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Jugular Venous Reflux and White Matter Abnormalities in Alzheimer's Disease: A Pilot Study

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Abstract. To determine whether jugular venous reflux (JVR) is associated with cerebral white matter changes (WMCs) in individuals with Alzheimer's disease (AD), we studied 12 AD patients 24 mild cognitive impairment (MCI) patients, and 17 elderly age- and gender-matched controls. Duplex ultrasonography and 1.5T MRI scanning was applied to quantify cerebral WMCs [T2 white matter (WM) lesion and dirty-appearing-white-matter (DAWM)]. Subjects with severe JVR had more frequently hypertension ($p=0.044$), more severe WMC, including increased total ($p=0.047$) and periventricular DAWM volumes ($p=0.008$), and a trend for increased cerebrospinal fluid volumes ($p=0.067$) compared with the other groups. A significantly decreased (65.8%) periventricular DAWM volume ($p=0.01$) in the JVR-positive AD individuals compared with their JVR-negative counterparts was detected. There was a trend for increased periventricular and subcortical T2 WMC lesion volumes in the JVR-positive AD individuals compared with their JVR-negative counterparts ($p=0.073$). This phenomenon was not observed in either the control or MCI groups. In multiple regression analysis, the increased periventricular WMC lesion volume and decreased DAWM volume resulted in 85.7% sensitivity and 80% specificity for distinguishing between JVR-positive and JVR-negative AD patients. These JVR-WMC association patterns were not seen in the control and MCI groups. Therefore, this pilot study suggests that there may be an association between JVR and WMCs in AD patients, implying that cerebral venous outflow impairment might play a role in the dynamics of WMCs formation in AD patients, particularly in the periventricular regions. Further longitudinal studies are needed to confirm and validate our findings.

Keywords: Alzheimer's disease, Doppler ultrasonography, jugular veins, leukoaraiosis, magnetic resonance imaging

INTRODUCTION

White matter changes (WMCs) with punctate and/or confluent WM lesions in the periventricular and/or

deep subcortical regions, revealed as hyperintensities on T2-weighted or fluid-attenuated inversion recovery (FLAIR) magnetic resonance imaging (MRI), are strongly related to aging [1]. These age-related WMCs are associated with occurrences of different types of dementia including Alzheimer's disease (AD) and predict faster cognitive decline not only in the general population, but also in AD patients [2, 3].

Jugular venous reflux (JVR), characterized by a retrograde flow in internal jugular veins (IJVs) during

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Valsalva maneuvers (VM) or spontaneously at rest, is found more frequently in the elderly [4–6]. Recently, it has been found that JVR is associated with age-related WMCs, particularly in posterior brain regions [7]. Retrograde-transmitted venous pressure associated with JVR in the elderly is thought to result in accumulated insults to the cerebral vasculature, small vessel structure/function, and/or the brain-blood barrier, leading to WM damage [7, 8]. Compared with non-demented individuals, WMCs are more prevalent and severe in AD patients [9–14]. In this study, we tested the hypothesis that JVR might be involved in the dynamics of WMC formation in AD by evaluating the relationship between the severity of JVR and WMC in patients with AD.

MATERIAL AND METHODS

Subjects

Between December 2008 and April 2010, Taiwanese residents consecutively admitted to a memory clinic at Taipei Veterans General Hospital, Taiwan due to subjective memory complaints were assessed for inclusion. Neurologists performed clinical and neurologic evaluations in all participants. A standard set of neuropsychological assessments including the Mini-Mental State Examination (MMSE) and Clinical Dementia Rating (CDR) scales were used.

Subjects eligible for participation in the current study were 55 years of age or older, had a CDR score ≤ 1 (as an assurance that they could cooperate during the VM for JVR detection), and were willing to receive brain MRI and neck duplex ultrasonography. Exclusion criteria for all subjects were a past history of stroke, ischemic heart disease, congestive heart disease, valvular heart disease, cardiac arrhythmia, pulmonary diseases, or malignancy, and having brain MRI of insufficient quality for performing quantitative volumetric analysis.

Vascular risk factors were defined according to international guidelines and prospectively identified using all available information including medical charts, laboratory results, patient interviews, and neurological examinations (Supplementary Material) [15].

Subjects were classified into AD, mild cognitive impairment (MCI), or control groups according to the criteria of National Institute of Neurological and Communicative Disorders and Stroke/Alzheimer's Disease and Related Disorders Association and by Petersen et al.'s study revised by the Stockholm consensus group [16–18]. The hospital's Institutional Review Boards

approved the study, and each included participant or his/her caregiver provided informed consent.

Color-coded duplex ultrasonography: JVR determination

Ultrasonographic determination of JVR has been described previously (Supplementary Materials) [6, 7]. JVR could be detected spontaneously at rest or only during VM. We graded the severity of JVR into three groups. Subjects with no JVR on both IJV sides were classified as a no-JVR group. Subjects with JVR detected on either or both IJV sides during VM only, or subjects with spontaneous JVR on one IJV side and no JVR on the other IJV side were graded as the mild-JVR group. Subjects with JVR detected on one IJV side spontaneously at rest in combination with JVR detected during VM or spontaneously at rest in the other side were graded as the severe-JVR group [7].

