

Association Between Large Numbers of Enlarged Perivascular Spaces in Basal Ganglia and Motor Performance in Elderly Individuals: A Cross-Sectional Study

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Background and Objective: Motor dysfunction is common in the elderly, and is associated with adverse consequences. Enlarged perivascular spaces in basal ganglia (BG-EPVSs) are considered an MRI marker of cerebral small-vessel diseases. However, the consequences of BG-EPVSs are largely unknown. In the present study, we aimed to explore the association between large numbers of BG-EPVSs and motor performance.

Methods: We prospectively recruited elderly individuals in the Neurology Department of our hospital from December 1, 2020 to January 31, 2022. Participants with >20 BG-EPVSs on the unilateral side of the slice containing the most EPVSs were classified as the BG-EPVS group (n=99) and the rest as controls (n=193). Motor performance was assessed by quantitative gait analysis, Tinetti test, timed up-and-go (TUG) test, and the Short Physical Performance Battery (SPPB). Spearman correlation analysis and multivariate linear regression analysis were performed to investigate the association between BG-EPVSs and motor performance.

Results: Compared with the control group, the BG-EPVS group had lower gait speed and cadence, shorter stride length, longer TUG duration, and lower Tinetti gait test, Tinetti balance test, and SPPB scores ($P<0.01$). Spearman correlation analysis showed that BG-EPVSs were negatively related to gait speed, gait cadence, stride length, and Tinetti gait test, Tinetti balance test, and SPPB scores ($\rho=-0.539$ to -0.223 , $P<0.001$) and positively related to TUG duration ($\rho=0.397$, $P<0.001$). Regression analysis indicated that BG-EPVSs were an independent risk factor of lower gait speed, shorter stride length, poor balance, and poor general physical performance after adjusting for confounders ($\beta=-0.313$ to -0.206 , $P<0.01$).

Conclusion: Large numbers of BG-EPVSs were independently related to poor gait, balance, and general physical performance in elderly individuals, which provides information about the consequences of BG-EPVSs and risk factors for motor dysfunction.

Keywords: cerebral small-vessel diseases, enlarged perivascular spaces, motor, gait, balance, physical performance

Introduction

Motor dysfunction is common in the elderly, and is related to adverse consequences, eg, falls, institutionalization, and death.^{1,2} There is important clinical significance in exploring the risk factors and mechanisms of motor dysfunction in elderly individuals. Normal motor function depends on the coordination of multiple brain regions.^{3,4} The basal ganglia (BG) are an important region related to the control of normal motor function, including gait, balance, and general physical performance.⁵

Perivascular spaces (PVSs), or Virchow–Robin spaces, are PV compartments surrounding small cerebral penetrating vessels serving as a protolymphatic system, and play an important role in interstitial fluid and solute clearance in brain.⁶ They dilate with the accumulation of interstitial fluids. Enlarged PVSs (EPVSs), visible on magnetic resonance imaging (MRI), appear as punctate or linear signal intensities similar to cerebrospinal fluid on all MRI sequences.⁷ EPVSs in BG (BG-EPVSs) are recognized as an MRI marker of cerebral small-vessel diseases (CSVDs).^{8,9}

Many studies have demonstrated that CSVDs are associated with vascular dementia, depression, motor dysfunction, and increased risk of stroke.^{10–12} Some scholars have proposed that CSVDs deserve more attention for their relevance and impact on clinical practice.¹³ However, the consequences of BG-EPVSs are largely unknown. Several BG-EPVSs are generally considered clinically silent. Recent studies have shown a relationship between EPVSs and cognitive disturbances;^{14,15} but others have found EPVSs not to be related to cognitive impairment.¹⁶ Therefore, the association between EPVSs and cognitive impairment is attributed to a direct effect of EPVSs or accompanying CSVD markers, eg, white-matter hyperintensity (WMH), cerebral microbleeds (CMBs), and lacunae, and remains incompletely understood.

Studies have shown that WMH, CMBs, and lacunae are related to physical performance¹⁷ and gait disturbances.^{10,18,19} However, the relationship between BG-EPVSs and motor function is unclear. We speculated that a large number of BG-EPVSs might be associated with motor disturbances via disrupting the function of the BG. Therefore, we explored the association between large numbers of BG-EPVSs and motor performance in elderly individuals.

Methods

Figure 1 presents a flowchart of the research process.

