Orthostatic hypotension is not associated with small vessel disease progression or cognitive decline

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Introduction: Cerebral hypoperfusion is thought to play an important role in the etiology of cerebral small vessel disease (SVD). Orthostatic hypotension (OH) is assumed to be a cause of cerebral hypoperfusion by causing recurrent hypoperfusion episodes, and might thus be related to progression of SVD. Here, we investigated whether presence of OH is associated with the progression of SVD MRI-markers and cognitive decline over a time period of 9 years in a cohort of sporadic SVD patients.

Methods: This study included SVD patients from the RUN DMC study, a prospective longitudinal single-center cohort study. In total, 503 patients were included at baseline (2006), from whom 351 participated at first follow-up (2011), and 293 at second follow-up (2015). During all visits, patients underwent MRI and cognitive testing. Association between presence of OH at baseline and progression of SVD-markers on MRI and cognitive decline over time was estimated using linear mixed-effects models.

Results: Of the 503 patients who participated at baseline, 46 patients (9.1%) had OH. Cross-sectional analysis of the baseline data showed that OH was associated with higher white matter hyperintensity (WMH) volume ($\beta = 0.18, p = 0.03$), higher mean diffusivity (MD; $\beta = 0.02, p = 0.002$), and with presence of microbleeds (OR 3.12, 95% CI 1.16–8.68). Longitudinally, OH was however not associated with a progression of total WMH volume ($\beta = -0.17, p = 0.96$) or with higher MD ($\beta = -0.001, p = 0.49$). There was no association between OH and cognitive performance, both at baseline and over time.

Conclusion: In this longitudinal observational study, there was no evidence that presence of OH is associated with progression of SVD-markers or cognitive decline over time. Our findings indicate that OH may not be causally related to SVD progression over time.

Introduction

Cerebral small vessel disease (SVD) is a heterogeneous disease with high inter-individual variability in progression and clinical symptoms over time [1, 2]. Unraveling risk factors contributing to SVD-progression provides a better understanding of the pathophysiology of SVD. Current knowledge regarding risk factors of SVD is predominantly based on longitudinal cohort studies and includes traditional vascular risk factors (e.g., hypertension and diabetes). However, the variation in clinical course over time raises the possibility that other still unverified mechanisms beyond traditional risk factors may also play an eminent role in SVD progression.

Previous studies suggest that cerebral hypoperfusion play an important role in the etiology of SVD [3, 4]. Orthostatic hypotension (OH), characterized by a significant drop of blood pressure after postural change, occurs commonly in the elderly and has been proposed as a risk factor for cognitive decline and dementia [5–8]. OH is assumed to be a clinical marker for cerebral hypoperfusion, wherever the cerebral
autoregulation is impaired, by causing recurrent transient hypoperfusion episodes [9]. We therefore hypothesized that OH is causally related to progression of white matter hyperintensities (WMH) [10, 11], more specifically of those in the periventricular regions. Especially those areas are thought to be particularly vulnerable for hypoperfusion [12], supporting the hypothesis that spatial distribution of WMH may reflect differential etiologies [12, 13].

Current knowledge regarding relation between OH and SVD is limited to inconclusive cross-sectional studies [14–17], while longitudinal studies investigating progression of SVD over a long time-period are completely lacking. In this study, we aimed to investigate the effect of OH on progression of SVD-markers on MRI and cognitive decline over time over a time-period of 9 years.

Methods

Study population

This study is part of the RUN DMC study, a prospective cohort study among 503 individuals with SVD, that investigates risk factors and clinical consequences of SVD. The detailed study protocol has been published previously [18]. In short, we included community-dwelling consecutive patients, who were referred to our outpatient clinic and who underwent diagnostic cerebral imaging for different clinical reasons (e.g., TIA, cognitive/motor disturbances). Inclusion criteria were age between 50 and 85 years and presence of SVD on neuroimaging (i.e., WMH or lacunes). Of the 503 patients who participated at baseline in 2006, 361 underwent repeated MRI assessment at first follow-up in 2011, and 296 patients at second follow-up in 2015 (Fig. 1). Of those, 10 patients were excluded because of insufficient scan quality at first follow-up, and three at second follow-up, yielding a sample of 503 patients for neuroimaging analyses at baseline, 351 at first follow-up, and 293 at second follow-up. In total, 273 subjects underwent repeated cognitive and neuroimaging assessments at all three time-points and could be used for longitudinal analyses.

The Medical Review Ethics Committee region Arnhem-Nijmegen approved the study and all participants gave written informed consent.