MRI acquisition

All study participants were scanned on 1.5T MRI (Excite II; GE Medical Systems, Milwaukee, WI). MRI brain sequences included axial two dimensional (2D) T1- and T2-weighted imaging (WI), FLAIR and three dimensional (3D) T1 spoiled gradient echo (SPGR) pulse sequences. The images were acquired with the following sequence parameters: for 3D T1 the repetition time (TR) was 8.5 ms, echo time (TE) 1.8 ms, inversion time (TI) 400 ms, flip angle (FA) 15° , slice thickness 1.5 mm, field of view (FOV) 260 mm \times 260 mm and matrix 256 \times 256; for T2 the TR was 3700 ms, TE 102 ms, echo train length 19, slice thickness 3 mm, FOV 240 mm \times 240 mm and matrix 256 \times 256; for FLAIR: TR 9000 ms, TE 120 ms, TI 2250 ms, slice thickness 3 mm, FOV 240 mm \times 240 mm and matrix 256 \times 256.

MRI image analysis

The MRI analysis personnel were blinded to the subjects' demographic and clinical characteristics.

We classified WMCs as belonging to two groups of signal abnormalities based on FLAIR/T2-WI signal intensity: T2 WM lesion and dirty-appearing-whitematter (DAWM). The T2 WM lesion number and volume on T2-WI were outlined using a semi-automated edge detection contouring/thresholding technique as described previously [19]. Regional localization of WM lesion volume was determined in the periventricular and deep subcortical regions. We also

outlined areas of DAWM and calculated their number and volume. The DAWM was defined as a non-focal area of signal increase on the FLAIR/T2-WI, with a subtly increased signal intensity compared with the signal intensity of normal appearing WM, as previously proposed [20]. The DAWM showed relatively diffuse border compared with focal T2 WM lesions and was tapered off toward the normal appearing WM (Fig. 1). The regional localization of DAWM volume was also determined in the periventricular and deep subcortical regions. The data are presented separately for number and volume of total and regional T2 WM lesions as well as DAWM.

In addition, we used Structural Image Evaluation, using Normalisation, of Atrophy, cross-sectional (SIENAX, version 2.6) to estimate normalized lateral ventricular volume (NLVV), and normalized cerebrospinal fluid (CSF) volume, as previously described [21].

Statistical analysis

Statistical analysis was undertaken using a combination of Statistical Package for the Social Sciences (SPSS, IBM, Armonk, New York, USA) and in-house algorithms written in Matlab (Mathworks, Natick, Mass) with the aim of evaluating the impact of JVR on the respective MRI variables.

Parametric and non-parametric analyses were performed on the respective study cohorts to identify significant differences between the various study groups, including disease groups, JVR severity score, and JVR presence. In order to identify MRI variables that were associated with the presence of JVR, we obtained plots of the logit versus logistic function of the multiple regression analysis in the respective study groups.

Given the exploratory pilot nature of the study, values of $p < 0.05$ were considered statistically significant and $p < 0.1$ was considered a trend.

RESULTS

Eighty-four subjects [57–93 years of age; mean(SD): 79.77(7.41) years; 33 women; 31 control subjects, 33 MCI subjects, and 20 AD subjects] were eligibly enrolled according to our criteria. To match age and gender in each disease group, 53 subjects [64–91 years of age; mean(SD): 81.5 (4.50) years; 22 women] were enrolled for further analysis. There were 17 control subjects, 24 MCI subjects, and 12 AD subjects (Table 1). All the AD patients met the

diagnostic criteria for AD and CDR = 1; and all MCI patients met the diagnostic criteria for MCI and had a CDR = 0.5. Twenty two subjects (41.5%) had right-sided JVR (all detected during VM) and 23 subjects (43.4%) had left-sided JVR (9 detected spontaneously at rest and 14 detected during VM, Table 2). There were 12 subjects (22.6%) with bilateral JVR; among them, 5 had unilateral spontaneous JVR at rest with contralateral VM-induced JVR and 7 had bilateral VM-induced JVR. Twenty subjects (37.7%) presented with JVR on neither side.

Table 1 shows the comparisons of clinical characteristics and MRI variables between control, MCI, and AD groups. From this it can be seen that for all but three study variables, there was no significant difference between the respective groups. The only exceptions to this were: the MMSE score, which was significantly lower in the AD group ($p < 0.001$); the number of years in education, which was on average approximately 3 years less in the MCI and AD groups ($p = 0.033$); and hyperlipidemia, which had a higher incidence in the AD group ($p = 0.014$).

The clinical characteristics and MRI variables between the three JVR-graded groups are presented in Table 2, which also shows the frequencies of vascular risk factors. The three JVR-graded groups were closely age-matched and had similar clinical characteristics, with the exception of the incidence of hypertension, which was significantly greater in the mild and severe JVR groups ($p = 0.044$). There was a general trend for an increased WMC number and volume in the severe-JVR group that reached significance for the total DAWM ($p = 0.047$) and periventricular DAWM volume ($p = 0.008$). The more severe JVR group also exhibited a trend for increased CSF volumes compared with the other groups ($p = 0.067$).

Further investigation, using a 2-tailed Mann Whitney U-test (Table 3), revealed a significantly decreased (65.8%) periventricular DAWM volume ($p = 0.01$) in the JVR-positive AD individuals compared with their JVR-negative counterparts. There was also a trend for an increased periventricular and subcortical T2 WMC lesion volume in the JVR-positive AD individuals compared with their JVR-negative counterparts ($p = 0.073$). This phenomenon was not observed in the either the control or MCI groups.

