Cerebral small vessel disease phenotype and 5-year mortality in asymptomatic middle-to-old aged individuals

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The present study aimed to determine whether a recently proposed cerebral small vessel disease (CSVD) classification scheme could differentiate the 5-year all-cause mortality in middle-to-old aged asymptomatic CSVD. Stroke-free and non-demented participants recruited from the community-based I-Lan Longitudinal Aging Study underwent baseline brain magnetic resonance imaging (MRI) between 2011 and 2014 and were followed-up between 2018 and 2019. The study population was classified into control (non-CSVD) and CSVD type 1–4 groups based on MRI markers. We determined the association with mortality using Cox regression models, adjusting for the age, sex, and vascular risk factors. A total of 735 participants were included. During a mean follow-up of 5.7 years, 62 (8.4%) died. There were 335 CSVD type 1 (57.9 ± 5.9 years), 249 type 2 (65.6 ± 8.1 years), 52 type 3 (67.8 ± 9.2 years), and 38 type 4 (64.3 ± 9.0 years). Among the four CSVD types, CSVD type 4 individuals had significantly higher all-cause mortality (adjusted hazard ratio = 5.0, 95% confidence interval 1.6–15.3) compared to controls. This novel MRI-based CSVD classification scheme was able to identify individuals at risk of mortality at an asymptomatic, early stage of disease and might be applied for future community-based health research and policy.

CNS (central nervous system) or cerebral small vessel disease (CSVD) causes 25% of stroke1. It is age-related and also considered as an important etiology of geriatric syndromes such as dementia, gait disturbance, and mood disorders2,3. The etiologies of age-related CSVD are heterogeneous and include two most common forms, arteriosclerosis/ lipohyalinosis and amyloid accumulation (cerebral amyloid angiopathy; CAA)1–3. These brain microvascular pathologies might result in brain parenchymal damage through ischemia, edema or hemorrhage, particularly in the cerebral white matter1–3. Owing to developments in magnetic resonance imaging (MRI), we can now identify the presence of CSVD by visualizing the associated brain parenchyma changes at pre-mortem2,4. MRI markers of SVD are also heterogeneous and include ischemic lesions, such as white matter hyperintensities (WMH) and lacune(s), and hemorrhagic lesions, the cerebral microbleeds (CMB)2,5. These CSVD-related brain abnormalities usually co-occur in different etiologies, and their clinical courses are variable6. Clinical significances of each MRI CSVD marker have been revealed in several studies, particular in patients with stroke or dementia1,2,5,6. However, since each individual with CSVD usually has different combinations and severity of MRI markers, the known clinical risks of each single MRI marker might be hard to be applied in predicting individual's clinical outcomes.

There is no effective treatment or method for the prevention of CSVD despite its acknowledged significance. It could be due to a lack of phenotyping methods that could stratify CSVD in response to its heterogeneity in clinical and neuroimaging manifestations and etiology5. Current CSVD indices are often derived by summing

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the different MRI markers that are present in an individual (CSVD score)\textsuperscript{12}. These scoring methods may reflect the CSVD burden and predict prognosis\textsuperscript{2}. However, they do not provide information about the underlying pathogenesis of CSVD. Recently, we developed a stratification scheme to classify a community-based, asymptomatic middle-to-old aged population into non-CSVD and four CSVD subtypes based on the following criteria in order: (1) bleeding or non-bleeding, (2) CMB locations, and (3) the severity and combination of WMH and lacune (Fig. 1b)\textsuperscript{7}. Common and distinct patterns of the clinical and neuroimaging manifestations were found in the four stratified CSVD subgroups\textsuperscript{7}. This MRI-based stratification scheme highlights the distinct features of SVD pathogenesis of CSVD. 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Figure 1. (a) Flowchart of included participants and analyzed neuroimaging data. (b) Methodological sequence for phenotyping age-related cerebral small vessel disease. The stratification scheme had three steps in the following order: (1) presence of cerebral microbleeds, (2) presence of severe WMH (defined as > 50th percentile of WMH/TIV ratio), and (3) a combination of lacunes with severe WMH or the geographic patterns of cerebral microbleeds (mixed or strictly lobar) if cerebral microbleeds were present. Participants without cerebral microbleeds and severe WMH were classified as the control group. CSVD, cerebral small vessel disease; FLAIR, fluid-attenuated inversion-recovery; SWI, susceptibility-weighted images; TIV, total intracranial volume; WMH, white matter hyperintensities.
(TIV, summation of gray matter volume, white matter volume and cerebrospinal fluid volume) and WMH volume simultaneously.