Orthostatic hypotension

Systolic and diastolic blood pressures (SBP/DBP) were measured three times (separated by 1–2 min) after 5 min of supine rest. Then, after one minute of standing, the blood pressure was measured again, with the BP cuff positioned at heart level. Orthostatic hypotension was defined as either a SBP decrease of at least 20 mmHg and/or DBP decrease of at least 10 mmHg after one minute of standing up (difference in blood pressure measurements between the mean supine and the standing position) according to accepted criteria [19].

Vascular risk factors

Hypertension was defined as a SBP of ≥ 140 mmHg and/or DBP of ≥ 90 mmHg and/or the use of blood pressure lowering agents. Diabetes mellitus and hypercholesterolemia were considered to be present if the patient was taking oral glucose-lowering drugs or insulin or lipid-lowering drugs, respectively.

Use of OH-inducing drugs was defined as use of the following group of drugs: anti-depressants, alpha-blockers, beta-blockers, calcium-blockers, diuretics and angiotensin-system-blockers.

Cognition

Cognitive function was assessed by a standardized neuropsychological test battery during all waves of data collection and has been described in detail elsewhere [20]. Briefly, global cognitive function was evaluated by MMSE and a cognitive index was calculated covering the following cognitive domains: verbal and visuospatial memory, executive function, psychomotor speed, fluency and attention.
The cognitive index is a compound score that was calculated as the mean of the z-scores of the 1-letter subtask of the Verbal Series Attention Test, the Paper–Pencil Memory Scanning Task, the Verbal fluency Tasks, the mean of the reading subtask of the Stroop test, the mean of the Symbol–Digit Substitution Task, and the mean of the added score on the three learning trials of the Rey Auditory verbal Learning Test and the delayed recall of this last test.

The executive function was calculated as the interference score of the Stroop Test by dividing SAT-scores of the color-word task by the mean SAT-scores of the reading and color naming tasks of the Stroop Test, the verbal fluency task, and the SAT-scores of the Verbal Series Attention Test. Psychomotor speed was calculated as the mean of the z-scores of the 1-letter subtask of the Paper-Pencil Memory Scanning Task, the reading and color naming tasks of the Stroop Test, and the Symbol Digit Substitution Task.

MRI protocol

Images were acquired at three time points on a 1.5-Tesla MRI (2006: Siemens, Magnetom Sonata; 2011 and 2015: Siemens, Magnetom Avanto) and included the following whole brain scans: 3D T1 MRAGE imaging (isotropic voxel size 1.0 mm\(^3\)); FLAIR sequences (2006: voxel size 0.5 \(\times\) 0.5 \(\times\) 5.0 mm, interslice gap 1.0 mm; 2011 and 2015: voxel size 0.5 \(\times\) 0.5 \(\times\) 2.5 mm; interslice gap 0.5 mm); and a transversal T2*-weighted gradient echo sequence (2006: isotropic voxel size 2.5 mm\(^3\); 4 unweighted scans, 30 diffusion weighted scans at \(b = 900\) s/mm\(^2\); 2011 and 2015: 8 unweighted scans, 60 diffusion weighted scans at \(b = 900\) s/mm\(^2\)). The same head coil was used at all 3 time points. Full acquisition details have been described previously [18].

Brain volumetry

Gray matter volume (GMV), white matter volume (WMV) and CSF volumes were assessed by using Statistical Parametric Mapping, SPM12 (https://www.fil.ion.ucl.ac.uk/spm) unified segmentation routines on the T1 MRAGE images corrected for WMH, as has been described in detail previously [21]. Intracranial volume (ICV) was calculated by summing GMV, WMV, and CSF volumes, and total brain volume. To correct for inter-scan effects, we corrected for differences in ICV between baseline and follow-up. We normalized all volumes to baseline ICV to account for head size.

MRI-markers of small vessel disease

MRI-markers of SVD were rated according to the Standards for Reporting Vascular Changes on Neuroimaging (STRIVE) criteria [22]. WMH volumes were calculated by a semiautomatic WMH segmentation method, which has been described in detail elsewhere [23]. Segmentations were visually checked for segmentation errors by one trained rater, blinded for clinical data. In addition, we distinguished periventricular WMH (P WMH) and deep WMH (D WMH) using FMRIB Software Library (FSL; www.fmrib.ox.ac.uk/fsl). More specifically, we generated ventricle masks and calculated the geometrical distance from this mask in each voxel using Brain Intensity Abnormality Classification Algorithm (BI ANCA) and the command distancemap [12, 24, 25]. Based on previously validated criteria in literature [12], we considered voxels within 10 mm distance of the ventricles as PWMH and calculated the corresponding volumes. WMH volumes were corrected for interscan differences in ICV and then normalized to baseline ICV.

