerosis is an inflammatory disease of the central nervous system, believed to arise from a dysfunctional immune-mediated response in a genetically susceptible host.¹ In 2009, “chronic cerebrospinal venous insufficiency” was proposed to play an etiologic role in multiple sclerosis.²⁻⁴ Despite an abundance of published literature on this topic,²⁻²⁸ a causal link has not been established. Recent meta-analyses have suggested a strong association between an ultrasound-based diagnosis of chronic cerebrospinal venous insufficiency and multiple sclerosis,^{26,28} yet there has been significant heterogeneity across studies.^{26,27} A factor contributing to this heterogeneity appears to be the involvement of investigators who support endovascular procedures as a treatment for multiple sclerosis.²⁷ Furthermore, these meta-analyses have been predicated on the assumption that valid diagnostic criteria for chronic cerebrospinal venous insufficiency exist.

We aimed to explore the validity of the chronic cerebrospinal venous insufficiency the-

ory by using extracranial ultrasonography and gadolinium-enhanced magnetic resonance venography to compare the proportion of venous outflow abnormalities between patients with multiple sclerosis and healthy individuals. Our primary hypothesis was that if chronic cerebrospinal venous insufficiency is associated with multiple sclerosis, we would detect significant evidence of venous outflow obstruction in patients relative to controls.

Methods

Study design

This cross-sectional study involved participants who were evaluated at the University of Calgary. The protocol was approved by the institution’s Conjoint Health Research Ethics Board, and participants provided written, informed consent.

Study population

Hundreds of patients followed at the Calgary Multiple Sclerosis Clinic volunteered to participate in the study. We consecutively screened and enrolled

individuals from this group who met the inclusion criteria until we reached predetermined quotas for each subtype of multiple sclerosis. We recruited age-matched (by decile) and sex-matched controls from volunteers who expressed interest in the study. Eligible patients had clinically isolated syndromes,²⁹ relapsing–remitting multiple sclerosis proven by McDonald²⁹ and Poser³⁰ criteria, secondary progressive multiple sclerosis, primary progressive multiple sclerosis^{29,31} or antibody-positive neuromyelitis optica.³² Participants provided consent themselves or with the aid of guardian. Exclusion criteria are described in Appendix 1, available at www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.131431/-/DC1.

Screening

Participants underwent a standardized clinical interview and data collection at baseline (Figure 1). This phase served to establish the diagnosis and confirm eligibility for all participants.

Clinical assessment

All participants underwent grading according to the Expanded Disability Status Scale.³³ This grading was performed by a neurologist (F.C., W.J.D. or J. Mah) blinded to the imaging results.

Ultrasound procedures

Certified technologists³⁴ blinded to the clinical status of participants performed the ultrasonography,

using a Phillips IU-22 unit, with a linear-array, broad-bandwidth 9–3 MHz transducer on the manufacturer's setting for "upper extremity venous." Technical parameters were optimized for accurate results. Scanning was performed with participants in the supine and then sitting positions. Participants were instructed to breathe normally and to avoid performing a Valsalva manoeuvre.

The internal jugular and vertebral veins were assessed with grey-scale (B-mode) imaging in both longitudinal and transverse planes. Standardized measurements of each vein included vessel diameter and cross-sectional area. Grey-scale analysis of the internal jugular valve was performed in 2 planes to look for anomalies. When valve identification was challenging with the linear-array probe, a curvilinear-array probe with a small footprint was used.