Participants and Grouping

We prospectively reviewed medical records of elderly patients with CSVDs without acute cerebral infarction in the Neurology Department of Beijing Chaoyang Hospital affiliated with Capital Medical University from December 1, 2020 to January 31, 2022. CSVDs were defined as the presence of WMH and/or CMBs and/or lacunae of presumed vascular origin on brain MRI. Inclusion criteria were age 60 years or above and agreement to participate.

Exclusion criteria were: dementia, including Alzheimer's disease, frontotemporal dementia, or dementia with Lewy bodies; Parkinson's disease (PD), PD-plus syndrome, including multiple-system atrophy, corticobasal degeneration, and progressive supranuclear palsy; a history of severe stroke (largest lesion diameter >20 mm) due to difficulties and inaccurate assessments on CSVD MRI markers, or large-vessel cerebrovascular diseases, defined as internal carotid, middle cerebral, or basilar intracranial artery stenosis >50%; lacunar syndrome within 6 months after the event to avoid acute effects on outcomes, or patients with stroke sequelae 6 months after cerebral infarction onset; traumatic, toxic, or infectious brain injury, brain tumor, or brain metastases, non-CSVD-rated white-matter lesions, eg, multiple sclerosis and irradiation-induced gliosis; inability to walk for 30 m unaided; inability finish the tests because of prominent visual, hearing, language impairment, or psychiatric disease; conditions not related to CSVDs affecting motor function (eg, diabetic peripheral neuropathy, joint fusion, severe arthritis, joint replacement, or lumbar spondylopathy); heart failure, myocardial infarction, or angina pectoris disorders during the previous 3 months, severe nephrosis/liver diseases, and life expectancy <6 months; MRI contraindications/known claustrophobia/poor imaging quality affecting CSVD assessments because the patient's head moved when performing MRI; and any problem in acquiring quantitative gait data.

All enrolled patients were divided into two groups based on the number of BG-EPVSs on axial T_2 -weighted images. Patients with >20 BG-EPVSs on the unilateral side of the slice containing the most EPVSs comprised the BG-EPVS group (Figure 2A) and the rest were controls (Figure 2B). The cutoff of 20 BG-EPVSs has been used to represent “high grade” in previous studies.²⁰

Demographic and Clinical Assessments

Age, sex, BMI, and history of hypertension, diabetes, hyperlipidemia, stroke, smoking, and alcohol consumption were collected. All blood samples were collected in the morning and sent to the clinical laboratory of our hospital for measurement of serum indices. Laboratory tests included total cholesterol (TC), triglycerides, high-density lipoprotein (HDL), low-density lipoprotein (LDL), HbA_{1c}, blood urea nitrogen (BUN), creatinine, and homocysteine.

MRI Examinations and Assessment of EPVSs, WMH, CMBs, and Lacunae

MRI was performed on a 3 T MRI scanner (Prisma; Siemens, Erlangen, Germany) in the Radiology Department of our hospital. The standardized MRI sequences used were axial T_1 -weighted, T_2 -weighted, and diffusion-weighted imaging, fluid-attenuated inversion recovery, and susceptibility-weighted imaging. Imaging markers of CSVDs, ie, EPVSs, WMH,