Number of lacunes and microbleeds were rated manually on FLAIR/T1-weighted and T2*-weighted MRI scans according to the STRIVE criteria by two trained raters blinded for clinical data. Follow-up FLAIR images were resliced to match the baseline scans to prevent differences in partial volume effects between baseline and follow-up scans. Inter-rater and interrater reliability were excellent [24].

Diffusion tensor imaging (DTI) analysis has been described elsewhere [26]. Diffusion data were first preprocessed by using local algorithm [27] to correct for cardiac and head motion as well as eddy currents. Using DTIFit within the Functional MRI of the Brain diffusion toolbox, we created fractional anisotropy (FA) and mean diffusivity (MD) images, which were imported into TBSS pipeline [28]. We then calculated the mean MD within the skeleton for each subjects.

Statistical analysis

First, we identified differences in baseline clinical and MRI-characteristics between patients with and without OH, by using chi-square test or Mann-Whitney test, where appropriate.

Second, we investigated the association between presence of OH and cognitive index, executive function, psychomotor speed, as well as MRI-characteristics of SVD, by using multivariable linear regression for the continuous outcome variables and logistic regression for dichotomous outcome variables, adjusting for age, sex, presence of hypertension, hypercholesterolemia and use of any OH-inducing drugs.

Finally, we studied the effect of baseline presence of OH on the change of cognition (hereby again focusing on cognitive index, executive function, and psychomotor speed) and progression of WMH (stratified by its location: total WMH, DWMH and PWMH) over time. We fitted linear-mixed effects (LME) models via R package “lme4”. LME models allow for the simultaneous modeling of fixed (population-average) and random (subject-specific) effects, allowing us to investigate change in outcome variables in the entire population while accounting for individual differences in rates of WMH progression. We specified the following fixed-effects: presence of OH, baseline age, sex, hypertension, usage of any OH-inducing drugs, an interaction term between follow-up time and presence of OH/absolute SBP changes/absolute DBP changes. We included random intercepts and slopes of time per subject. To evaluate a possible quadratic relationship, indicating nonlinear progression of SVD, we compared model fit between the full model and the full model with a quadratic term during follow-up. We evaluated change in Aikake information criterion (AIC) by one-way ANOVA. Correction for multiple comparisons was done by FDR correction. All statistical analyses were carried out in R Programming Language version 4.1.1.

Results

Of the 503 patients who participated at baseline, 46 patients (9.1%) had OH. Differences in baseline characteristics between the two groups are presented in Table 1. Patients with OH were older than patients without OH (mean ± SD 76.5 years ± SD 8.9, \(p = 0.01\)) and showed worse cognitive performance at baseline compared to non-OH patients (median cognitive index: 0.01 vs –0.20, \(p = 0.04\)). Patients with OH had also a significant higher prevalence of hypertension (\(p = 0.02\)) and hypercholesterolemia (\(p = 0.002\)).

OH and small vessel disease burden

OH was cross-sectionally associated with higher WMH (\(\beta = 0.18, p = 0.03\)), DM WMH (\(\beta = 0.24, p = 0.02\)) and PWMH (\(\beta = 0.17, p = 0.04\)) volumes at baseline, adjusted for sex, age, hypertension, hypercholesterolemia and OH-inducing drugs. Patients with OH had a higher MD (\(\beta = 0.02, p = 0.002\)) and were more likely to have microbleeds compared to patients without OH (OR 2.37 95% CI 1.16–4.68). Table 2 shows the associations between having OH and MRI-characteristics of SVD.

OH and progression of SVD

Of the 46 OH-patients at baseline, 28 patients participated at first follow-up and 19 at second follow-up. Mean follow-up duration was 5.3 (SD 0.2) years until first follow-up, and 8.7 (SD 0.2) years until the second follow-up.
Table 1
Baseline characteristics of the patients with and without OH.