We assessed venous patency using colour Doppler sonography. Pulsed-wave analysis documented flow direction, velocity and volume. We considered spectral and colour Doppler sonography results to be abnormal if flow appeared to be accelerated or if the normal variability of venous waveforms was lost. Colour Doppler sonography in the longitudinal plane was performed in the lower internal segment of the jugular vein, just above the level of the valve. If reflux was detected, a pulsed-wave spectral tracing was performed to measure duration. The cross-sectional area was viewed in grey scale in the transverse plane, with

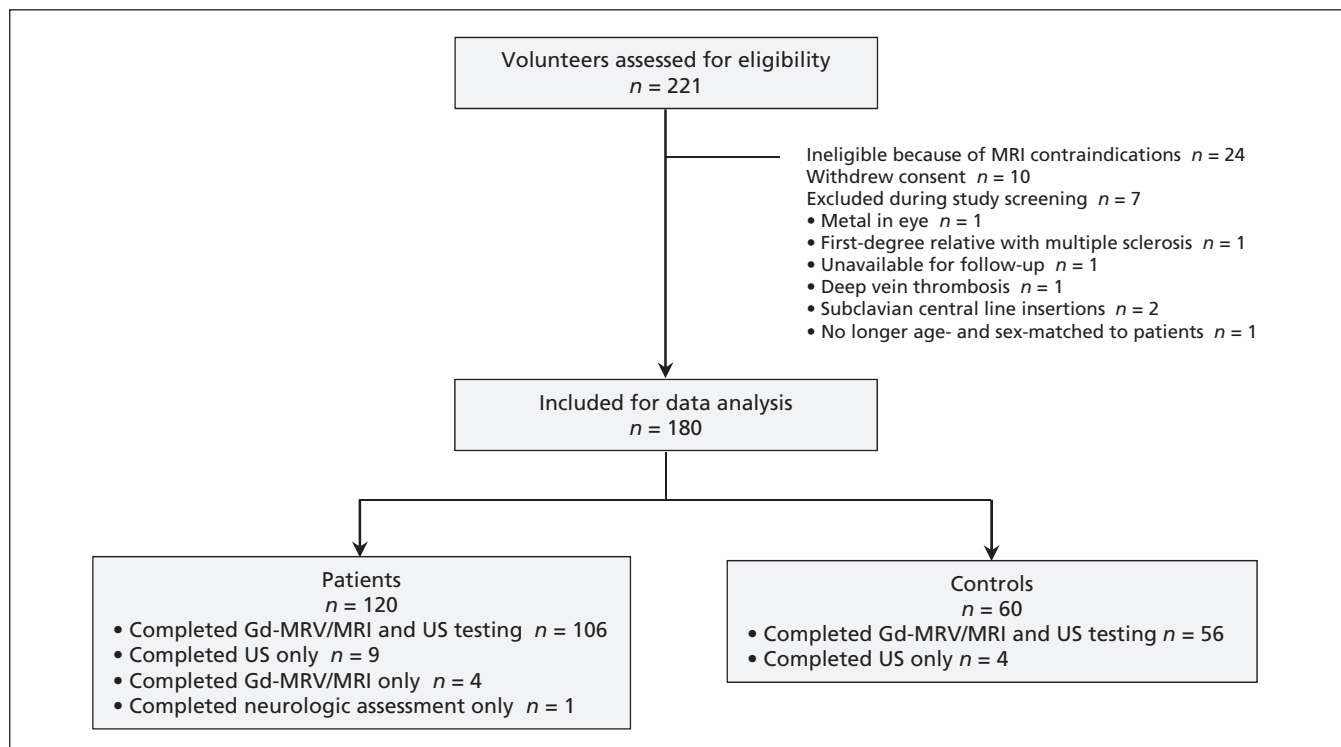


Figure 1: Enrolment of patients with multiple sclerosis and healthy controls, reasons for exclusions and types of data obtained. Gd = gadolinium, MRI = magnetic resonance imaging, MRV = magnetic resonance venography, US = ultrasonography.

measurement at the superior, inferior and mid sections of the internal jugular veins, with the participant in the supine and sitting positions.