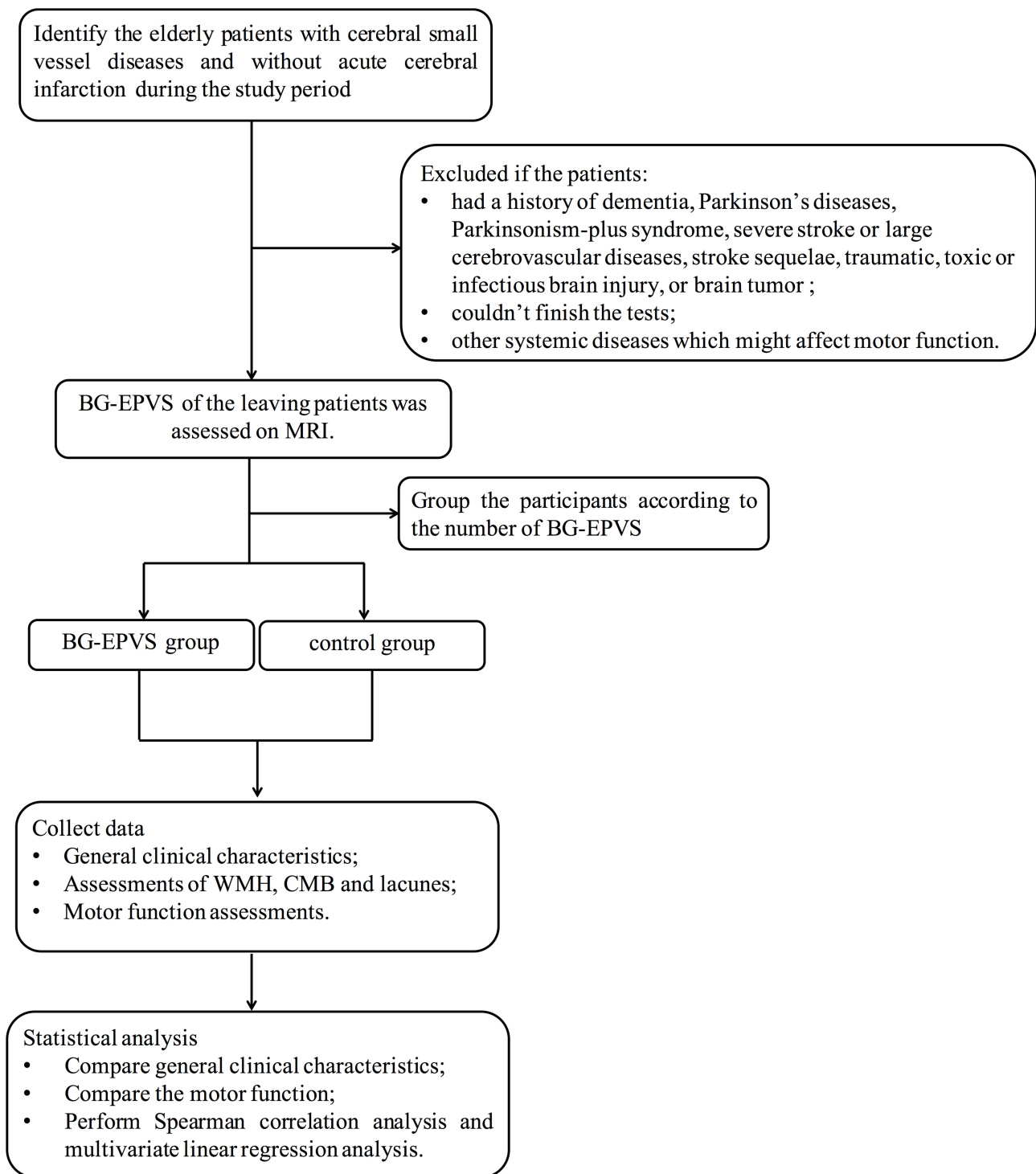


Figure 1 Flowchart of study process.

CMBs, and lacunae, were defined as per the Standards for Reporting Vascular Changes on Neuroimaging criteria.⁸ BG-EPVSs were counted in the slice containing the most EPVSs. Periventricular and deep WMH were evaluated separately and combined as Fazekas scores. A detailed description of the Fazekas scale has been published.²¹ CMBs and lacunae were counted.

Assessments of EPVSs, WMH, CMBs, and lacunae were performed by two experienced neurologists blinded to clinical data. Random scans of 50 individuals were independently examined by the two neurologists blinded to each

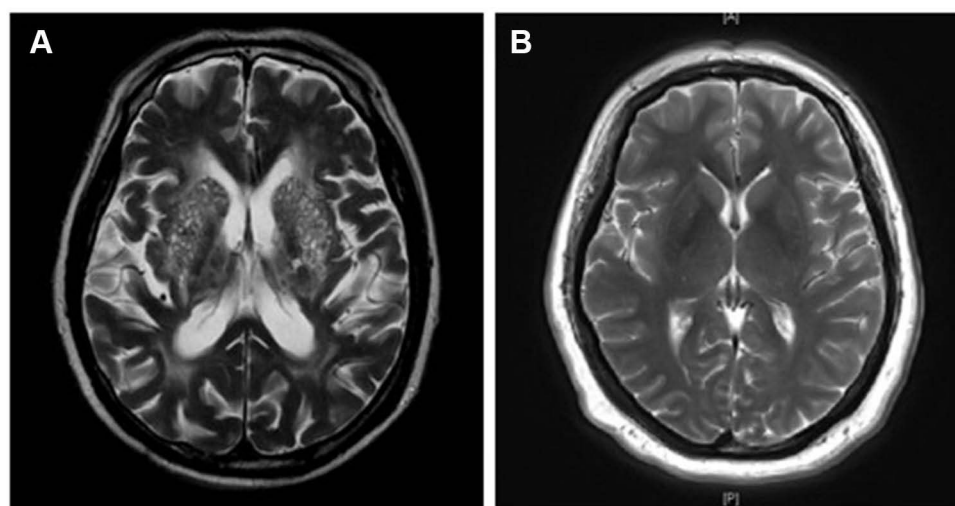


Figure 2 MRI of BG-EPVS and control groups. Patients with >20 BG-EPVSs on the unilateral side of the basal ganglia slice containing the most EPVSs were defined as the BG-EPVS group (A) and the rest as controls (B).