**Detection and assessment of MRI SVD markers.** CMBs were defined as small, rounded or circular, hypointense lesions within the brain parenchyma with clear margins and ≤ 10 mm in size on SWI. Microbleed mimics, such as vessels, calcification, partial volume, air-bone interfaces, and hemorrhages within or adjacent to an infarct, were carefully excluded. We used the microbleed anatomical rating scale to measure the presence, amount, and topographic distribution of CMBs. Intra-rater reliability was assessed by evaluating CMBs in 20 randomly sampled images at a separate time (K, 0.83; 95% confidence interval [CI], 0.79–0.90). We also reassessed CMBs in the 25 randomly sampled images previously assessed by Dr. Chung and another investigator (K, 0.82; 95% CI, 0.79–0.88). CMBs were classified into deep, infratentorial, and lobar cate-
Determination of CSVD types. The CSVD stratification scheme (CSVD types) had three steps in order: checking for the presence of (1) CMB, (2) severe WMH (defined as >50th percentile of WMH/total intracranial volume ratio [0.07%]), and (3) a combination of lacunes with severe WMH or a certain geographic pattern of CMB (mixed or strictly lobar) when CMB is present (Fig. 1b). The differentiation between bleeding and nonbleeding SVD was achieved in the first step of the stratification scheme. We further stratified bleeding SVDs into different subtypes with CMBs of specific geographic features. Regarding the non-bleeding SVDs, subjects with severe WMHs were further divided depending on whether a lacune was present. Participants without CMB and severe WMH were allocated to the robust (control) group. There were two types of nonbleeding SVD (WMH without or with lacune; CSVD type 1 and 2) and two types of bleeding SVD (mixed or strictly lobar CMB; CSVD type 3 and 4).

CSVD burden. We used a simple CSVD score to represent the CSVD burden\(^1\). One point was given for the presence of any lacune, severe WMH, and CMB; thus, the simple CSVD score ranged from 0 to 3.

Statistical methods. Analyses were performed using SPSS version 22.0. (IBM, Armonk, NY, USA). All data are presented as mean (standard deviation) for continuous variables and number (percentage) for discrete variables. Group comparisons were made using the nonparametric Kruskal–Wallis test with post-hoc analyses. When appropriate, chi-square or Fisher's exact tests were performed for categorical variables.

The follow-up time for each individual was calculated from the date of initial recruitment until the date of the phone interview. The incidence rate of all-cause mortality was determined from the incidence per person year. A Kaplan–Meier survival curve was plotted, and the log-rank test was applied to test the difference in survival between groups. We then used the Cox regression analysis to calculate the crude and adjusted hazard ratios (HRs) and 95% CIs for the occurrence of all-cause mortality in each CSVD group compared to the control group. The covariates included the age, sex, and vascular risk factors (presence of hypertension, diabetes mellitus, and dyslipidemia, and cigarette smoking). There was no significant 13 in the HR changes with time in CSVD types \(p = 0.061\) and CSVD scores \(p = 0.057\), which showed that the assumption of proportionality was not violated. However, due to the border line statistical-significance, we also put follow-up time as one of the covariates in regression analyses.

Results
Among the initial sampling population of the ILAS recruited between August 2011 and July 2014, 760 individuals had received comprehensive MRI modalities for CSVD detection and evaluation. We excluded nine individuals with incidentally found brain tumors and 16 individuals with problematic images due to head motion. The flow chart of the study population is shown in Fig. 1a.

In 735 individuals with eligible brain MRI images, 335 (45.6%) were categorized into robust (control group) and 249 (33.9%), 52 (7.1%), 61 (8.3%), and 38 (5.2%) were classified into CSVD type 1, type 2, type 3, and type 4 groups, respectively (Table 1). The demographics and imaging characteristics of the five groups are shown in Table 1. The Kruskal–Wallis nonparametric analyses showed that age and the presence of hypertension were significantly different among the five groups. Post-hoc analyses showed that individuals in each CSVD group were older than those in the control group. Nevertheless, individuals in CSVD types 1, 2, and 3 groups (however, not in CSVD type 4) had a higher prevalence of hypertension than those in the control group.