Ultrasound criteria

Two radiologists (D.L, D.B.), who were blinded to participants' clinical status and magnetic resonance imaging results, interpreted the ultrasound results. They determined the presence of obstruction of venous outflow using the following criteria proposed for diagnosis of chronic cerebrospinal venous insufficiency^{3,4} (excluding reflux in deep cerebral veins, because we did not perform transcranial ultrasonography):

1. Reflux > 0.88 seconds in the internal jugular and vertebral veins (sitting or supine)
2. B-mode evidence of stenosis of the internal jugular vein, defined as local reduction of cross-sectional area < 50% or $\leq 0.3 \text{ cm}^2$ in the supine position
3. Flow not Doppler-detectable in the internal jugular and vertebral veins
4. Reverted postural control of the main cerebral venous outflow

Magnetic resonance venography: procedures and criteria

We performed gadolinium-enhanced magnetic resonance venography according to the procedures described in Appendix 1 (available at www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.131431/-/DC1). At the start of this study, there were no established criteria to diagnose chronic cerebrospinal venous insufficiency by this method. Consequently, we evaluated extracranial venous outflow and burden of disease according to the criteria presented in Appendix 1. Three radiologists (J. Modi, J.N.S. or M.G.), blinded to participants' clinical status and ultrasound results, interpreted these results.

Statistical analyses

The primary objective of the study was to determine the proportion of patients and controls with ultrasound and magnetic resonance venography evidence of venous outflow obstruction. Secondary outcomes included magnetic resonance imaging measures of brain inflammation and Expanded Disability Status Scale scores. The selection of 120 patients and 60 controls was a sample of convenience because the rate of venous outflow obstruction in patients versus controls in a Canadian population is unknown. The primary analysis compared the number of patients with venous outflow obstruction as determined by ultrasound and gadolinium-enhanced magnetic resonance venography versus controls with McNemar's test.

Results

Demographic characteristics

After screening, 180 participants were included in the study (Figure 1). Among the 120 patients, multiple sclerosis had been present for a median duration of 10.5 years and was predominantly of the relapsing–remitting subtype (86 patients [72%]) (Table 1).

Ultrasound results

Evaluations of reflux, cross-sectional area and velocity of internal jugular venous flow illustrated variable measurements that did not differ between groups (Appendix 2, available at www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.131431/-/DC1). A high proportion of patients (67/115 [58%]) and controls (38/60 [63%]) met 1 or more of the proposed criteria for the diagnosis of chronic cerebrospinal venous insufficiency,⁴ with no differences between groups ($p = 0.6$) (Table 2). A minority of

Table 1: Characteristics of study participants

Characteristic	No. (%) of participants*	
	Patients <i>n</i> = 120	Controls <i>n</i> = 60
Demographic		
Age, yr, mean \pm SD	45.7 \pm 11.9	45.3 \pm 11.2
Sex, female	89 (74)	45 (75)
Ethnic origin, white	110 (92)	56 (93)
Clinical factors		
Family history of MS	18 (15)	NA
Duration of MS, yr, median (IQR)	10.5 (5–18)	NA
EDSS, median (IQR)	2.25 (1.5–3.5)	0 (0)
MS subtype		
Relapsing–remitting	86 (72)	NA
Secondary progressive	17 (14)	NA
Primary progressive	12 (10)	NA
Clinically isolated syndrome	4 (3)	NA
Neuromyelitis optica	1 (1)	NA
Current disease-modifying treatment		
None	58 (48)	NA
Glatiramer acetate (Copaxone)	42 (35)	NA
Interferon β 1a (Rebif)	12 (10)	NA
Interferon β 1a (Avonex)	3 (2)	NA
Interferon β 1b (Betaseron)	3 (2)	NA
Fingolimod (Gilenya)	2 (2)	NA

Note: EDSS = Expanded Disability Status Scale (where steps 1.0–4.5 refer to patients with MS who are fully ambulatory, steps 5.0–9.5 are defined by impairment in neurologic function with progressive disability corresponding to higher numbers on the scale, and step 10.0 refers to death due to MS)³³; IQR = interquartile range, MS = multiple sclerosis, NA = not applicable, SD = standard deviation.

