Association between Brachial-Ankle Arterial Stiffness and Cerebral Small Vessel Disease Load in Patients with Acute Non-Cardioembolic Ischemic Stroke

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Abstract

Background: Increased arterial stiffness may lead to vessel damage and atherosclerosis, with the former occurring more frequently in the microvessels of end organs. We investigated the association between arterial stiffness, measured via brachial-ankle pulse wave velocity (baPWV), and cerebral small vessel disease (CSVD), visualized on magnetic resonance imaging (MRI) as white matter hyperintensities (WMHs), lacunar infarctions, and cerebral microbleeds (CMBs).

Methods: This study included 88 patients with ischemic stroke who were diagnosed with non-cardioembolic acute cerebral infarction. All patients underwent baPWV and brain MRI. The number of CMBs and lacunar infarctions was evaluated. The locations of CMBs and lacunar infarctions were divided into infratentorial, lobar, and deep regions, including the basal ganglia, thalamus, and internal capsule. Moreover, WMHs were divided into deep and periventricular regions, and their severity was assessed.

Results: Increased baPWV was associated with lacunar infarctions and WMHs (p<0.05), but not with CMBs. On multivariate analysis, the association between baPWV and lacunar infarctions in the deep regions was stronger than that between baPWV and the other imaging markers of CSVD (p<0.01).

Conclusions: These findings suggest that increased arterial stiffness may be associated with CSVD severity. Moreover, lacunar infarctions in the deep regions, such as the basal ganglia and thalamus, were related to arterial stiffness.

Keywords: Ankle brachial index, Cerebral small vessel diseases, Magnetic resonance imaging, White matter, Lacunar infarcts

1. INTRODUCTION

The arterial pulse wave velocity (PWV), a known independent predictor of cardiovascular disease, is widely used in clinical settings to determine changes in blood vessels caused by atherosclerosis. PWV increases as arteriosclerosis worsens, and increased arterial stiffness exacerbates vascular damage and atherosclerosis. It mainly affects the small vessels of the end organ. The brachial-ankle pulse wave velocity (baPWV), most commonly used for its simple and reproducible measurements, reflects the central and peripheral arterial stiffness.

Cerebral small vessel disease (CSVD) is a major form of ischemic stroke. Although its pathogenesis is unclear, arterial stiffness associated with high blood pressure plays a major role in its pathophysiology. The markers of CSVD include lacunar infarction, white matter hyperintensity (WMH), cerebral microbleed (CMB), and dilated perivascular space (PVS). Although increased arterial stiffness has been linked to lacunar infarction, WMH, and CMB, previous studies have reported inconsistent results.

To understand their relation, we compared the severity of imaging markers of CSVD, such as WMH, lacunar infarction, and CMB, visualized using brain magnetic resonance imaging (MRI), and arterial stiffness, measured using baPWV, in the patients with acute ischemic stroke.
2. MATERIALS AND METHODS

2.1. Materials

Between March 2018 and July 2020, 628 consecutive patients with acute ischemic stroke who were admitted within 7 days of symptom onset were enrolled. All patients were hospitalized with neurological symptoms and were diagnosed with cerebral infarction on brain MRI. A total of 193 patients with embolic cerebral infarction of cardiac origin, such as atrial fibrillation, a trial flutter, patent foramen ovale, artificial heart valve, mitral valve stenosis, thrombus in the left atrium, or atrial appendage, were excluded because of the difficulty in measuring baPWV. A total of 233 patients who did not have their baPWV measured and 62 patients with cardiac disorders, such as congestive heart failure, myocardial infarction within 6 months, and dilated cardiomyopathy, were also excluded. We further excluded 52 patients who had poor quality MRI data and missing data in medical records. A total of 88 patients were finally enrolled in the study.

2.2. Methods

The patients' clinical data were obtained through medical records and interviews, including their medical history, past history, and risk factors for cerebrovascular disease. Blood tests, electrocardiogram, and echocardiography were also conducted. Hypertension was defined as a systolic blood pressure of >140 mmHg or a diastolic blood pressure of >90 mmHg after the acute period of stroke and current use of antihypertensive medication, regardless of the blood pressure. Diabetes mellitus was defined as diabetes treatment, fasting glucose level >126 mg/dL after hospitalization, oral glucose tolerance test result >200 mg/dL at 2 hours, or hemoglobin A1c level >6.5 %A1c. Hyperlipidemia was defined as fasting total cholesterol level >240 mg/dL, low-density lipoprotein level >160 mg/dL, or use of lipid-lowering medication.