| Clinical Characteristics | Total sample | Without OH | With OH | p value |
|--------------------------|--------------|------------|---------|---------|
|                        | n = 503     | n = 457    | n = 46  |         |
| Demographics            |             |            |         |         |
| Sex (male,%)            | 284 (56.5)  | 257 (56.2) | 27 (58.7)| 0.87    |
| Age, mean ± SD          | 65.6 ± 8.8  | 65.3 ± 8.6 | 68.9 ± 7.6| 0.01    |
| MMSE                    | 28           | 29         | 28      | 0.11    |
| (IQR)                   | (27-29)      | (27-29)    | (26-29) |         |
| Cognitive index         |              |            |         |         |
| -0.04 (-0.6-0.5)        |              | -0.01 ± 0.8| -0.18 ± 0.7| 0.11    |
| Executive function, mean ± SD | 0.0 ± 0.8 | 0.02 ± 0.9 | 0.23 ± 0.9| 0.04    |
| Psychomotor speed, mean ± SD | 0.9       | 0.9        |         |         |
| OH-inducing drugs       |              |            |         |         |
| Anti-depressants, n (%)  | 62 (12.3)    | 55 (12.0)  | 7 (15.2)| 0.70    |
| Alpha-blockers, n (%)    | 2 (0.4)      | 2 (0.4)    | 0 (0.0)| 1.00    |
| Beta-blockers, n (%)     | 181 (36.0)   | 158 (34.6) | 23 (50.0)| 0.06    |
| Calcium-blockers, n (%)  | 51 (10.1)    | 43 (9.4)   | 8 (17.4)| 0.15    |
| Diuretics, n (%)         | 124 (24.7)   | 111 (24.3) | 13 (28.3)| 0.68    |
| Angiotensin-system blocker, n (%) | 160 (31.8) | 140 (30.6) | 20 (43.5) | 0.11    |
| Any OH-inducing drugs, n (%) | 303 (60.2) | 269 (58.9) | 34 (73.9)| 0.07    |
| Risk factors SVD        |              |            |         |         |
| Ever smoking, n (%)      | 353 (70.2)   | 317 (69.4) | 36 (78.3)| 0.28    |
| Hypertension, n (%)      | 369 (73.4)   | 328 (71.8) | 41 (89.1)| 0.02    |
| Diabetes, n (%)          | 66 (13.1)    | 58 (12.7)  | 8 (17.4)| 0.50    |
| Hypercholesterolemia, n (%) | 237 (47.1) | 205 (44.9) | 32 (69.6)| 0.002   |
| BMI, kg/m², mean ± SD    | 27.2 ± 4.1   | 27.2 ± 4.1 | 27.0 ± 4.2| 0.53    |
| MRI Characteristics      |              |            |         |         |
| WMH (ml; IQR)           | 8.6          | 3.3        | 8.9     | <0.001  |
| (1.2-11.5)              | (1.2-10.5)   | (3.3-17.6)|         |         |
| DWMI (ml; IQR)          | 2.9          | 0.9        | 2.5     | 0.002   |
| (0.3-2.9)               | (0.3-2.6)    | (0.6-6.8)|         |         |
| PWMI (ml; IQR)          | 5.7          | 2.27       | 6.02    | <0.001  |
| (0.8-8.0)               | (0.8-7.1)    | (2.6-13.0)|         |         |
| GMV (ml; SD)            | 606.2 ± 87   | 607.5 ± 90 | 594 ± 90| 0.09    |
| WMV (ml; SD)            | 545.7 ± 94   | 546.2 ± 94 | 439.4 ± 94| 0.07    |
| MOWhite matter          | 46.0         | 46.7       | 34.6    |         |
| Presence of microbleeds, n (%) | 83 (13.6) | 68 (15.0) | 15 (32.6)| 0.002   |
| Presence of lacunes, n (%) | 132 (26.2) | 111 (24.3) | 21 (45.7)| 0.002   |

Table 2
Multivariable cross-sectional association between presence of OH and cognitive and radiological correlates in SVD.

| Cognition | β (95% CI) | p value |
|-----------|------------|---------|
| Cognitive index | -0.056 (-0.25-0.14) | 0.57    |
| Executive function | -0.002 (-0.21-0.21) | 0.99    |
| Psychomotor speed | -0.04 (-0.27-0.19) | 0.75    |
| MRI Characteristics |             |         |
| WMH*       | 0.18 (0.02-0.34) | 0.03    |
| PWMH*      | 0.24 (0.04-0.43) | 0.02    |
| PWMI*      | 0.17 (0.01-0.33) | 0.04    |
| GMV        | 3.59 (-8.34-15.52) | 0.56    |
| WM         | 0.61 (-0.02-0.01) | 0.41    |
| MD white matter* | 0.02 (0.01-0.03) | 0.002  |
| OR (95% CI) |             |         |
| Presence of microbleeds* | 2.37 (1.16-4.68) | 0.02    |
| Presence of lacunes | 1.88 (0.97-3.61) | 0.06    |