Figure 2 shows a plot of the logit versus logistic function of the multiple regression analysis for the AD patients group using the periventricular T2 WMC volume ($B = 0.0004$) and periventricular DAWM volume ($B = -0.0039$) versus the JVR positivity score. This resulted in sensitivity of 85.7% and specificity

Table 1
Clinical characteristics and MRI variables between control, MCI, and AD groups

	Control (n = 17)	MCI (n = 24)	AD (n = 12)	p
Age, years, mean (SD)	81.4 (3.8)	81.4 (4.0)	81.8 (6.4)	0.968
Females, n (%)	5 (29.4)	11 (45.8)	6 (50.0)	0.457 ^a
Education, years, mean(SD)	13.4 (1.8)	10.9 (4.4)	9.8 (4.1)	0.033
MMSE	28.1 (1.6)	26.2 (1.9)	20.4 (3.1)	<0.001
Hypertension, n (%)	11 (64.7)	12 (50.0)	7 (58.3)	0.639 ^a
Diabetes mellitus, n (%)	4 (23.5)	5 (20.8)	3 (25.0)	0.956 ^a
Hyperlipidemia, n (%)	3 (17.6)	2 (8.3)	6 (50.0)	0.014 ^a
Smoking, n (%)	1 (5.9)	1 (4.2)	0 (0.0)	0.709 ^a
JVR				
Positive ^c , n (%)	11 (64.7)	15 (62.5)	7 (58.3)	0.941 ^a
Severity score ^d , mean (SD)	1.8 (0.7)	1.9 (0.8)	1.8 (0.8)	0.864 ^b
WMC number changes, mean (SD)				
T2 WM lesions + DAWM	28.1 (16.6)	44.9 (30.6)	42.6 (18.6)	0.091
T2 WM lesions	25.4 (17.3)	25.3 (16.0)	24.8 (19.9)	0.996
DAWM	18.2 (21.2)	33.4 (38.5)	31.3 (11.6)	0.246
WMC volume changes, mean(SD)				
T2 WM lesions + DAWM	9.5 (12.8)	11.9 (14.7)	10.6 (16.4)	0.875
T2 WM lesions	8.6 (12.8)	10.6 (14.7)	9.5 (16.3)	0.909
Subcortical	2.5 (5.6)	2.2 (3.4)	2.2 (4.2)	0.975
Periventricular	6.1 (7.8)	8.4 (13.8)	7.4 (12.1)	0.828
DAWM	0.9 (0.9)	1.3 (1.1)	1.1 (0.6)	0.483
Subcortical	0.5 (0.6)	0.7 (0.7)	0.4 (0.3)	0.252
Periventricular	0.4 (0.4)	0.6 (0.7)	0.7 (0.5)	0.504
Normalized LV volume	77.4 (27.0)	79.5 (24.8)	91.8 (25.3)	0.294
CSF volume	374.5 (54.4)	364.0 (63.8)	367.0 (63.4)	0.860

JVR, jugular vein reflux; n, number; SD, standard deviation; %, percentage; WM, white matter; WMC, white matter changes; MCI, mild cognitive impairment; AD, Alzheimer's disease; DAWM, dirty-appearing white matter; LV, lateral ventricle; CSF, cerebrospinal fluid. *p* value determined by one-way ANOVA unless otherwise stated. ^a*p* value determined using chi square test. ^b*p* value determined using Kruskal-Wallis test. ^cThe presence of JVR in either side would be defined as JVR-positive. ^dJVR severity score: 1 = no JVR; 2 = mild JVR; 3 = severe JVR.

236 of 80% for distinguishing between JVR-positive and
237 JVR-negative AD patients. Similar analysis revealed
238 that none of the MRI variables were significantly asso-
239 ciated with JVR positivity score in the control and MCI
240 groups.

241 DISCUSSION

242 The present pilot study was of limited scope, having
243 a small sample size with only few subjects who pre-
244 sented with severe JVR. Therefore, our findings should
245 be interpreted with caution and no firm conclusions
246 should be made regarding the general applicability
247 of our findings at this moment in time. Future stud-
248 ies including larger sample size and more MCI and
249 AD patients are needed to confirm the validity of our
250 findings.

251 The principal finding of the present study is that JVR
252 appears to be associated with changes in the DAWM,
253 particularly in the periventricular region. Increased
254 DAWM volume, particularly in the periventricular
255 region, was associated with a more severe JVR score.
256 DAWM is defined as a region of intermediate signal

257 intensity between that of T2 WM lesions and that
258 of normal-appearing white-matter [20, 23–27]. The
259 measurement of DAWM is relatively unexplored terri-
260 tory in neurodegenerative disorders and aging. To the
261 best of our knowledge, the present study is the first
262 of its kind to explore measurement of the DAWM in
263 elderly. Most of the work on DAWM has been per-
264 formed in patients with multiple sclerosis [20, 23–27],
265 and although the concept of DAWM remains largely
266 under investigation, it appears to have potential as a
267 diagnostic metric. Image-pathology correlation stud-
268 ies reveal DAWM, compared with T2 WM lesions, to
269 be associated with mild blood-brain barrier breakdown
270 and myelin loss [20, 23–27]. Therefore, it can be sug-
271 gested that DAWM may represent an earlier stage of
272 WM pathology in elderly [25, 26], which is followed
273 at a later stage, by T2 WM lesion formation [35–38].
274 Indeed, using WMC lesion volumes and DAWM it
275 was possible to distinguish between the JVR-positive
276 and JVR-negative AD patients, with high degree of
277 accuracy (sensitivity = 85.7% and specificity = 80%)
278 and minimal overlap (Fig. 2). These findings sug-
279 gest that in the AD group, DAWM areas may have
280 converted into T2 WM lesions at a greater rate in

Table 2
Clinical characteristics and MRI variables between three JVR-graded groups