other's readings. Intrarater agreement for EPVSs, WMH, CMBs, and lacunae was assessed on a random sample of 50 individuals with a month's interval between the first and second readings. Intrarater and interrater agreement had values of $\kappa=0.80\text{--}0.98$, indicating good reliability. Disagreement was resolved through discussion with the other authors.

Motor-Function Assessments

Motor-function assessments covered gait, balance, and general physical performance based on quantitative gait analysis, timed up-and-go (TUG) test and clinical scales consisting of the Tinetti test and Short Physical Performance Battery (SPPB). Quantitative gait parameters were acquired by a MiniSun IDEEA, which has an excellent test–retest reliability and validity.^{22,23} The IDEEA is equipped with Windows-based software that analyzes gait data and provides parameters of gait speed, gait cadence (number of steps per minute), and stride length (distance between the heel points of two consecutive footprints from the same foot). The participants were instructed to walk 8 m wearing the device at their usual gait speed in low-heeled shoes before the formal test. Then, the formal 8 m test was performed.

The TUG test was performed and the duration recorded using a 3 m walk. The Tinetti test contains 17 items — nine for balance (score 0–16) and eight for gait (score 0–12) — with a maximum score of 28.²⁴ The Tinetti test was independently scored by the two neurologists blinded to clinical data, and then the scores were averaged. Interrater reliability was good ($\kappa=0.9$). General physical performance was measured with the SPPB. This scale consists of three measurements of gait speed (score 0–4), standing balance (score 0–4), and chair stand (score 0–4). The gait speed subscale uses a 4 m walk. Total SPPB scores range from 0 to 12 points. Measurement and scoring methods of the SPPB have been described previously.¹⁷

Statistical Analysis

Continuous variables are presented as means \pm SD or medians (interquartile range) according to conformation with normal distribution. Categorical variables are presented as numbers and percentages. Continuous variables with both normal distribution and homogeneity of variance were compared with Student's *t* test, and otherwise with Wilcoxon's rank-sum test. For comparison of categorical variables, χ^2 tests were used. Spearman correlation analysis was performed to analyze correlations between large numbers of BG-EPVSs and motor function. Partial correlation analysis and multivariate linear regression analysis were performed to analyze if BG-EPVSs were independently related to motor function after adjusting for confounding factors. Analysis was performed with SPSS 21.0, and statistical significance was accepted at $P<0.05$.

Results

Participants' General Clinical Characteristics

During the study period, 442 elderly patients with CSVDs and without acute cerebral infarction were identified. In sum, 52 patients were excluded because of a history of severe stroke and large-vessel diseases, 16 because of stroke sequelae, seven because of a diagnosis of PD. Another 40 were excluded because they had severe arthritis, joint fusion, lumbar spondylopathy, or visual impairment affecting walking, four because of poor imaging quality not clear enough to assess CSVD markers and 28 patients did not agree to participate. Three patients were excluded because of problems in acquiring quantitative gait parameters. Finally, 292 elderly patients were enrolled and finished the tests, of which 99 were classified into BG-EPVS group and 193 served as controls according to the number of BG-EPVSs.

The general clinical characteristics of all participants are presented in Table 1. Mean age was 72 ± 7.7 years, and 171 (58.56%) were men. The BG-EPVS group was older than the controls (75 ± 8 vs 70 ± 7 , $P<0.001$). Male sex, hypertension, diabetes, and stroke in the BG-EPVS group were more predominant than the control group ($P<0.05$). The BG-EPVS group had higher levels of BUN (6.2 ± 1.86 vs 5.8 ± 2.00 , $P=0.047$), creatinine (76 ± 23.1 vs 68 ± 20.9 , $P=0.010$), and homocysteine (16 ± 6.40 vs 13 ± 5.63 , $P=0.004$). There were no statistical differences in BMI, hyperlipidemia, smoking, alcohol consumption, TC, triglycerides, HDL, LDL, or HbA_{1c} between the groups. In terms of other CSVD markers, the BG-EPVS group had more serious WMH and more lacunae and CMBs ($P<0.001$).