Additional characteristics of the CSVD markers in each classified subtype are presented in Table 1. In individuals with bleeding SVD, particularly type 3, the presence of severe WMH was also prominent (80.3% and 57.9% in types 3 and 4, respectively). The WMH volume ratios were significantly higher in each SVD group than in the control group. Lacunes were also present in bleeding SVD; 42.6% and 13.2% of types 3 and 4 had at least one lacune. The mean CSVD score and the distribution of the CSVD score category in each group are shown in Table 1. CSVD burden was different among the five groups; CSVD type 3 had the highest CSVD scores (Table 1). Notably, there was no participant with isolated lacune (with lacune but no severe WMH or CMB).

Mortality and CSVD types. We contacted all recruited participants by phone call and recorded their all-cause mortality status. The follow-up time did not differ between the groups (Table 1). During a mean follow-up of 5.7 (0.7) years, 62 (8.4%) died. The survival curves for each group are shown in Fig. 3. Table 2 demonstrates the incidence and HR of all-cause mortality in each group and the group comparisons. The results showed that after age and sex adjustment, only the CSVD type 4 group had a significantly higher all-cause mortality rate than the control group (HR 4.1, 95% confidence interval 1.4–12.1, \(p = 0.011\)), and the significance remained after adjustment for vascular risk factors (Table 2; HR 5.0, 95% confidence interval 1.6–15.3, \(p = 0.005\)). Since the ages were different between controls and CSVD groups, we also selected participants in the control group with age-matched with CSVD groups for the regression analyses (Table 2). The results showed that higher mortality
Mortality and CSVD burden. We also analyzed the associations between the all-cause mortality rate and CSVD burden using CSVD scores. There were 335, 275, 97, and 28 individuals with CSVD scores of 0, 1, 2, and 3, respectively. The results of Cox regression analyses did not show a dose-dependent relationship between the CSVD score and all-cause mortality, e.g. higher CSVD scores did not have a higher all-cause mortality rate (Table 2). In addition, after age and sex adjustment, only individuals with CSVD score 2 showed a significantly higher all-cause mortality rate compared to the those with CSVD score 0. In contrast, the CSVD score 1 and 3 groups had similar all-cause mortality rates as the CSVD score 0 group (Table 2).

Subgroup analysis of CSVD type 4. Within CSVD type 4, the associations between each demographic/neuroimaging characteristic and all-cause mortality are shown in Table 3. We did not find any factors associated with the all-cause mortality rate in the CSVD type 4 group. However, a higher number of CMBs (CMB ≥ 2) had a trend showing higher rate of mortality though statistically non-significant. Again, the CSVD burden was not associated with all-cause mortality in this subgroup.

Table 1. Comparisons between the control and four cerebral small vessel disease groups. BP, blood pressure; CMB, cerebral microbleed; CSVD, cerebral small vessel disease; HgbA1c, hemoglobin A1c; LDL, low-density lipoprotein cholesterol; MRI, magnetic resonance imaging; SD, standard deviation; SL, strictly lobar; WMH, white matter hyperintensities.  

| CSVD MRI markers | Non-bleeding CSVD | Bleeding CSVD |
|------------------|------------------|---------------|
|                  | Isolated WMH    | WMH with lacune | Mixed CMBs | SL CMBs | p value |
| Number           | 335 (45.6%)     | 249 (33.9%)    | 52 (7.1%)  | 61 (8.3%) | 38 (5.2%) |
| Age, years, mean (SD) | 57.9 (5.9) | 65.6 (8.1)    | 67.8 (9.2) | 67.1 (10.3) | 64.3 (9.0) | <0.001 |
| Sex, men, n (%)   | 132 (39.4%)     | 120 (48.2%)    | 26 (50.0%) | 31 (50.8%) | 17 (44.7%) | 0.241  |
| Follow-up time, year | 5.8 (0.8)     | 5.7 (0.6)     | 5.7 (0.8)  | 5.5 (0.7)  | 5.7 (0.6)  | 0.233  |