*Except where indicated otherwise.

patients (23/115 [20%]) and controls (6/60 [10%]) fulfilled 2 or more diagnostic criteria[†] ($p = 0.1$) (Table 2). Notably, baseline characteristics did not differ for these participants relative to those who fulfilled fewer criteria (Appendix 3, available at www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.131431/-/DC1). When we analyzed the results for individual ultrasound criteria, we found no differences between patients and controls (Table 2).

Magnetic resonance venography results

Patients showed greater “lesional” burden than controls. Lesions were detected in controls but were not accompanied by brain atrophy or gadolinium enhancement suggestive of underlying demyelination (Table 3). Post hoc analysis showed that these nonspecific lesions were fewer in number (relative to lesions observed in patients), and they were considered age-appro-

Table 2: Ultrasound criteria for chronic cerebrospinal vascular insufficiency in patients with multiple sclerosis and healthy controls

Criterion*	No. (%) of participantst		p value	OR (95% CI)‡
	Patients n = 115	Controls n = 60		
Criterion 1: reflux (IJV, VV: right and left, upright and supine)			0.3	
None	68 (59)	30 (50)		
1 vein	25 (22)	22 (37)		
2 veins	11 (10)	5 (8)		
3 veins	9 (8)	3 (5)		
4 veins	2 (2)	0 (0)		
Criterion 3: IJV stenosis (right and left)			0.4	
Both normal	89 (77)	52 (87)		
1 abnormal	17 (15)	6 (10)		
2 abnormal	9 (8)	2 (3)		
Criterion 4: no Doppler-detectable flow (IJV, VV: right and left, upright and supine)			> 0.9	
Normal	113 (98)	59 (98)		
1 vein	2 (2)	1 (2)		
Criterion 5: negative CSA in IJV (right and left)			0.2	
Change in CSA > 0	94 (82)	54 (90)		
Change in CSA ≤ 0	21 (18)	6 (10)		
No. of ultrasound criteria met				
≥ 1	67 (58)	38 (63)	0.6	0.9 (0.4–1.6)
≥ 2	23 (20)	6 (10)	0.1	2.2 (0.8–7.2)
≥ 3	6 (5)	1 (2)	0.5	3.2 (0.3–151.8)
≥ 4	0 (0)	0 (0)	NA	NA
0	48 (42)	22 (37)	0.6	
Exactly 1	44 (38)	32 (53)	0.08	
Exactly 2	17 (15)	5 (8)	0.3	
Exactly 3	6 (5)	1 (2)	0.4	
Exactly 4	0 (0)	0 (0)	NA	

Note: CI = confidence interval, CSA = cross-sectional area, IJV = internal jugular vein, NA = not applicable, OR = odds ratio, VV = vertebral vein

*All criteria as presented by Zamboni and colleagues.^{3,4} Criterion 1 = reflux in the IJVs and/or VVs in sitting or supine position, defined as reflux > 0.88 s; criterion 3 = Doppler B-mode evidence of IJV stenosis, defined as local reduction of cross-sectional area < 50% and/or ≤ 0.3 cm² in supine position; criterion 4 = flow not Doppler-detectable in the IJVs and/or VVs; criterion 5, reverted postural control of the main cerebral venous outflow. For the current study, criterion 2 (transcranial Doppler assessment of internal cerebral veins) was not performed.

†Except where indicated otherwise.

‡Fisher's exact test for patients and controls meeting 1 or more US criteria, 2 or more US criteria, 3 or more US criteria, or 4 or more US criteria.