Arterial stiffness and the severity of CSVD in the brain were evaluated using baPWV and brain MRI, respectively. After the patient had rested in a supine position for at least 5 min, baPWV was measured using a volumeplethysmography apparatus (VP-1000; Collin, Co., Ltd., Komaki, Japan), which simultaneously records the PWV, arterial blood pressure, electrocardiogram, and heart sounds at the upper arms and ankles on both sides. The baPWV was calculated using time-phase analysis between the right arm and volume waveforms at both ankles. The distance between the right brachium and the ankle was estimated according to the body height. The average baPWV values obtained from both sides were then used for further analysis. The patients were then divided into tertiles depending on their baPWV: there were 29 patients with a baPWV of <1,699 cm/s, 30 patients with a baPWV of 1,699 - 2,023 cm/s, and 29 patients with a baPWV of >2,023 cm/s.

Axial T1- and T2-weighted, fluid-attenuated inversion recovery, and gradient-echo T2*-weighted images were collected using a 1.5-T MRI (GE HealthCare, Waukesha, WI). The number of CMBs and lacunar infarctions was identified using the T1/T2-weighted images, respectively. Using the Microbleed Anatomical Rating Scale, the locations of the CMBs and lacunar infarctions were reclassified into infratentorial, lobar, and deep regions. WMHs were evaluated using the Clinical Research for Dementia of South Korea visual rating score. Specifically, periventricular WMHs were evaluated on P1 (cap and band, <5 mm), P2 (cap or band, 5-10 mm), and P3 (cap or band, >10 mm). Moreover, deep WMHs were evaluated as D1 (<10 mm), D2 (10-25 mm), and D3 (>25 mm) based on the maximum diameter. WMHs were categorized asperiventricular, deep, and mixed based on their location and were graded as minimal, moderate, and severe (1, 2, and 3 points, respectively). The assessments were performed by a neurologist.

2.3. Statistics

For statistical analyses, patients were divided into the tertiles according to baPWV. Continuous variables were analyzed using the Kruskal-Wallis test, and categorical variables were analyzed using linear-by-linear association. Multivariate analysis was performed to determine the variables associated with baPWV. First, we adjusted for age, sex, and systolic blood pressure (model 1). In model 2, we additionally adjusted for heart rate, hemoglobin level, erythrocyte sedimentation rate, and glomerular filtration rate. Statistical Package for the Social Sciences Version 19.0 (SPSS Inc., Chicago, IL, USA) was used. A p-value of <0.05 was considered significant.
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3. RESULTS

3.1. Clinical Characteristics According to baPWV Differences

Among the three groups divided by tertiles, sex, age, systolic and diastolic blood pressure, pulse pressure, heart rate, hemoglobin level, erythrocyte sedimentation rate, and glomerular filtration rate were significantly different (p<0.05). However, factors such as presence of hypertension, diabetes mellitus, and hyperlipidemia; smoking; alcohol consumption; prior history of stroke or coronary heart disease; serum creatinine level; fasting glucose level; total cholesterol level; and body mass index were not different among the groups (Table 1).

Table 1: Baseline demographics and characteristics of patients with acute ischemic stroke according to pulse wave velocity