	No JVR (<i>n</i> = 20)	Mild JVR (<i>n</i> = 28)	Severe JVR (<i>n</i> = 5)	<i>p</i>
Age, years, mean (SD)	81.1 (3.4)	81.7 (5.0)	81.6 (6.3)	0.882
Females, <i>n</i> (%)	8 (40.0)	13 (46.4)	1 (20.0)	0.535 ^a
Disease classification, <i>n</i> (%)				0.864 ^b
Control	6 (30.0)	10 (35.7)	1 (20.0)	
MCI	9 (45.0)	12 (42.9)	3 (60.0)	
AD	5 (25.0)	6 (21.4)	1 (20.0)	
Education, years, mean (SD)	12.3 (4.0)	10.9 (3.8)	11.2 (4.2)	0.496
MMSE	25.1 (4.0)	25.9 (3.1)	24.8 (4.7)	0.705
Hypertension, <i>n</i> (%)	8 (40.0)	17 (60.7)	5 (100.0)	0.044 ^a
Diabetes mellitus, <i>n</i> (%)	7 (35.0)	5 (17.9)	0 (0.0)	0.168 ^a
Hyperlipidemia, <i>n</i> (%)	4 (20.0)	6 (21.4)	1 (20.0)	0.992 ^a
Smoking, <i>n</i> (%)	0 (0.0)	2 (7.1)	0 (0.0)	0.395 ^a
WMC number changes, mean (SD)				
T2 WM lesions + DAWM	35.4 (20.8)	38.1 (26.9)	58.0 (26.6)	0.193
T2 WM lesions	23.1 (15.9)	25.7 (19.1)	31.0 (3.9)	0.642
DAWM	28.6 (27.4)	24.3 (32.2)	46.6 (13.6)	0.301
WMC volume changes, mean (SD)				
T2 WM lesions + DAWM	7.3 (8.9)	11.3 (15.2)	21.9 (22.2)	0.119
T2 WM lesions	6.2 (8.9)	10.4 (15.2)	19.8 (22.6)	0.150
Subcortical	1.5 (3.2)	2.6 (4.7)	4.1 (6.0)	0.441
Periventricular	4.7 (7.4)	7.9 (12.6)	15.7 (17.0)	0.159
DAWM	1.1 (0.8)	0.9 (1.0)	2.1 (1.0)	0.047
Subcortical	0.5 (0.4)	0.6 (0.7)	1.0 (0.3)	0.296
Periventricular	0.7 (0.5)	0.4 (0.4)	1.1 (1.0)	0.008
Normalized LV volume	81.6 (25.9)	77.8 (24.4)	102.8 (27.4)	0.136
CSF volume	373.0 (53.0)	355.1 (61.6)	420.7 (53.3)	0.067

JVR, jugular vein reflux; *n*, number; SD, standard deviation; %, percentage; WM, white matter; WMC, white matter changes; MCI, mild cognitive impairment; AD, Alzheimer's disease; DAWM, dirty-appearing white matter; LV, lateral ventricle; CSF, cerebrospinal fluid. Volumes are expressed in milliliters. *p* value determined by one-way ANOVA unless otherwise stated. ^a*p* value determined using chi square test. ^b*p* value determined using Kruskal-Wallis test.

281 subjects with more severe JVR (Table 3). However, 304
282 future longitudinal studies are necessary to elucidate 305
283 the chronological JVR effects on WMC formation in 306
284 AD patients.

285 The present study suggests that periventricular 307
286 WMCs may be associated with JVR (Tables 2 and 3). 308
287 This regional-association was not shown in our pre- 309
288 vious study [7], possibly because: (1) the volumetric 310
289 and DAWM approaches used in this study were more 311
290 sensitive to small group differences than the WMC 312
291 visual-rating scale used previously [20, 22–27]; or 313
292 (2) there were differences in the severe-JVR popula- 314
293 tions of the groups that were used in the two studies 315
294 [7]. Why the periventricular region in AD patients 316
295 could be influenced by JVR is something unknown 317
296 at this time. However, there is evidence that venous 318
297 drainage anomalies can influence CSF dynamics, slow- 319
298 ing down CSF absorption into the superior sagittal 320
299 sinus [39]. The lateral ventricles represent the major 321
300 reservoir of CSF in the brain. It has been proposed 322
301 that retrograde-transmitted venous pressure, caused by 323
302 JVR, may induce venous hypertension in the superior 324
303 sagittal sinus, resulting in decreased CSF absorption 325
326

and larger lateral ventricle volume [28–31] and CSF 304
volume, as evidenced in the present study. However, 305
this association has to be further investigated. 306

307 The etiology of WMC in AD is still uncertain. Our 308
309 pilot results suggest that DAWM may be a precursor 310
311 to T2 lesion formation and that transition between the 312
313 two might be accelerated by the presence of JVR. It 314
315 may be that JVR alters the intracranial biomechanics 316
317 interacting synergistically with other AD pathologies. 318
319 That these associations were not observed in either the 320
321 MCI subjects or controls may suggest that JVR may 322
323 play some role in late AD pathophysiology. It has been 324
325 reported that retrogradely transmitted venous hyper- 326
tension in AD patients associated with JVR might: (1) decrease cerebral perfusion, damage small vessel structure/function, and possibly damage the blood-brain barrier [8]; or (2) impair CSF absorption, thus reducing CSF net flow, enlarging ventricles, and possibly lowering amyloid- β clearance with the result that it accumulates in the periventricular tissues [32–34]. Either one or both of these mechanisms might contribute to WMC formation in AD patients. However, at this stage it is not possible to ascertain this, and

Table 3
MRI variables classified according to JVR status (i.e., positive or negative)