Data represents β estimates (95% CI), OR values (95% CI) or p-values. *Log transformed. All models were adjusted for sex, age, hypertension, hypercholesterolemia, and use of any OH inducing drugs. 8 non-OH patients and 1 OH-case were excluded due to insufficient scan quality. 4 patients without OH were excluded for missing data regarding microbleeds. OH: Orthostatic Hypotension, WMH: White Matter Hyperintensities, DWMH: Deep White Matter Hyperintensities, PWMH: Periventricular White Matter Hyperintensities, GMV: gray Matter Volume, WMV: White Matter Volume, MD: Mean Diffusivity.

Table 2. Furthermore, no differences in change of cognitive index, executive function or psychomotor speed was observed over time between patients with and without OH (Table 4).

Discussion

We found that the presence of OH was associated with higher WMH volume, impaired structural integrity and higher prevalence of microbleeds at the cross sectional level. However, OH was not associated with progression of MRI-markers of SVD over time or with cognitive decline in individuals with sporadic small vessel disease. Similar longitudinal findings were found when analysing the effect of absolute changes in SBP and DBP, between supine and standing position, rather than the dichotomous definition of OH.

The vascular hypothesis of autonomic dysfunction proposes that recurrent episodic cerebral hypoperfusion causes vascular oxidative stress, which results in alterations in cerebral vasculature (e.g. impaired vasodilation) [29]. This could lead to chronic ischaemia that on its turn may partly explain the occurrence and progression of WMH lesions [30, 31]. Although blood pressure variability is increasingly recognized to be associated with SVD-burden [31, 32], though other studies have failed to find these associations [14, 16].

We found OH not to be associated with WMH progression over time nor with cognitive decline. This might be explained by the assumption that OH does not lead to harmful (episodic) reductions in cerebral perfusion, due to compensatory cerebral autoregulation [14, 29]. OH is defined by a blood pressure reduction that persists after 1 min (or more) of standing. Within that timeframe, there is sufficient time for cerebral autoregulation to adapt to this reduction in blood pressure and return CBF to normal [34, 14, 29]. Indeed, it is only the initial phase of the postural response where blood pressure changes occur too fast for cerebral autoregulation to adapt [5]. Therefore, a likely possibility is that OH is only detrimental in case of impaired cerebral autoregulation [14]. Another explanation could be that not per se the severity of drop in blood pressure that matters, but rather the duration of delayed recovery
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Table 3
Longitudinal association between OH and progression of white matter hyperintensities, stratified by location.

|                | Total WMH | p value | DWIH | p value | PWMH | p value | MD white matter | p value |
|----------------|-----------|---------|------|---------|------|---------|-----------------|---------|
| Baseline age (years) | 0.458     | <0.001  | 0.128 | <0.001  | 0.330| <0.001  | 0.002           | <0.001  |
| Sex (male)      | -0.078    | 0.99    | -0.240| 0.75    | 0.433| 0.69    | -0.003          | 0.69    |
| Hypertension    | 1.479     | 0.60    | 0.320 | 0.82    | 0.79 | 0.80    | 0.006           | 0.60    |
| OH             | 3.633     | 0.98    | 1.368 | 0.98    | 2.290| 0.98    | 0.018           | 0.98    |
| FUP-time*age   | 0.041     | <0.001  | 0.002 | 0.46    | 0.027| <0.001  | 5.172e05        | 0.26    |
| FUP-time*OH    | -0.061    | 0.97    | -0.035| 0.97    | -0.101| 0.97   | 0.001           | 0.97    |
| FUP-time*SBP   | 0.002     | 0.66    | 0.001 | 0.66    | -0.002| 0.58   | 1.181e05        | 0.66    |
| FUP-time*DBP   | 0.004     | 0.97    | -0.001| 0.97    | 0.002 | 0.97   | 8.554e06        | 9.97    |

Fixed effects results from linear mixed-effects models explaining progression of total WMH, DWMH, PWMH and MD. Data represents β-estimates (95% CI) or p-values. We studied the interaction between FUP time and OH and postural SBP/DBP changes, in the different models. In all models a quadratic term was included. WMH: white matter hyperintensities; DWMH: deep white matter hyperintensities; PWMH: periventricular white matter hyperintensities; MD: Mean Diffusivity; FUP: Follow-Up; OH: Orthostatic Hypotension; SBP: Systolic Blood Pressure; DBP: Diastolic Blood Pressure.