|                         | baPWV (<1699 cm/s) (n=29) | baPWV (1699-2023 cm/s) (n=30) | baPWV (>2023 cm/s) (n=29) | p-value |
|-------------------------|---------------------------|-------------------------------|---------------------------|---------|
| Sex (Male/Female)       | 23/6                      | 21/9                          | 14/15                     | 0.038   |
| Age (years)             | 58.8±10.2                 | 70.6±9.7                      | 70.6±8.5                  | <0.001  |
| Hypertension, n (%)     | 13 (44.8)                 | 14 (46.7)                     | 15 (51.7)                 | 0.601   |
| Diabetes mellitus, n (%)| 3 (10.3)                  | 10 (33.3)                     | 9 (31.0)                  | 0.070   |
| Hyperlipidemia, n (%)   | 5 (17.2)                  | 4 (13.3)                      | 4 (13.8)                  | 0.713   |
| Current Smoking, n (%)  | 11 (37.9)                 | 9 (30.0)                      | 6 (20.7)                  | 0.152   |
| Alcohol, n (%)          | 11 (37.9)                 | 13 (43.3)                     | 10 (34.5)                 | 0.789   |
| Prior stroke, n (%)     | 4 (13.8)                  | 3 (10.0)                      | 5 (17.2)                  | 0.704   |
| Coronary artery disease, n (%) | 2 (6.9) | 2 (6.7) | 2 (6.9) | 0.971 |
| Systolic BP (mmHg)      | 133.3±12.4                | 146.0±22.4                    | 162.8±26.4                | <0.001  |
| Pulse pressure (mmHg)   | 51.9±9.3                  | 62.1±13.4                     | 66.9±15.5                 | <0.001  |
| Diastolic BP (mmHg)     | 81.4±8.4                  | 87.0±22.9                     | 91.8±10.6                 | 0.004   |
| Heart rate (beats/min)  | 65.0±8.5                  | 63.3±15.3                     | 72.5±12.1                 | 0.021   |
| Hemoglobin (g/dL)       | 14.3±1.4                  | 13.4±1.7                      | 13.1±1.2                  | 0.017   |
| ESR (mm/hr)             | 13.3±9.4                  | 20.8±21.7                     | 28.2±24.6                 | 0.017   |
| Creatine (mg/dL)        | 0.8±0.2                   | 1.0±1.1                       | 1.01±0.7                  | 0.510   |
| GFR (mL/min)            | 101.8±23.7                | 98.5±23.3                     | 80.8±37.2                 | 0.015   |
| Fasting blood sugar (mg/dL) | 105.7±33.3 | 125.3±102.8 | 108.4±21.6 | 0.282 |
| Total cholesterol (mg/dL) | 193.0±45.4 | 174.1±63.1 | 192.5±42.0 | 0.412 |
| Body mass index (kg/m²) | 23.6±2.0                  | 23.9±2.7                      | 22.9±2.8                  | 0.645   |

Values are presented as mean±standard deviation or number (%).

baPWV: brachial-ankle pulse wave velocity; BP: blood pressure; ESR, erythrocyte sedimentation rate; GFR, glomerular filtration rate.

3.2. Relation between baPWV and Brain Imaging Markers of CSVD

There were no significant differences in the degree and location of CMBs among the three groups divided by tertiles (Table 2). There were significant differences in the total number of lacunar infarctions and deep lacunar infarctions (p<0.01). Further, there was no difference in the number of lacunar infarctions in the infarctorial and lobar areas (Table 2). WMHs were significantly more severe in the higher baPWV group, both in the deep and periventricular regions (p<0.05). There was also a significant difference in the total WMH grade (p=0.042; Table 2). Therefore, baPWV was associated with WMHs and the severity of lacunar infarction, but not with CMBs.

3.3. Multivariate Analysis of the Association between CSVD and baPWV

Multivariate analysis was performed by adjusting for variable factors. In model 1, baPWV was significantly associated with the total number of lacunar infarctions, deep and lobar lacunar infarctions, and periventricular WMHs (p<0.05; Table 3). However, in model 2, which adjusted for more variables, only deep lacunar infarction was significantly associated with baPWV (p<0.01; Table 3).
Table 2: Imaging markers of small vessel disease according to tertiles between brachial-ankle pulse wave velocity

|                    | baPWV (≤1699 cm/s) (n=29) | baPWV (1699-2023 cm/s) (n=30) | baPWV (>2023 cm/s) (n=29) | p-value |
|--------------------|---------------------------|--------------------------------|---------------------------|---------|
| Cerebral microbleeds |                           |                                |                           |         |
| Total              | 1.24±4.62                 | 1.50±4.02                      | 1.24±2.53                 | 0.488   |
| Infratentorial     | 0.14±0.58                 | 0.23±0.57                      | 0.28±0.92                 | 0.531   |
| Deep               | 0.69±2.97                 | 1.00±3.64                      | 0.69±1.56                 | 0.463   |
| Lobar              | 0.41±1.15                 | 0.27±0.74                      | 0.24±0.69                 | 0.559   |
| Lacunar infarctions|                           |                                |                           |         |
| Total              | 1.66±2.41                 | 2.10±1.81                      | 4.10±3.71                 | 0.001   |
| Infratentorial     | 0.17±0.76                 | 0.33±0.61                      | 0.41±0.87                 | 0.106   |
| Deep               | 1.03±1.80                 | 1.50±1.68                      | 3.03±2.96                 | 0.001   |
| Lobar              | 0.45±0.95                 | 0.23±0.50                      | 0.66±1.14                 | 0.349   |
| WM hyperintensities|                           |                                |                           |         |
| Periventricular WM | 1.03±0.50                 | 1.23±0.43                      | 1.48±0.57                 | 0.008   |
| Deep WM            | 1.07±0.53                 | 1.33±0.55                      | 1.48±0.51                 | 0.016   |
| Total grade        | 1.07±0.47                 | 1.17±0.38                      | 1.38±0.56                 | 0.042   |