Association Between Motor Function and BG-EPVSs

Table 2 presents the motor performance of the groups. The BG-EPVS group had lower gait speed and cadence, shorter stride length (Figure 3), and longer TUG duration. Tinetti gait test and balance test scores were lower in the BG-EPVS group ($P<0.05$). Gait speed, standing balance, chair stand, and total scores on the SPPB were lower in BG-EPVS group ($P<0.05$). Spearman correlation analysis showed that large numbers of BG-EPVSs were negatively

Table 1 General clinical characteristics of participants

	All	BG-EPVS group	Control group	P
n	292	99	193	—
Age ^a , years	72 ± 7.7	75 ± 8.0	70 ± 7.0	<0.001
Male, n (%)	171 (58.56)	71 (71.72)	100 (51.81)	0.001
BMI ^a , kg/m ²	25 ± 3.7	25 ± 3.6	25 ± 3.8	0.931
Smokers, n (%)	91 (31.16)	36 (36.36)	55 (28.50)	0.169
Alcohol drinkers, n (%)	48 (16.44)	17 (17.17)	31 (16.06)	0.809
Hypertension, n (%)	205 (70.21)	77 (77.78)	128 (66.32)	0.043
Diabetes, n (%)	77 (26.37)	35 (35.35)	42 (21.76)	0.013
Hyperlipidemia, n (%)	105 (35.96)	35 (35.35)	70 (36.27)	0.877
Stroke, n (%)	69 (23.63)	40 (40.40)	29 (15.03)	<0.001
TC ^a , mmol/L	4.4 ± 0.99	4.3 ± 1.00	4.4 ± 1.00	0.153
Triglycerides ^a , mmol/L	1.6 ± 0.91	1.5 ± 0.88	1.6 ± 0.90	0.633
HDL ^a , mmol/L	1.2 ± 0.41	1.1 ± 0.29	1.2 ± 0.46	0.266
LDL ^a , mmol/L	2.8 ± 0.93	2.7 ± 0.95	2.8 ± 0.93	0.255
HbA _{1c} ^a , %	6.4 ± 1.15	6.5 ± 1.28	6.3 ± 1.06	0.143
BUN ^a , mmol/L	5.8 ± 2.00	6.2 ± 1.86	5.7 ± 2.05	0.047
Creatinine ^a , umol/L	71 ± 21.9	76 ± 23.1	68 ± 20.9	0.010
Homocysteine ^a , μmol/L	14 ± 5.81	16 ± 6.40	13 ± 5.63	0.004
WMH ^b , Fazekas score	3 (2–4)	4 (3–5)	2 (2–3)	<0.001
Lacunae ^b , n	0 (0–2)	2 (0.5–4)	0 (0–1)	<0.001
CMBs ^b , n	0 (0–1)	1 (0–5.5)	0 (0–1)	<0.001

Notes: ^aContinuous variables are expressed as means \pm SD and were compared using Student's *t*-test. ^bContinuous variables are expressed as medians (interquartile range) and were compared using Wilcoxon's rank-sum test.

Abbreviations: TC, total cholesterol; HDL, high-density lipoprotein; LDL, low-density lipoprotein; BUN, blood urea nitrogen; WMH, white-matter hyperintensity; CMBs, cerebral microbleeds.

Table 2 Motor performance of BG-EPVS and control groups

	BG-EPVS group	Control group	P
Quantitative gait parameters			
Speed ^a , m/min	39.8±13.75	51.3±12.84	<0.001
Cadence ^a , steps/min	97±14.9	102±12.33	0.004
Stride length ^a , m	0.8±0.20	1.0±0.18	<0.001
TUG duration ^a , seconds	18.9±10.66	13.0±4.09	<0.001
Tinetti test			
Gait ^b score	11 (8.5–12)	12 (12–12)	<0.001
Balance ^b score	14 (10–16)	16 (16–16)	<0.001
SPPB			
Gait speed ^b score	1 (1–2)	2 (2–3)	<0.001
Standing balance ^b score	1 (1–3)	4 (3–4)	<0.001
Chair stand ^b score	2 (2–3)	4 (3–4)	<0.001
Sum score ^b	6 (4–8)	10 (8–11)	<0.001