	JVR-negative	JVR-positive	<i>p</i>
Controls, <i>n</i> (%)	6 (30.0)	11 (70.0)	NA
WMC number changes, mean (SD)			
T2 WM lesions + DAWM	24.2 (16.7)	30.3 (17.0)	0.509
T2 WM lesions	21.5 (14.3)	27.5 (19.1)	0.713
DAWM	13.8 (10.3)	20.5 (25.4)	0.831
WMC volume changes, mean (SD)			
T2 WM lesions + DAWM	3.9 (3.4)	12.6 (15.1)	0.216
T2 WM lesions	3.3 (3.1)	11.5 (15.2)	0.256
Subcortical	0.7 (0.9)	3.5 (6.9)	0.538
Periventricular	2.6 (2.5)	8.0 (9.1)	0.216
DAWM	0.6 (0.4)	1.1 (1.1)	0.612
Subcortical	0.3 (0.2)	0.6 (0.7)	0.675
Periventricular	0.4 (0.3)	0.4 (0.4)	0.904
Normalized LV volume	69.0 (23.1)	82.0 (28.9)	0.525
CSF volume	348.0 (49.7)	389.0 (53.4)	0.216
MCI, <i>n</i> (%)	9 (45.0)	15 (55.0)	NA
WMC number changes, mean (SD)			
T2 WM lesions + DAWM	35.4 (19.9)	50.5 (35.0)	0.633
T2 WM lesions	26.3 (18.2)	24.7 (15.1)	0.743
DAWM	37.9 (37.6)	30.7 (40.0)	0.455
WMC volume changes, mean (SD)			
T2 WM lesions + DAWM	11.4 (12.0)	12.2 (16.4)	0.952
T2 WM lesions	10.1 (12.2)	10.9 (16.4)	1.000
Subcortical	2.7 (4.6)	2.0 (2.6)	0.905
Periventricular	7.4 (10.4)	9.0 (15.7)	0.858
DAWM	1.3 (1.0)	1.3 (1.2)	0.698
Subcortical	0.7 (0.6)	0.7 (0.8)	0.952
Periventricular	0.6 (0.6)	0.5 (0.8)	0.426
Normalized LV volume	85.8 (29.6)	75.6 (21.6)	0.474
CSF volume	377.3 (54.2)	356.0 (69.4)	0.512
AD, <i>n</i> (%)	5 (25.0)	7 (75.0)	NA
WMC number changes, mean (SD)			
T2 WM lesions + DAWM	48.8 (22.5)	38.1 (15.5)	0.505
T2 WM lesions	19.2 (15.5)	28.9 (22.7)	0.530
DAWM	29.4 (8.6)	32.6 (13.9)	0.861
WMC volume changes, mean (SD)			
T2 WM lesions + DAWM	4.2 (2.2)	15.2 (20.7)	0.202
T2 WM lesions	2.8 (1.8)	14.4 (20.4)	0.073
Subcortical	0.3 (0.4)	3.5 (5.2)	0.073
Periventricular	2.5 (1.6)	10.9 (15.2)	0.073
DAWM	1.4 (0.7)	0.8 (0.5)	0.106
Subcortical	0.3 (0.3)	0.4 (0.4)	0.755
Periventricular	1.1 (0.4)	0.4 (0.2)	0.010
Normalized LV volume	88.9 (20.6)	93.8 (29.7)	0.876
CSF volume	395.4 (53.0)	346.8 (66.2)	0.268

JVR, jugular vein reflux; *n*, number; SD, standard deviation; %, percentage; WM, white matter; WMC, white matter changes; MCI, mild cognitive impairment; AD, Alzheimer's disease; DAWM, dirty-appearing white matter; LV, lateral ventricle; CSF, cerebrospinal fluid; NA, not available. Volumes are expressed in milliliters. *p* values determined by 2-tailed Mann Whitney U-test.

327 further investigation is required before any causal link
328 between JVR in AD pathology can be claimed. Our
329 findings may serve as an important pilot data in this
330 direction.

331 It is noticeable that JVR appears to be associated
332 with certain cardiovascular risk factors, e.g., hyperten-
333 sion. Dolic et al. [40] studied 240 healthy individuals
334 and found that the presence of heart disease, especially

heart murmurs, obesity, and cigarette smoking were
associated with an increased prevalence of extracra-
nial venous abnormalities. These results, in addition
to the findings from the present study, may imply that
cardiovascular or systemic metabolic disorders may
be involved in the pathophysiology of venous (espe-
cially IJV) hemodynamic abnormalities. Besides, it is
known that patients with hypertension have less venous