Values are presented as mean ± standard deviation.

baPWV, brachial-ankle pulse wave velocity; WM, white matter.

Table 3: Multivariable analysis on the relationship between brachial-ankle pulse wave velocity and cerebral small vessel disease load

|                    | baPWV, Model 1 |                                | baPWV, Model 2 |                                |
|--------------------|----------------|--------------------------------|----------------|--------------------------------|
|                    | β              | p-value                        | β              | p-value                        |
| Cerebral microbleeds|                |                                |                |                                |
| Total              | 0.005          | 0.957                          | -0.001         | -0.990                         |
| Infratentorial     | 0.093          | 0.270                          | 0.120          | 0.206                          |
| Deep               | 0.000          | 1.000                          | -0.006         | 0.947                          |
| Lobar              | -0.055         | 0.515                          | -0.078         | 0.413                          |
| Lacunar infarcts   |                |                                |                |                                |
| Total              | 0.226          | 0.007                          | 0.175          | 0.077                          |
| Infratentorial     | -0.012         | 0.893                          | -0.017         | 0.856                          |
| Deep               | 0.265          | 0.001                          | 0.304          | 0.001                          |
| Lobar              | 0.181          | 0.035                          | 0.085          | 0.408                          |
| WM hyperintensities|                |                                |                |                                |
| Periventricular WM | 0.179          | 0.044                          | 0.182          | 0.068                          |
| Deep WM            | 0.117          | 0.203                          | 0.114          | 0.297                          |
| Total grade        | 0.128          | 0.150                          | 0.099          | 0.332                          |

baPWV, brachial-ankle pulse wave velocity; WM, white matter. Values represent the standardized regression coefficients (β). Model 1: Adjusted for age, sex, and systolic blood pressure. Model 2: adjusted for age, sex, systolic blood pressure, heart rate, hemoglobin level, erythrocyte sedimentation rate, and glomerular filtration rate.

4. DISCUSSION

This study investigated the relationship between arterial stiffness and the imaging markers of CSVD in patients with acute ischemic stroke and found that increased baPWV was associated with the severity of deep lacunar infarction and deep and periventricular WMHs, but not with CMBs. However, in multivariate analysis, only deep lacunar infarction was independently associated with baPWV.

Arterial stiffness increases pulse pressure,
leading to the development and progression of cardiovascular diseases. It is also a known independent predictor of CVD. Age, body weight, body mass index, total cholesterol level, low-density lipoprotein level, triglyceride level, fasting blood sugar level, and intima-media thickness of the internal carotid artery are known to be associated with baPWV. In acute ischemic stroke, the blood pressure rises due to cerebral blood flow autoregulation and stress, which may affect the arterial stiffness during the acute period. Although the biochemical changes affecting arterial stiffness during an acute ischemic stroke are not well known, arterial stiffness is believed to increase due to inflammatory activation, vascular endothelial dysfunction, and other toxic effects. Arterial stiffness may have different measurement values even in the same patient due to daily blood pressure fluctuations. However, arterial stiffness is determined by several factors, including the structural component of the arterial wall, tension in the vascular smooth muscle, and mean blood pressure. In this study, arterial stiffness was measured using baPWV, and in the acute stages of ischemic stroke, this can be affected by increased blood pressure due to stroke, size and location of the stroke, and date of the test. However, arterial stiffness is not solely determined by high blood pressure. Especially in the acute period, where few antihypertensive drugs are used, it may be less responsive to medication. Other previous studies conducted in the acute stage have not analyzed the effects of arterial stiffness on acute blood pressure changes due to the difficulty in correcting for or analyzing blood pressure, which continuously changes. All the previous studies included patients with ischemic stroke. As such, the effect of blood pressure may have been due to some effect in all patients. In this study, baPWV was associated with age, systolic and diastolic blood pressure, pulse pressure, heart rate, glomerular filtration rate, hemoglobin level, and erythrocyte sedimentation rate, but not with serum lipid level.