Table 3: Results of magnetic resonance imaging and gadolinium-enhanced venography

MR result	No. (%) of participants				p value
	Patients n = 110		Controls n = 56		
MR imaging					
No. of white matter lesions					
0	5 (5)		29 (52)		< 0.001
1–10	34 (31)		22 (39)		
11–20	33 (30)		5 (9)		
21–30	38 (35)		0 (0)		
Enhancing lesions	12 (11)		0 (0)		0.2
Parenchymal atrophy	29 (26)		0 (0)		< 0.001
Corpus callosum atrophy	25 (23)		0 (0)		< 0.001
MR venography (TRICKS)*					
	Right	Left	Right	Left	Right / Left
IJV dominance	29 (26)	12 (11)	12 (21)	3 (5)	0.3/0.3
IJV flattening C1–C3 level, %					
	n = 109	n = 109			
0–24	32 (29)	32 (29)	14 (25)	17 (30)	0.9/0.7
25–49	44 (40)	42 (39)	23 (41)	17 (30)	
50–74	10 (9)	11 (10)	5 (9)	6 (11)	
75–100	23 (21)	24 (22)	14 (25)	16 (29)	
IJV flattening C4–C5 level, %					
0–24	94 (86)	87 (80)	51 (91)	46 (82)	0.9/0.8
25–49	5 (5)	9 (8)	1 (2)	5 (9)	
50–74	3 (3)	1 (1)	1 (2)	1 (2)	
75–100	7 (6)	12 (11)	3 (5)	4 (7)	
IJV flattening C6–T1 level, %					
0–24	55 (50)	87 (80)	29 (52)	46 (82)	0.9/0.9
25–49	21 (19)	9 (8)	11 (20)	5 (9)	
50–74	8 (7)	1 (1)	2 (4)	1 (2)	
75–100	25 (23)	12 (11)	14 (25)	4 (7)	
Collateral vein score					
Normal	25 (23)	25 (23)	12 (21)	13 (23)	> 0.9/0.8
Mild	45 (41)	39 (36)	25 (45)	23 (41)	
Moderate	29 (27)	37 (34)	14 (25)	15 (27)	
Prominent	10 (9)	8 (7)	5 (9)	5 (9)	
Intracranial venous stenosis	0 (0)		0 (0)		NA
Transverse sinus dominance	23 (21)	12 (11)	13 (23)	4 (7)	0.4/0.8
MR pre/post Gd RAGE					
IJV narrowing† at C1–C3 level, %					
0–24	35 (32)	38 (35)	14 (25)	17 (30)	0.8/0.4
25–49	44 (40)	43 (39)	25 (45)	18 (32)	
50–74	9 (8)	11 (10)	4 (7)	6 (11)	
75–100	21 (19)	17 (16)	13 (23)	15 (27)	
IJV narrowing† at C4–C6 level, %					
0–24	103 (94)	98 (90)	55 (98)	49 (88)	0.8/0.9
25–49	5 (5)	7 (6)	1 (2)	5 (9)	
50–74	1 (1)	1 (1)	0 (0)	0 (0)	
75–100	0 (0)	3 (3)	0 (0)	2 (4)	
IJV narrowing† at C7–T1 level, %					
0–24	61 (56)	69 (63)	31 (55)	32 (57)	0.7/0.7
25–49	21 (19)	11 (10)	10 (18)	8 (14)	
50–74	8 (7)	8 (7)	2 (4)	3 (5)	
75–100	19 (17)	21 (19)	13 (23)	13 (23)	
Mass or lymph nodes	0 (0)		0 (0)		NA

Note: Gd = gadolinium, IJV = internal jugular vein, MR = magnetic resonance, NA = not applicable, RAGE = rapid gradient echo, TRICKS = time-resolved imaging of contrast kinetics.

*For one patient, some MR venography sequences were unreadable, so data were available for only 109 patients.

†Narrowing commonly by extrinsic compression from the transverse vertebral process.

appropriate for the controls.¹ There were no differences in measures of venous outflow between patients and controls (Table 3).

Interpretation

We observed no differences in the proportion of venous outflow abnormalities, as measured by ultrasonography or gadolinium-enhanced magnetic resonance venography, between patients with multiple sclerosis and healthy controls.