Vascular risk factors

| Hypertension, n (%) | 77 (23.0%) | 106 (42.6%) | 28 (53.8%) | 25 (41.0%) | 9 (23.7%) | <0.001 |
| Diabetes, n (%)     | 22 (6.6%)  | 46 (18.5%)  | 14 (26.9%) | 11 (18.0%) | 8 (21.1%) | <0.001 |
| Dyslipidemia, n (%) | 15 (4.5%)  | 14 (5.6%)   | 2 (2.8%)   | 7 (11.5%)  | 3 (7.9%)  | 0.241  |
| Cigarette smoking, n (%) | 67 (20.0%) | 75 (30.2%) | 18 (34.6%) | 17 (27.9%) | 9 (23.7%) | 0.029  |
| Systolic BP, mmHg, mean (SD) | 124.3 (14.7) | 132.0 (16.3) | 134.7 (19.3) | 132.6 (18.0) | 129.2 (21.3) | <0.001 |
| LDL, mg/dl, mean (SD) | 120.1 (33.6) | 117.3 (31.2) | 114.0 (27.8) | 113.6 (29.7) | 112.5 (27.4) | 0.319  |
| HgbA1c, %, mean (SD) | 5.8 (0.6)  | 6.0 (0.8)   | 6.6 (1.4)  | 6.1 (1.0)  | 6.0 (0.8)  | <0.001 |

CSVD score category, n (%)<0.001

| CSVD score category, n (%) | Control | Non-bleeding CSVD | Bleeding CSVD |
|---------------------------|---------|------------------|---------------|
|                           | 335 (100%) | 0 0 0 0 | 0 0 0 0 |
| 1                         | 0 249 (100%) | 0 0 0 0 | 10 (16.4%) | 16 (42.1%) |
| 2                         | 0 0 52 (100%) | 0 0 0 0 | 27 (44.3%) | 18 (47.4%) |
| 3                         | 0 0 0 0 | 24 (39.3%) | 4 (10.5%) |
Discussion
This study evaluated the association between CSVD and 5-year mortality in an asymptomatic (stroke-free and non-demented) middle-to-old aged population with two MRI marker-based classification methods, CSVD phenotypes, and burden. The results showed that among all CSVD types, the CSVD type 4 group was significantly associated with a higher rate of all-cause mortality independent of age, sex, and vascular risk factors. However, the CSVD burden measured by the simple CSVD score did not show a positive association with all-cause mortality. Only CSVD scores of 2, but not scores 1 and 3, were significantly and independently associated with a higher rate of all-cause mortality.

Our CSVD stratification scheme used three common MRI markers (Fig. 1b). The first step of the stratification scheme differentiated CSVD into bleeding and nonbleeding subtypes. Bleeding CSVDs were further stratified into different subtypes with CMBs of specific topographic features that have distinct underlying microvasculopathy: strictly lobar CMBs considering as CAA and mixed CMBs as arteriosclerosis/lipohyalinosis.
This study showed that the association between CSVD and 5-year all-cause mortality in the same population was also mediated by the CSVD types. In the CSVD type 4, the presentation of possible CAA as the underlying CSVD, at a mean age of 64.3 years, predicted a 4–fivefold higher all-cause mortality than in the non-CSVD group. Notably, these individuals lacked hemorrhagic stroke, the typical diagnostic criteria of CAA. These results indicate that this CSVD phenotyping method is able to differentiate survival outcomes in different underlying microvasculopathies, in an asymptomatic early stage of disease.

Previous studies have studied the mortality rate in patients with CMBs of different topographic patterns. They showed that mixed CMBs were associated with cardiovascular mortality, while lobar CMBs were associated with stroke-specific mortality compared to patients without CMBs; these results correspond with the notion that mixed CMBs are considered to reflect hypertensive arteriopathy and, therefore, systemic vascular disease, while lobar CMBs are indicative of CAA and primarily restricted to the brain. Although not provided in the literature, the strokes related to lobar CMB-associated mortality were probably intracerebral hemorrhage, a major clinical consequence of CAA. Therefore, we postulated that causes of mortality in the CSVD type 4 in this study were primarily hemorrhagic strokes. In the CSVD type 3 group, patients with mixed CMBs, had higher mortality than the control group (14.8% vs. 5.4%; Table 1), and the significance diminished after adjustment for age, sex, and vascular risk factors (Table 2). Our population had low cardiovascular risk or otherwise optimal medical control (Table 1), therefore, it might reduce the expected higher cardiovascular mortality in patients with CSVD type 3 in this study.