Arterial stiffness impairs the elasticity of the blood vessel walls, increases pulse pressure, and subsequently causes systemic hypertension. In particular, small arterioles in the brain have low vascular resistance, making them vulnerable to high pulsatile systemic blood pressure, which may be damaging. Increased pulsatile stress disrupts the vascular endothelial cells and smooth muscles, causing blood vessel rupture. Systemic arterial stiffness affects earlier cerebral vessels more than other organs. Because CSVD is difficult to observe directly, brain imaging markers for CSVD are used. Although lacunar infarction, WMH, CMB, dilated PVS, and cerebral atrophy are imaging markers of CSVD, their pathophysiological differences are different. Lacunar infarction and WMH are ischemic lesions caused by decreased perfusion, while CMB is a hemorrhagic lesion. In contrast, dilated PVS is caused by the extravasation of interstitial fluids due to the increased permeability of the blood-brain barrier. As such, the relevant aspects of arterial stiffness and imaging markers may differ.

A recent meta-analysis showed that arterial stiffness was associated with lacunar infarction, WMH, and CMB. Carotid-femoral PWV has been found to be associated with lacunar infarction, PVS of the basal ganglia, a wide range of WMHs, such as Fazekas 3. However, it was not associated with deep CMB, PVS of the central semioval, and deep WMHs. It is notable that not all the imaging markers of CSVD are associated with arterial stiffness. Based on the findings of the few studies on patients with acute ischemic stroke, baPWV is associated with chronic lacunar infarction, WMH, deep CMB, and CSVD. Aortic PWV is associated with WMH volume and chronic lacunar infarction, but not with deep CMB. In one community-based sample, PWV was related to PVS in the white matter, larger WMH volume, and lobar CMB. However, it was not associated with lacunar infarction. Studies have shown inconsistent results regarding the relationship between arterial stiffness and CMB. In one study, PWV was associated with deep CMBs, but not with lobar CMBs. It is believed that the difference between the study group and the research method led to inconsistent results. Unlike other studies, our study was conducted on patients with acute ischemic stroke. However, our results showed that chronic lacunar infarction and WMH were associated with baPWV, consistent with the results of previous studies. In particular, multivariate analysis, which adjusted for variables that may affect baPWV, showed that only deep lacunar infarction was significantly associated. CMB was not associated with baPWV, and no analysis of PVS or acute CSVD
was performed. CMB may be associated with elevated blood pressure, but it can also be caused by brain amyloid angiopathy. Therefore, mechanism other than arterial stiffness may have affected it. This may not have reached statistical significance because of the low frequency of CMBs. CMBs and lacunar infarctions were based on the number of lesions. Furthermore, WMH was analyzed by dividing the grades according to severity, and since the number of study groups was small, it did not seem to be significant.

The limitations of this study are as follows. First, the study group was limited to patients with acute ischemic stroke, and there was no comparison with a normal control group. Second, the number of subjects was small, and other markers of CSVD, such as PVS, were not analyzed. There was also a lack of accurate assessment of peripheral vascular disease, leading to selection bias. Third, there was no analysis according to the mechanism of cerebral infarction, and there is a possibility that the acute cerebral infarction may have affected the variable. Fourth, the imaging evaluation was conducted by one person, and no inter-examiner reliability evaluation was performed. Prospective research on more patients will be needed in the future.

In conclusion, increased baPWV in patients with non-cardioembolic acute ischemic stroke was associated with lacunar infarction and WMH. Among the several markers of CSVD, deep lacunar infarction was a good marker of arterial stiffness. Systemic arterial stiffness in some way affects CSVD, and baPWV may be valuable as a screening tool for CSVD.

REFERENCES

[1] Mancia G, De Backer G, Dominiczak A, Cifkova R, Fagard R, Germano G, et al. 2007 Guidelines for the management of arterial hypertension: the task force for the management of arterial hypertension of the European Society of Hypertension (ESH) and of the European Society of Cardiology (ESC). J Hypertens 2007; 25:1105-1187.