Table 4
Longitudinal association between OH and change of cognitive index, executive function and psychomotor speed.

|                 | Cognitive index | p value | Executive function | p value | Psychomotor speed | p value |
|-----------------|-----------------|---------|--------------------|---------|-------------------|---------|
| Baseline age (years) | -0.040       | <0.001  | -0.037            | <0.001  | -0.043            | <0.001  |
| Sex (male)      | -0.037       | 0.69    | 0.047             | 0.69    | -0.077            | 0.69    |
| Hypertension    | 0.004        | 0.99    | -0.050            | 0.82    | -0.017            | 0.95    |
| OH             | -0.074       | 0.98    | -0.041            | 0.98    | -0.054            | 0.98    |
| FUP-time*age   | -0.003       | <0.001  | -0.003            | <0.001  | -0.002            | <0.001  |
| FUP-time*OH    | 0.022        | 0.47    | 0.005             | 0.97    | 0.030             | 0.47    |
| FUP-time*SBP   | 3.32e04      | 0.58    | 3.68e06           | 0.66    | 3.987e04          | 0.58    |
| FUP-time*DBP   | 5.840e04     | 0.97    | -4.322e04         | 0.97    | 8.859e04          | 0.97    |

Fixed effects results from linear mixed-effects models explaining change of cognitive index, executive function and psychomotor speed over time. Data represents β-estimates with (95% CI) or p-values. We studied the interaction between FUP time and OH and postural SBP/DBP changes, in the different models. In all models a quadratic term was included. WMH: white matter hyperintensities; DWMH: deep white matter hyperintensities; PWMH: periventricular white matter hyperintensities; MD: Mean Diffusivity; FUP: Follow-Up; OH: Orthostatic Hypotension; SBP: Systolic Blood Pressure; DBP: Diastolic Blood Pressure.

from OH that mediates the pathologic effect of OH [17].

In contrast to our results, there is some support for clinical evidence of OH earlier in life and SVD pathological changes in the brain vasculature in a postmortem study [30]. However, ‘reversed causality’ may have played a role in this study as they included only dementia and/or Parkinson patients. The neuronal loss observed during these diseases may affect brain structures involved in blood pressure control leading to autonomic dysfunction. In addition, recent evidence suggests that incomplete recovery of BP at 1 min of standing was related to cognitive decline and mortality in people with clinical Alzheimer’s disease [35].

Regional separation of WMH is recently reported to be an imported avenue in SVD research, given that PWMH and DWMH have not only been associated with different cardiovascular risk factors, but also with different underlying microstructural and neuropathological properties, suggesting at least partially distinct pathogenic mechanisms [12, 13, 30]. DWMH has been more attributed to SVD related hypoxic/ischemic damage, while PWMH is thought to be more related to hemodynamic factors (e.g. hypoperfusion) with an accompanying inflammatory component [12, 13]. However, we found no significant differences of the effect of OH on the progression of these two distinct regional white matter lesions.

This is the first study providing longitudinal data regarding the effect of OH on SVD progression over a 9-year follow-up with a large sample size of SVD patients. By performing serial extensive MRI sequences at three time points, we were able to examine the progression of WMH by location, over time. Furthermore, the 10 mm distance rule we used for WMH stratification gives most comparable results to the current literature in terms of showing the best separation in risk factors and patho-

logical features between PWMH and DWMH [12].

Nevertheless, some limitations need to be addressed when interpreting the results of this study. First, we had a relatively small number of patients with OH. This attrition bias due to loss of participants may have influenced our results. Second, we were not able to disentangle patients with possibly more severe (symptomatic) OH from those with (asymptomatic) OH only based on blood pressure decrease during our research assessment. OH accompanied by symptoms as dizziness, syncope and falls may indicate more severe OH with, given the symptoms,
possible cerebral hypoperfusion. However, for the present study these clinical data were not available. Patients with more severe OH could also be identified by applying a more strict definition of OH, like often applied in multiple system atrophy (SBP drop by 30 mmHg or DBP drop by 15 mmHg within 3 min) [36]. Unfortunately, we would then reach even smaller numbers of patients with OH for statistical analyses.

In conclusion, in this longitudinal observational study, there was no evidence that presence of OH or SBP/DBP postural variability is associated with more severe WMH progression over time. Our data indicate that OH may not be causally related to SVD progression over time: orthostatic hypotension as a new cardiovascular risk factor, J. Am. Coll. Cardiol. 40 (2002) 133-141.

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