The initial studies of CSVD regarding their clinical outcome usually focused on only one MRI marker, particularly WMH or CMBs. However, individuals with CSVD usually have not only one but several coexisting MRI markers. Attempting to capture the overall effect of CSVD on the brain (CSVD burden), researchers have proposed the CSVD score, which is yielded by summing up the number of simultaneous MRI markers’ appearance in a person. The original CSVD score was computed by counting the presence of severe WMH, lacune, CMB, and dilated perivascular space (PVS) as an ordinal score of 0–4. Since PVS data are not routinely assessed as commonly as the other MRI markers in clinical or research settings, measurement without PVS is also generated as the simple CSVD score. Two previous large studies evaluated the association between CSVD score and mortality; one was in patients with acute ischemic stroke, and another was in a stroke-free and non-demented community-based population. In patients with acute ischemic stroke, only patients with the highest CSVD score (score 4) but not score 1, 2, or 3 had a significantly higher rate of all-cause mortality than those with CSVD score 0. In another study with an asymptomatic population, similar to the present study, only a CSVD score of 2 showed significant association with all-cause mortality but not the other CSVD scores. The lack of significant association between a higher CSVD score and all-cause mortality in an asymptomatic population might be explained by a low prevalence of mortality rate, small number of patients with a high CSVD score, or both. Therefore, it is suggested that the CSVD score might not be feasible to predict survival outcomes in asymptomatic or preclinical stages of CSVD. Notably, in the subgroup analyses of CSVD type 4, a higher CSVD score was also not correlated with a higher all-cause mortality. Among the three MRI markers, only the number of CMBs showed a positive association with all-cause mortality in the CSVD type 4 group.

### Table 3. Associations between each demographic and neuroimaging factor and all-cause mortality in cerebral small vessel disease type 4

| Demographic or Neuroimaging Factor | HR (95% CI) | p value |
|-----------------------------------|------------|---------|
| Age > 63 year-old*                 | 1.7 (0.4–7.8) | 0.476   |
| Sex, female                       | 1.0 (0.2–4.6) | 0.976   |
| Hypertension                      | 1.0 (0.2–5.4) | 0.959   |
| Diabetes Mellitus                 | 0.6 (0.1–5.1) | 0.640   |
| Hyperlipidemia                    | 1.2 (0.1–11.2) | 0.850   |
| Cigarette smoking                 | 1.6 (0.9–3.1) | 0.126   |
| The presence of severe WMH (n = 22) | 1.4 (0.3–7.8) | 0.686   |
| The presence of lacune (n = 5)    | 0.04 (0–2911.6) | 0.576   |
| The number of CMB ≥ 2 (n = 5)     | 4.0 (0.7–22.2) | 0.109   |
| **CSVD burden (versus CSVD score 1)** |            |         |
| CSVD score 2 (n = 18)             | 1.7 (0.3–9.3) | 0.537   |
| CSVD score 3 (n = 4)              | 0.04 (0–142,489.2) | 0.668   |

*The median age of the CSVD type 4 group.
again indicate that the nature (bleeding or nonbleeding; arteriosclerosis or CAA) might be more important than the number of CSVD markers when considering the clinical significance of CSVD.

There were limitations to the present study. First, we did not obtain information regarding the cause of death. Second, the present findings were from a community-based community that had no history of stroke or dementia. Whether the results could be generalized to other populations with evident clinical events requires further validation. Third, we did not assess and thus not include PVS in the evaluation of CSVD. Finally, the presence of severe WMH was defined as > 50th percentile of WMH/total intracranial volume ratio in our study population. We used automatic volumetric measurement to scale WMH since it offers a reliable and objective alternative to visual rating scales. The cut-off points of the WMH volume ratio for defining severe WMH in different populations might be different.

In conclusion, we showed that the MRI-based CSVD classification scheme could evaluate the 5-year all-cause mortality in stratified CSVD types in stroke-free and non-demented populations. This CSVD stratification method, which could identify asymptomatic individuals at risk of mortality, may be of clinical diagnostic value. The results showed that patients with CSVD type 4, possible CAA, had a higher all-cause mortality and provided insight into the early disease stage of CAA.

Data availability
Clinical and neuroimaging raw data are available from the corresponding author on request.

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Author contributions
W.J.L., C.P.C., L.N.P. collected clinical data, performed analysis and drafted the manuscript. K.H.C., P.L.L., and C.P.L. defined the MRI protocols and analyzed the neuroimaging data. C.P.L., K.H.C., L.K.C. and P.N.W.
critically revised the manuscript. WJ Lee performed statistical analysis. All authors have approved the submitted manuscript.

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**Competing interests**
The authors declare no competing interests.

**Additional information**
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