Previously, Zamboni and colleagues^{3,4} reported that diagnosis of chronic cerebrospinal venous insufficiency required fulfillment of at least 2 of 5 ultrasound criteria. They detected greater prevalence of several criteria among patients than among controls: reflux (71% v. 0%), B-mode evidence of internal jugular vein stenosis (37% v. 0%), absent flow detectable by Doppler ultrasonography in the internal jugular or vertebral veins (52% v. 3%) and reversed postural flow in the internal jugular vein (55% v. 11%).³ When these criteria were applied in the evaluation of 109 patients with multiple sclerosis and 177 controls, each patient was deemed to meet at least 2 criteria, whereas none of the control participants did so.⁴ The sensitivity, specificity, and positive and negative predictive values for the proposed criteria were each 100%.⁴ Numerous groups have since been unable to replicate these findings.^{5-8,13,16,19,21} Doepp and associates⁵ reported that no patients with multiple sclerosis and no controls fulfilled more than 1 ultrasound criterion for chronic cerebrospinal venous insufficiency. Baretto and colleagues²¹ noted that 30% of their study's participants fulfilled 1 criterion, 5% met 2 criteria, and none satisfied more than 2 criteria for the diagnosis. The diagnostic criteria proposed for chronic cerebrospinal venous insufficiency are overly inclusive and nonspecific. This would explain why the condition was detected in 56% of patients with multiple sclerosis and 23% of controls in a prior study.⁹ It also accounts for our observation that 10% of the normal population would apparently carry the diagnosis.

The lack of reproducibility of the original findings reported by Zamboni and colleagues^{3,4} may relate to “methodological flaws” and the “pathophysiologic implausibility” of the ultrasound features used to define chronic cerebrospinal venous insufficiency.²⁰ The definition of venous reflux used by Zamboni and colleagues has been challenged because spectral Doppler measurements of duration were not implemented.²¹ Furthermore, the 0.88-second threshold used to identify reflux was adopted from a study that examined internal jugular vein valve insufficiency during the Valsalva manoeuvre and was not based on validated methodology.³⁵ Our findings indicate that the phenomenon characterized as “reflux” in the proposed criteria consists of low-velocity retrograde flow, which is almost exclusively observed in the periphery of the inferior internal jugular vein, near its confluence with the valve or subclavian vein. This flow was frequently present at a length of less than 2 cm, was of small volume and did not reach a height above the thyroid cartilage in any participant. We observed no evidence of large-volume reflux that would have approached the intracranial veins. Therefore, the phenomenon termed “reflux” was trivial in our observations and likely represented an eddy current near recesses associated with the valves of the inferior jugular vein (Video 1). Because this finding is nonpathologic, it is not surprising that “reflux” was frequently detected in both controls (50%) and patients with multiple sclerosis (41%).

In the proposed criteria,^{3,4} venous “stenosis” was defined as a 50% reduction in the cross-sectional area of the internal jugular vein or a value less than or equal to 0.3 cm².^{3,4,20} Both these measures have subsequently been refuted on the basis of methodologic and anatomic considerations.^{20,21} The internal jugular vein is wider at its point of origin in the upper bulb and its confluence into the subclavian vein in the lower bulb.²⁰ The cross-sectional area of the medium sections can be readily imaged using B-mode ultrasonography and varies with posture and central venous pressure.²⁰ Given this inherent variability, defining internal jugular vein stenosis as a 50% reduction is arbitrary.²⁰ Similarly, the 0.3-cm² threshold was chosen on the basis of findings published by Lichtenstein and associates.³⁶ Yet, in that same report, 23% of participants had a cross-sectional area less than or equal to 0.4 cm², and measurements less than or equal to 0.3 cm² were observed without associated pathology.^{20,36} Identifying a 50% reduction in the cross-sectional area as evidence of internal jugular vein stenosis may lead to false-positive results, because the venous wall is thin and can easily be compressed, either



Please see the following video online at: www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.131431/-/DC1.