Notes: ^aContinuous variables are expressed as means ± SD and were compared using Student's *t*-test. ^bContinuous variables are expressed as medians (interquartile range) and compared using Wilcoxon's rank-sum test.

related to gait speed, gait cadence, and stride length and positively related to TUG duration (Table 3). Large numbers of BG-EPVSs were negatively related to Tinetti gait test and balance test scores and SPPB gait speed, standing balance, chair stand, and total scores. We further conducted partial correlation analysis to adjust for confounding factors of general clinical characteristics (age, sex, history of hypertension, diabetes, stroke, BUN, creatinine, and homocysteine) and other CSVD markers (WMH, lacunae, and CMBs). The negative correlation between large numbers of BG-EPVSs and gait speed, stride length, balance, and general physical performance did not change after adjustment (Table 3). Large numbers of BG-EPVSs were not independently related to TUG duration or gait cadence after adjusting for confounding factors.

To identify the effect of BG-EPVSs on motor function, we took the gait parameters, Tinetti test, and SPPB scores as dependent variables and conducted multiple linear regression analysis. This showed that large numbers of BG-EPVSs were an independent risk factor of lower gait speed, shorter stride length, poor balance, and poor general physical performance after adjusting for age, sex, history of hypertension, diabetes, and stroke, BUN, creatinine, homocysteine, Fazekas score, and number of lacunae and CMBs. Detailed results are presented in Table 4.

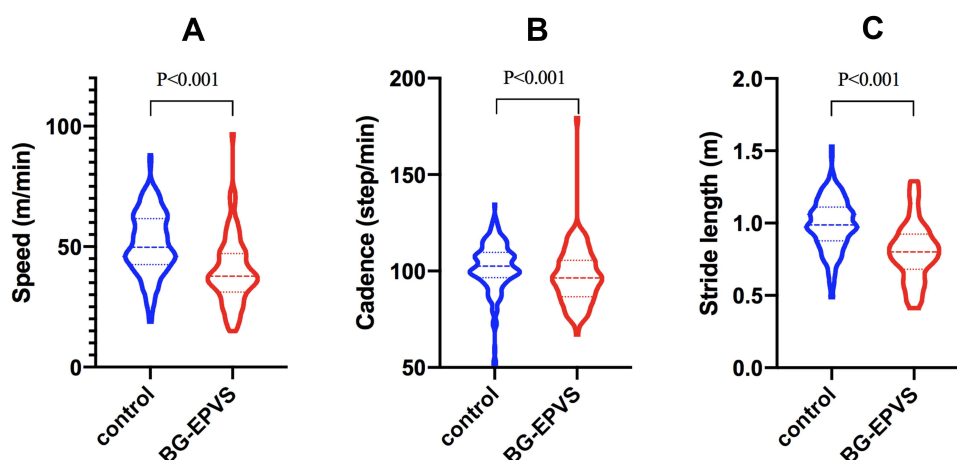
**Figure 3** Violin plot of quantitative gait parameters. (A–C) Gait speed, gait cadence, and stride length of BG-EPVS group and control group, respectively.

Table 3 Correlations between motor function and BG-EPVs

	ρ	P	ρ (adjusted for confounders)	P (adjusted for confounders)
Quantitative gait parameters				
Speed, m/min	-0.404	<0.001	-0.181	0.008
Cadence, steps/min	-0.223	<0.001	-0.052	0.445
Stride length, m	-0.426	<0.001	-0.216	0.001
TUG duration, seconds	0.397	<0.001	0.126	0.065
Tinetti test				
Gait score	-0.507	<0.001	-0.213	0.002
Balance score	-0.512	<0.001	-0.269	<0.001
SPPB				
Gait speed score	-0.396	<0.001	-0.248	<0.001
Standing balance score	-0.534	<0.001	-0.285	<0.001
Chair stand score	-0.420	<0.001	-0.180	0.008
Sum score	-0.539	<0.001	-0.333	<0.001

Note: Confounders included age, sex, hypertension, diabetes, stroke, blood urea nitrogen, creatinine, homocysteine, Fazekas score, and number of lacunae and CMBs.