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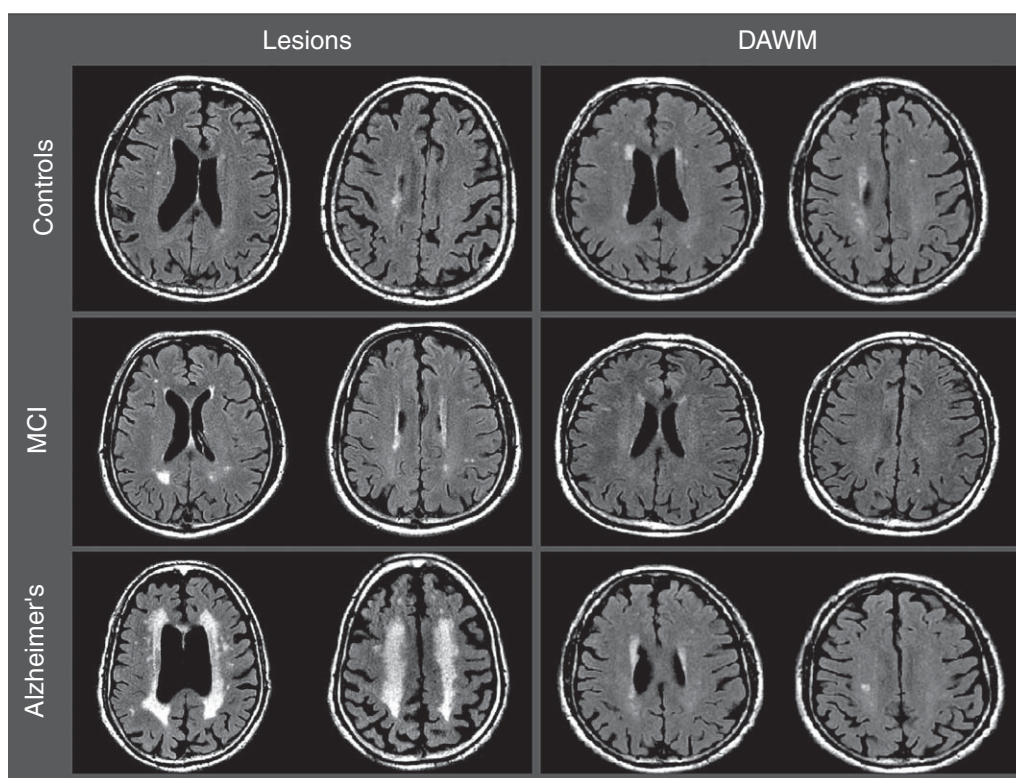


Fig. 1. Examples of periventricular and subcortical white matter T2 lesions and dirty appearing white matter (DAWM) in controls, mild cognitive impairment (MCI) patients, and Alzheimer's disease patients.

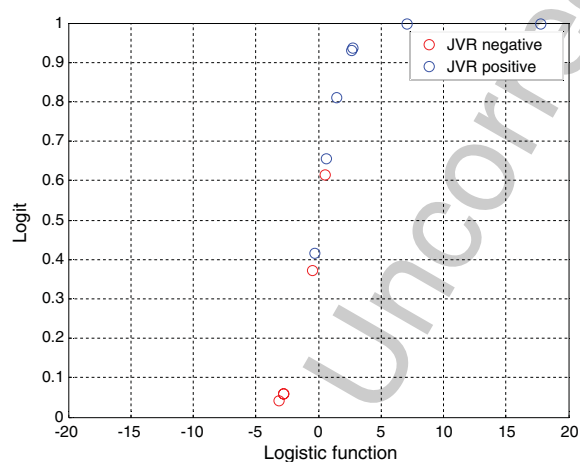


Fig. 2. Plot of the logit versus logistic function for the AD patients group using the variables periventricular T2 WMC volume ($B=0.0004$) and periventricular DAWM volume ($B=-0.0039$). Logit=0.5 is deemed to separate the JVR-positive subjects from the negative subjects. This results in one false positive and one false negative (sensitivity = 85.7% and specificity = 80%). JVR, jugular vein reflux; AD, Alzheimer's disease; WMC, white matter change; DAWM, dirty-appearing white matter.

distensibility [41, 42]. One characteristic specific for venous walls is that they have greater distensibility than arterial walls which makes them able to accommodate a greater range of increased venous pressure [43]. Decreased venous distensibility in hypertension might reduce this pressure buffer ability in IJV and repetitive higher venous pressure might damage IJV valvular structure and its competence, leading to more frequent JVR seen in patients with hypertension.

It should be noted that this pilot study had a relatively small sample size and that future larger case-control studies are needed to elucidate the differences of WMC severity between AD, MCI, and control groups. In addition, another limitation of the study is related to its cross-sectional nature, and further longitudinal studies are needed to determine temporal relationship between JVR and WMC lesion formation in relation to DAWM, particularly in the periventricular regions.

In conclusion, the findings of our pilot study showed that cerebral venous drainage impairment may play some role in the pathophysiology of WMC in the AD patients.

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DISCLOSURE STATEMENT

Authors' disclosures available online (<http://www.jalz.com/disclosures/view.php?id=1975>).

SUPPLEMENTARY MATERIAL

Supplementary material is available in the electronic version of this article: <http://dx.doi.org/10.3233/JAD-131112>.