Video 1 is an ultrasound video clip of the right internal jugular vein in the sagittal plane obtained with colour flow Doppler in a control participant. The video shows antegrade normal forward venous flow (toward the heart), which appears blue and phasic in the vein lumen. There is an intermittent small amount of peripheral red reversed flow against the deep vein wall inferiorly. This observation reflects only a very small volume of reversed blood flow, and the blood travels only a short distance (less than 1–2 cm) from the valve in the internal jugular vein in the root of the neck near the clavicle. This trivial reversed flow is brief, occurs a fair distance (about 15 cm) from the brain and was seen commonly in control participants (normal volunteers, without neurologic disease).

manually or by surrounding anatomic structures (Figure 2). The internal jugular vein is dilated at the craniocervical junction and distally, and it may appear narrowed within the region of the valves. We observed this phenomenon in our study, in which the overwhelming number of veins deemed to be “stenotic”^{3,4} had an appearance suggesting a smoothly collapsed vein rather than a focal stricture. We did not observe focal vein narrowing with upstream dilatation, accompanied by a focal intraluminal source of accelerated flow and collateral formation, in any participant. The “stenoses” detected in our study would not have been identified as such by qualified ultrasonographers. This impression is supported by the fact that gadolinium-enhanced magnetic resonance venography showed no evidence of impaired cerebral venous drainage in any of our participants, despite the fact that 34 “stenotic” internal jugular veins were found with the proposed ultrasound criteria (Table 2).

We rarely observed “flow not Doppler-detectable” in the internal jugular or vertebral veins of patients or controls. In the original work of Zamboni and colleagues, Doppler-detected flow was assessed “either with the pulse wave mode and the sample placed in the vessel at a 60 degree angle, or with the color coded mode.”⁴ Veins were examined using “both the transversal and/or the longitudinal cervical access.”⁴ These methods suggest insufficient attention to Doppler sonography techniques and physics. The presence of a cosine in the Doppler formula indicates that any time a structure is interrogated at an angle approaching 90°, the signal will diminish toward zero. Therefore, Doppler evaluation of a vessel in the transverse plane is apt to create a false-positive interpretation of absent flow. Several factors may have resulted in the reported absence of Doppler-detected flow in earlier pub-

lications, including scanning without angle correction, failing to employ Doppler in a manner that kept the plane of imaging as parallel as possible to the direction of flow, and poor optimization of scale, gain and velocity settings.

Limitations

Perhaps the most striking finding in our study was the range of venous outflow anomalies detected that did not reflect pathology, but instead demonstrated the large natural variance in intracranial and extracranial venous anatomy (Figure 3).³⁷ Venous capacitance vessels change appearance and flow depending upon patient position, intravascular volume status and ultrasonography technique.²⁰ Given that venous walls are collapsible under conditions of reduced flow, the complexity of the extracranial venous circulation may be difficult to understand in any but a qualitative fashion.³⁷ By extension, there are challenges in standardizing ultrasound examinations of the cerebral venous system. Our data indicate that venous peak velocity and cross-sectional area vary greatly with changes in posture.

We were unable to control for intravascular volume status, because invasive measurements of central venous pressure were not possible in this study. Many patients with multiple sclerosis manifest symptoms of bladder (detrusor) sphincter dyssynergia and may be reluctant to drink fluids, which may cause them to be relatively hypovolemic compared with controls. The effect of hydration on ultrasound findings in this patient population remains unclear.

In our study, magnetic resonance imaging was conducted with patients in the supine position only, because technology allowing upright scanning was not available. For this reason, the effects of posture on measures of venous patency could not be determined. Importantly, we noted