Table 4 Multiple linear regression analysis of motor function and BG-EPVs

	Model 1		Model 2		Model 3	
	β	P	β	P	β	P
Quantitative gait parameters						
Speed, m/min	-0.305	<0.001	-0.321	<0.001	-0.206	0.003
Cadence, steps/min	-0.114	0.069	-0.121	0.087	-0.080	0.327
Stride length, m	-0.342	<0.001	-0.333	<0.001	-0.217	0.001
TUG duration, seconds	0.286	<0.001	0.249	<0.001	0.125	0.064
Tinetti test						
Gait test score	-0.395	<0.001	-0.377	<0.001	-0.226	0.001
Balance test score	-0.423	<0.001	-0.395	<0.001	-0.269	<0.001
SPPB						
Gait speed score	-0.280	<0.001	-0.268	<0.001	-0.291	<0.001
Standing balance score	-0.459	<0.001	-0.394	<0.001	-0.285	<0.001
Chair stand score	-0.343	<0.001	-0.339	<0.001	-0.207	0.002
Sum score	-0.438	<0.001	-0.400	<0.001	-0.313	<0.001

Notes: Model 1: adjusted for age and sex. Model 2: model 1 + history of hypertension, diabetes, and stroke, blood urea nitrogen, creatinine, and homocysteine. Model 3: model 2 + Fazekas score and number of lacunae and CMBs.

Discussion

In the present study, we explored the association between large numbers of BG-EPVs and motor function in elderly individuals with CSVDs. Gait, balance, and general physical performance were covered in motor-function assessments. We found that large numbers of BG-EPVs were independently related to lower gait speed and shorter stride length after adjusting for other CSVD markers (WMH, CMBs, and lacunae). We also found that large numbers of BG-EPVs were independently related to poor balance and general physical performance.

Previous studies have investigated the relationship between motor function and other CSVD MRI markers (WMH, lacunar infarction, and CMBs).¹¹ The LADIS study adopted SPPB, single-leg stance time, and walking speed to assess motor function, and found that SPPB total score, single-leg stance time, and walking speed decreased with increased WMH and that frontal deep WMH was significantly associated with balance disturbances.²⁵ The RUN DMC study found

that WMH and lacunar infarcts were both independently associated with most gait parameters and that stride length was the most sensitive parameter related to WMH.²⁴ WMH in BG and lacunar infarcts in the thalamus were related to lower velocity. The study also found a higher number of CMBs in the BG was independently related to shorter stride length and poorer performance on Tinetti tests.²⁶ However, to the best of our knowledge, clinical studies specifically addressing the association between EPVSs and gait in CSVD patients are sparse.

Shin et al²⁷ explored the relationship between BG-EPVSs, EPVSs in white matter (WM-EPVSs), and motor function in patients with PD. They found that severe BG-EPVSs were associated with worse motor symptoms. Chung et al²⁸ investigated the association between BG-EPVSs and long-term motor outcomes in patients with PD. They found that the PD-EPVS⁺ group exhibited more severely decreased dopamine transporters, and the risk of freezing gait was higher in the PD-EPVS⁺ group. The PD-EPVS⁺ group required higher doses of dopaminergic medications for effective symptom control than the PD-EPVS⁻ group. In the present study, we explored the relationship between BG-EPVSs and motor performance in elderly CSVD patients. We also found that severe BG-EPVSs were independently related to worse gait performance and balance, similar to the results of previous studies. Sun et al²⁹ investigated the correlation between CSVD burden and motor performance in community-dwelling populations. They found that EPVSs were not associated with motor performance, different from our results. They classified patients with more than five BG-EPVSs on the BG slice containing the most EPVSs into the EPVS group. The different results might be attributed to different grouping methods and study populations.

Clinical experience and other studies have indicated that moderate or severe CSVDs can lead to clinical symptoms.^{24,30} Mild CSVDs are generally considered clinically silent. For example, gait disturbances are only attributed to moderate or severe WMH and/or the presence of more than three lacunar infarcts.²⁴ Therefore, in the present study we classified subjects with high-degree BG-EPVSs into the BG-EPVS group and the rest into control group. As we all know, subcortical WM is another common region of EPVSs, but the anatomical structure, mechanisms, and risk factors of BG-EPVSs and WM-EPVSs are different.^{6,9} WM-EPVSs are related to cerebral amyloid angiopathy, whereas BG-EPVSs are related to hypertensive arteriopathy and recognized as an MRI marker of CSVDs. Additionally, BG play an important role in motor-function control. Therefore, in the present study we explored the relationship between BG-EPVSs and motor function. The consequences of large WM-EPVSs should be explored in the future.