REFERENCES

- [1] Hachinski VC, Potter P, Merskey H (1987) Leuko-araiosis. *Arch Neurol* **44**, 21-23.
- [2] Debette S, Markus HS (2010) The clinical importance of white matter hyperintensities on brain magnetic resonance imaging: Systematic review and meta-analysis. *BMJ* **341**, c3666.
- [3] Brickman AM, Muraskin J, Zimmerman ME (2009) Structural neuroimaging in Alzheimer's disease: Do white matter hyperintensities matter? *Dialogues Clin Neurosci* **11**, 181-190.
- [4] Tanaka T, Uemura K, Takahashi M, Takehara S, Fukaya T, Tokuyama T, Satoh A, Ryu H (1993) Compression of the left brachiocephalic vein: Cause of high signal intensity of the left sigmoid sinus and internal jugular vein on MR images. *Radiology* **188**, 355-361.
- [5] Akkawi NM, Agosti C, Borroni B, Rozzini L, Magoni M, Vignolo LA, Padovani A (2002) Jugular valve incompetence: A study using air contrast ultrasonography on a general population. *J Ultrasound Med* **21**, 747-751.
- [6] Chung CP, Lin YJ, Chao AC, Lin SJ, Chen YY, Wang YJ, Hu HH (2010) Jugular venous hemodynamic changes with aging. *Ultrasound Med Biol* **36**, 1776-1782.
- [7] Chung CP, Wang PN, Wu YH, Tsao YC, Sheng WY, Lin KN, Lin SJ, Hu HH (2011) More severe white matter changes in the elderly with jugular venous reflux. *Ann Neurol* **69**, 553-559.
- [8] Chung CP, Hu HH (2010) Pathogenesis of leukoaraiosis: Role of jugular venous reflux. *Med Hypotheses* **75**, 85-90.
- [9] Scheltens P, Barkhof F, Valk J, Algra PR, van der Hoop RG, Nauta J, Wolters EC (1992) White matter lesions on magnetic resonance imaging in clinically diagnosed Alzheimer's disease. Evidence for heterogeneity. *Brain* **115**, 735-748.
- [10] Kalaria RN (2000) The role of cerebral ischemia in Alzheimer's disease. *Neurobiol Aging* **21**, 321-330.
- [11] Meyer JS, Kawamura J, Terayama Y (1992) White matter lesions in the elderly. *J Neurol Sci* **110**, 1-7.
- [12] Rezek DL, Morris JC, Fulling KH, Gado MH (1987) Periventricular white matter lucencies in senile dementia of the Alzheimer type and in normal aging. *Neurology* **37**, 1365-1368.
- [13] Yoshita M, Fletcher E, Harvey D, Ortega M, Martinez O, Mungas DM, Reed BR, DeCarli CS (2006) Extent and distribution of white matter hyperintensities in normal aging, MCI, and AD. *Neurology* **67**, 2192-2198.
- [14] Pettersen JA, Sathiyamoorthy G, Gao FQ, Szilagyi G, Nadkarni NK, St George-Hyslop P, Rogaeva E, Black SE (2008) Microbleed topography, leukoaraiosis, and cognition in probable Alzheimer disease from the Sunnybrook dementia study. *Arch Neurol* **65**, 790-795.
- [15] Goldstein LB, Adams R, Alberts MJ, Appel LJ, Brass LM, Bushnell CD, Culebras A, Degraba TJ, Gorelick PB, Guyton JR, Hart RG, Howard G, Kelly-Hayes M, Nixon JV, Sacco RL; American Heart Association/American Stroke Association Stroke Council; Atherosclerotic Peripheral Vascular Disease Interdisciplinary Working Group; Cardiovascular Nursing Council; Clinical Cardiology Council; Nutrition, Physical Activity, and Metabolism Council; Quality of Care and Outcomes Research Interdisciplinary Working Group; American Academy of Neurology (2006) Primary prevention of ischemic stroke: A guideline from the American Heart Association/American Stroke Association Stroke Council: Cosponsored by the Atherosclerotic Peripheral Vascular Disease Interdisciplinary Working Group; Cardiovascular Nursing Council; Clinical Cardiology Council; Nutrition, Physical Activity, and Metabolism Council; and the Quality of Care and Outcomes Research Interdisciplinary Working Group: The American Academy of Neurology affirms the value of this guideline. *Stroke* **37**, 1583-1633.
- [16] McKhann G, Drachman D, Folstein M, Katzman R, Price D, Stadlan EM (1984) Clinical diagnosis of Alzheimer's disease: Report of the NINCDS-ADRDA Work Group under the auspices of Department of Health and Human Services Task Force on Alzheimer's Disease. *Neurology* **34**, 939-944.
- [17] Petersen RC, Doody R, Kurz A, Mohs RC, Morris JC, Rabins PV, Ritchie K, Rossor M, Thal L, Winblad B (2001) Current concepts in mild cognitive impairment. *Arch Neurol* **58**, 1985-1992.
- [18] Artero S, Petersen R, Touchon J, Ritchie K (2006) Revised criteria for mild cognitive impairment: Validation within a longitudinal population study. *Dement Geriatr Cogn Disord* **22**, 465-470.
- [19] Zivadinov R, Rudick RA, De Masi R, Nasuelli D, Ukmar M, Pozzi-Mucelli RS, Grop A, Cazzato G, Zorzon M (2001) Effects of IV methylprednisolone on brain atrophy in relapsing-remitting MS. *Neurology* **57**, 1239-1247.
- [20] Seewann A, Vrenken H, van der Valk P, Bleser EL, Knol DL, Castelijns JA, Polman CH, Pouwels PJ, Barkhof F, Geurts JJ (2009) Diffusely abnormal white matter in chronic multiple sclerosis: Imaging and histopathologic analysis. *Arch Neurol* **66**, 601-609.
- [21] Zivadinov R, Heininen-Brown M, Schirda CV, Poloni GU, Bergsland N, Magnano CR, Durfee J, Kennedy C, Carl E, Hagemeyer J, Benedict RH, Weinstock-Guttman B, Dwyer MG (2012) Abnormal subcortical deep-gray matter susceptibility-weighted imaging filtered phase measurements in patients with multiple sclerosis: A case-control study. *Neuroimage* **59**, 331-339.
- [22] van Straaten EC, Fazekas F, Rostrup E, Scheltens P, Schmidt R, Pantoni L, Inzitari D, Waldemar G, Erkinjuntti T, Mäntylä R, Wahlund LO, Barkhof F; LADIS Group (2006) Impact of white matter hyperintensities scoring method on correlations with clinical data: The LADIS study. *Stroke* **37**, 836-840.
- [23] Ge Y, Grossman RI, Babb JS, He J, Mannon LJ (2003) Dirty-appearing white matter in multiple sclerosis: Volumetric MR imaging and magnetization transfer ratio histogram analysis. *AJNR Am J Neuroradiol* **24**, 1935-1940.
- [24] Chen SC, Chung HW, Liou M (2003) Measurement of volumetric lesion load in multiple sclerosis: Moving from normal-to dirty-appearing white matter. *AJNR Am J Neuroradiol* **24**, 1929-1930.
- [25] Moore GR, Laule C, Mackay A, Leung E, Li DK, Zhao G, Traboulsee AL, Paty DW (2008) Dirty-appearing white matter in multiple sclerosis: Preliminary observations of myelin phospholipid and axonal loss. *J Neurol* **255**, 1802-1811.