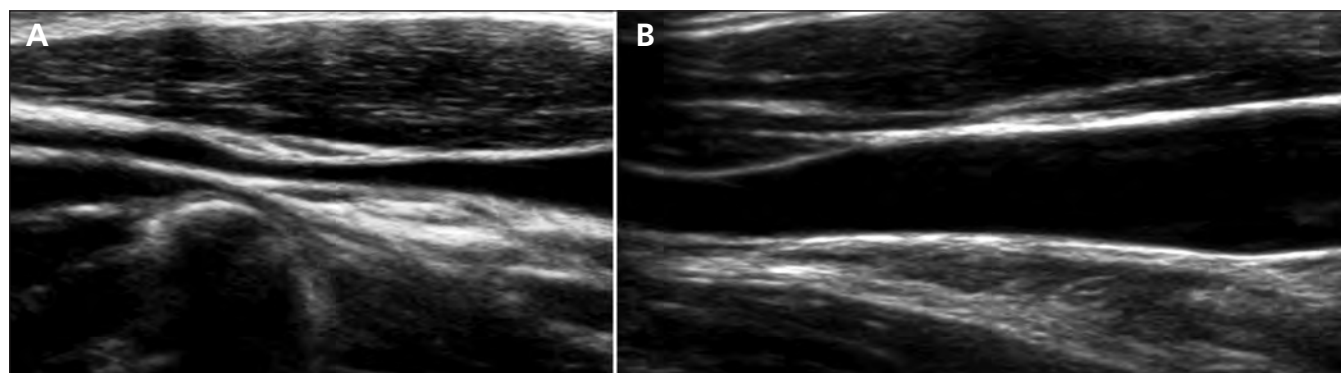


Figure 2: (A) Ultrasound image of the right internal jugular vein (sagittal plane), obtained with the participant lying supine. There is narrowing of the internal jugular vein that would be interpreted as abnormal according to the proposed ultrasound criteria for diagnosis of chronic cerebrospinal venous insufficiency.^{3,4} (B) Ultrasound image of the same vein obtained a few minutes later, from the same participant in the same position, shows that the vein is normal and does not have fixed narrowing. Observation of narrowing in Fig. 2A may have been due to compression by the transducer or may have been secondary to a change in central venous pressure (e.g., from a change in phase of respiration or in cardiac output).

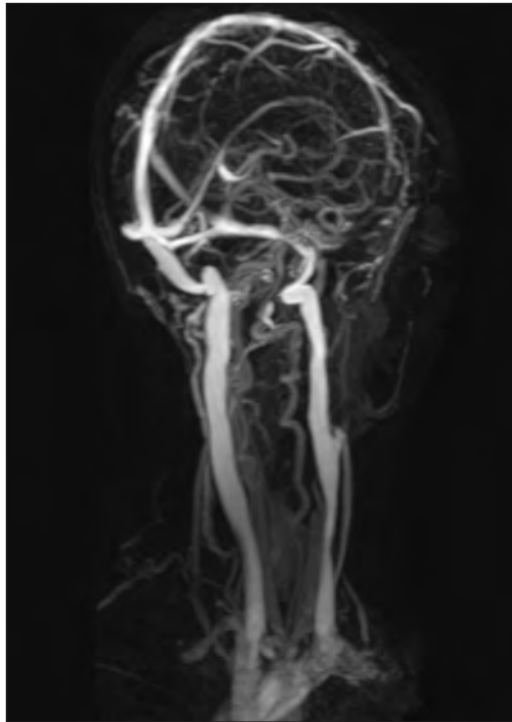


Figure 3: Gadolinium-enhanced magnetic resonance venography of a participant with 25% stenosis of the left internal jugular vein (arrow).

that the internal jugular vein is commonly partly compressed (25% narrowing) in its extracranial course (Figure 3), and there is a tendency toward right-sided dominance for venous drainage.

Our study had several strengths, including its prospective design, complementary magnetic resonance imaging protocols and blinding strategies. Yet interpretation of our findings may be limited by failure to control for participant intravascular volume status, head position, anatomic variation of the neck strap muscles and sample size.

Conclusion

We detected no link between chronic cerebrospinal venous insufficiency and multiple sclerosis. We also identified several methodologic concerns that challenge the validity of the criteria used to define chronic cerebrospinal venous insufficiency, and in turn we dispute the authenticity of this diagnosis.

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