Gait velocity is determined by both stride length and gait cadence. In the present study, gait velocity, stride length, and gait cadence were reduced in patients with large numbers of BG-EPVSs. However, BG-EPVSs were significantly related only to gait velocity and stride length and not to cadence after adjusting for WMH, lacunae, and CMBs. Therefore, we think that stride length may be a more sensitive indicator than gait cadence related to BG-EPVSs. Apart from the lower velocity, we found that patients with large numbers of BG-EPVSs were characterized by lower Tinetti balance and SPPB scores, which indicated large numbers of BG-EPVSs were related to poor balance and general physical performance.

The potential pathophysiological mechanisms underlying the association between large numbers of BG-EPVSs and motor function are not completely understood. Motor, balance, and gait abilities are based on the integrity of the complex motor-circuit network that connects the BG and brain-stem structures with the frontal cortex. This network's function may be challenged by a critical number of lesions in subcortical regions.²⁵ PVSs are thought to serve as a protolymphatic system and play an important role in maintaining neural homeostasis.⁶ Sulcal cerebrospinal fluid is cleared through arachnoid granulations or enters the parenchyma via the PVSs, where it combines with interstitial fluid prior to exiting the brain. This PV drainage system also allows for the clearance of toxic metabolites within the parenchyma and plays a role in the brain's immunological response.³¹ EPVSs are thought to be the result of PV blockage and disrupt the normal function of PVSs. Although a few EPVSs visible on MRI can be normal, the presence of many is not normal, and they have been shown to be associated with some age-related disorders, including cognitive dysfunction, Parkinson's syndrome, WMH, and lacunar infarction.^{6,7,32} Studies have found that the BG-thalamocortical circuit, brain-network efficiency, and loss of WM-microstructure integrity are involved in gait impairment in subjects with CSVDs.^{33,34} Recent findings showed that EPVSs lead to WM-microstructure damage by causing lymphatic circulation dysfunction. The glymphatic aberration results in the accumulation of toxic metabolic products that are harmful to the brain microenvironment.³⁵ Therefore, we speculate that large numbers of BG-EPVSs might disrupt the functioning of the

BG, brain network, and brain microenvironment, then lead to poor motor performance. This hypothesis should be tested and verified by multimodal MRI (eg, functional MRI and diffusion-tensor imaging) in the future.

Some limitations in the present study must be mentioned. Firstly, it was performed in a single center and the cohort may not represent the general population. Secondly, this was an observational study, and causal relationships between motor dysfunction and BG-EPVSS could not be established. Thirdly, the mechanisms underlying the association between BG-EPVSS and motor function were not explored. A multicenter prospective cohort study should be performed in the future with multimodal MRI being used to explore the mechanisms. Fourthly, we did not include brain-atrophy assessments, which are recognized as another potentially relevant manifestation of CSVDs (although still poorly characterized) and related to motor dysfunction in elderly individuals.^{36,37} Despite these limitations, this study is the first to find that large numbers of BG-EPVSS were associated with poor gait, balance, and general physical performance in elderly individuals with CSVDs by both quantitative and semiquantitative measures. These novel findings provide some information about the consequences of large numbers of BG-EPVSS and risk of motor disturbances in elderly people.

In addition, we would like to mention that the relationship between large numbers of BG-EPVSS and cognitive impairment should also be explored in elderly silent CSVD patients in the future. Many studies have demonstrated that other CSVD markers (WMH, CMBs, and lacunae) are related to cognitive impairment.¹⁰ Grau-Olivares et al found up to 57.8% of first acute lacunar stroke patients had mild neuropsychological disorders.³⁸ The incidence of cognitive impairment in patients with large numbers of BG-EPVSS is not clear, research on the relationship between large numbers of BG-EPVSS and cognitive impairment insufficient, and results have not been consistent.^{14,16}

Conclusion

We found that large numbers of BG-EPVSS were independently related to gait disturbances, poor balance, and general physical performance. Patients with large numbers of BG-EPVSS had lower gait speed and shorter stride length. Causal relationships and mechanisms should be further tested and explored by longitudinal studies in the future.

Data Sharing

Data that support the findings of this study are available from the corresponding author on reasonable request.

Ethical Standards

This study was approved by the Ethics Committee of Beijing Chaoyang Hospital affiliated with Capital Medical University and was conducted in accordance with the Declaration of Helsinki (ethics approval 2020-216). All participants provided informed written consent.

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Disclosure

The authors declare that they have no competing interests for this work, the materials or methods used, or the findings specified in this paper.

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