- 483 [26] Laule C, Vavasour IM, Leung E, Li DK, Kozlowski P, Traboulsee AL, Oger J, Mackay AL, Moore GR (2011) Pathological basis of diffusely abnormal white matter: Insights from magnetic resonance imaging and histology. *Mult Scler* **17**, 144-150. 519
- 484 520
- 485 [27] Laule C, Pavlova V, Leung E, Zhao G, MacKay AL, Kozlowski P, Traboulsee AL, Li DK, Moore GR (2013) Diffusely abnormal white matter in multiple sclerosis: Further histologic studies provide evidence for a primary lipid abnormality with neurodegeneration. *J Neuropathol Exp Neurol* **72**, 42-52. 521
- 486 522
- 487 523
- 488 524
- 489 [28] Schaller B, Graf R (2004) Cerebral venous infarction: The pathophysiological concept. *Cerebrovasc Dis* **18**, 179-188. 525
- 490 526
- 491 [29] Zamboni P, Menegatti E, Weinstock-Guttman B, Schirda C, Cox JL, Malagoni AM, Hojanacki D, Kennedy C, Carl E, Dwyer MG, Bergsland N, Galeotti R, Hussein S, Bartolomei I, Salvi F, Zivadinov R (2009) The severity of chronic cerebrospinal venous insufficiency in patients with multiple sclerosis is related to altered cerebrospinal fluid dynamics. *Funct Neurol* **24**, 133-138. 527
- 492 528
- 493 529
- 494 [30] Wu IH, Sheng WY, Hu HH, Chung CP (2011) Jugular venous reflux could influence cerebral blood flow: A transcranial Doppler study. *Acta Neurol Taiwan* **20**, 15-21. 530
- 495 531
- 496 [31] Chung CP, Hsu HY, Chao AC, Cheng CY, Lin SJ, Hu HH (2009) Jugular venous reflux affects ocular venous system in transient monocular blindness. *Cerebrovasc Dis* **29**, 122-129. 532
- 497 533
- 498 [32] Ott BR, Cohen RA, Gongvatana A, Okonkwo OC, Johanson CE, Stopa EG, Donahue JE, Silverberg GD, Alzheimer's Disease Neuroimaging Initiative (2010) Brain ventricular volume and cerebrospinal fluid biomarkers of Alzheimer's disease. *J Alzheimers Dis* **20**, 647-657. 534
- 499 535
- 500 536
- 501 [33] Mawuenyega KG, Sigurdson W, Ovod V, Munsell L, Kasten T, Morris JC, Yarasheski KE, Bateman RJ (2010) Decreased clearance of CNS beta-amyloid in Alzheimer's disease. *Science* **330**, 1774. 537
- 502 538
- 503 [34] Reed-Cossairt A, Zhu X, Lee HG, Reed C, Perry G, Petersen RB (2012) Alzheimer's disease and vascular deficiency: Lessons from imaging studies and Down syndrome. *Curr Gerontol Geriatr Res* **2012**, 929734. 539
- 504 540
- 505 [35] Fazekas F, Chawluk JB, Alavi A, Hurtig HI, Zimmerman RA (1987) MR signal abnormalities at 1.5 T in Alzheimer's dementia and normal aging. *AJR Am J Roentgenol* **149**, 351-356. 541
- 506 542
- 507 [36] De Groot CJ, Bergers E, Kamphorst W, Ravid R, Polman CH, Barkhof F, van der Valk P (2001) Post-mortem MRI-guided sampling of multiple sclerosis brain lesions: Increased yield of active demyelinating and (p)reactive lesions. *Brain* **124**, 1635-1645. 543
- 508 544
- 509 [37] Sachdev P, Chen X, Wen W (2008) White matter hyperintensities in mid-adult life. *Curr Opin Psychiatry* **21**, 268-274. 545
- 510 546
- 511 [38] Yue NC, Arnold AM, Longstreth WT, Jr., Elster AD, Jungreis CA, O'Leary DH, Poirier VC, Bryan RN (1997) Sulcal, ventricular, and white matter changes at MR imaging in the aging brain: Data from the cardiovascular health study. *Radiology* **202**, 33-39. 547
- 512 548
- 513 [39] Beggs CB (2013) Venous hemodynamics in neurological disorders: An analytical review with hydrodynamic analysis. *BMC Med* **11**, 142. 549
- 514 550
- 515 [40] Dolic K, Weinstock-Guttman B, Marr K, Valnarov V, Carl E, Hagemeyer J, Kennedy C, Kilanowski C, Hojnacki D, Ramanathan M, Zivadinov R (2012) Heart disease, overweight, and cigarette smoking are associated with increased prevalence of extra-cranial venous abnormalities. *Neurol Res* **34**, 819-827. 551
- 516 552
- 517 [41] Fitzpatrick MA, Hinderliter AL, Egan BM, Julius S (1986) Decreased venous distensibility and reduced renin responsiveness in hypertension. *Hypertension* **8**, II36-II43. 553
- 518 554
- 519 [42] Takeshita A, Mark AL (1979) Decreased venous distensibility in borderline hypertension. *Hypertension* **1**, 202-206. 555
- 520 556
- 521 [43] Klabunde RE (2004) *Cardiovascular physiology concepts*, eds. Lippincott Williams and Wilkins, Philadelphia, PA. 557
- 522 558