[2] Yamashina A, Tomiyama H, Takeda K, Tsuda H, Arai T, Hirose K, et al. Validity, reproducibility, and clinical significance of noninvasive brachial-ankle pulse wave velocity measurement. Hypertens Res 2002; 25:359-364.

[3] Laurent S, Boutouyrie P, Asmar R, Gautier I, Laloux B, Guize L, et al. Aortic stiffness is an independent predictor of all-cause and cardiovascular mortality in hypertensive patients. Hypertension 2001; 37:1236-1241.

[4] O'Rourke MF, Staessen JA, Vlachopoulos C, Duprez D, Plante GE. Clinical applications of arterial stiffness: definitions and reference values. Am J Hypertens 2002; 15:426-444.

[5] Pantoni L. Cerebral small vessel disease: from pathogenesis and clinical characteristics to therapeutic challenges. Lancet Neurol 2010; 9:689-701.

[6] van Sloten TT, Protogerou AD, Henry RM, Schram MT, Launer LJ, Stelouwer CD. Association between arterial stiffness, cerebral small vessel disease and cognitive impairment: A systematic review and meta-analysis. NeurosciBiobehav Rev 2015; 53:121-130.

[7] Kim YB, Park KY, Chung PW, Kim JM, Moon HS, Youn YC. Brachial-ankle pulse wave velocity is associated with both acute and chronic cerebral small vessel disease. Atherosclerosis 2016; 245:54-59.

[8] Riba-Llena I, Jiménez-Balado J, Castañé X, Girona A, López-Rueda A, Mundet X, et al. Arterial stiffness is associated with basal ganglia enlarged perivascular spaces and cerebral small vessel disease load. Stroke 2018; 49:1279-1281.

[9] Gregoire SM, Chaudhary UJ, Brown MM, Yousry TA, Kallis C, Jäger HR, et al. The Microbleed Anatomical Rating Scale (MARS): reliability of a tool to map brain microbleeds. Neurology 2009; 73:1759-1766.

[10] Moon SY, Na DL, Seo SW, Lee JY, Ku BD, Kim SY, et al. Impact of white matter changes on activities of daily living in mild to moderate dementia. EurNeurol 2011; 65:223-230.

[11] Laurent S, Katsahanis S, Fassot C, Tropeano AI, Gautier I, Laloux B, et al. Aortic stiffness is an independent predictor of fatal stroke in essential hypertension. Stroke 2003; 34:1203-1206.

[12] Mattace-Raso FU, van der Cammen TJ, Hofman A, van Popele NM, Bos ML, Schalekamp MA, et al. Arterial stiffness and risk of coronary heart disease and stroke: the Rotterdam Study. Circulation 2006; 113:657-663.

[13] Munakata M, Ito N, Nunokawa T, Yoshinaga K. Utility of automated brachial ankle pulse wave velocity measurements in hypertensive patients. Am J Hypertens 2003; 16:653-657.

[14] Tomiyama H, Yamashina A, Arai T, Hirose K, Koji Y, Chikamori T, et al. Influences of age and gender on results of noninvasive brachial-ankle pulse wave velocity measurement—a survey of 12517 subjects. Atherosclerosis 2003; 166:303-309.
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[15] Tuttolomondo A, Di Sciacca R, Di Raimondo D, Serio A, D’Aguanno G, Pinto A, et al. Arterial stiffness indexes in acute ischemic stroke: relationship with stroke subtype. *Atherosclerosis* 2010; 211:187-194.

[16] O’Rourke MF, Safar ME. Relationship between aortic stiffening and microvascular disease in brain and kidney: cause and logic of therapy. *Hypertension* 2005; 46:200-204.

[17] Faraci FM, Heistad DD. Regulation of large cerebral arteries and cerebral microvascular pressure. *Circ Res* 1990; 66:8-17.

[18] Zhai FF, Ye YC, Chen SY, Ding FM, Han F, Yang XL, et al. Arterial Stiffness and Cerebral Small Vessel Disease. *Front Neurol* 2018; 9:723.

[19] Song TJ, Kim J, Kim YD, Nam HS, Lee HS, Nam CM, et al. The distribution of cerebral microbleeds determines their association with arterial stiffness in non-cardioembolic acute stroke patients. *Eur J Neurol* 2014; 21:463–469.

[20] Henskens LH, Kroon AA, van Oostenbrugge RJ, Gronenschild EH, Fuss-Lejeune MM, Hofman PA, et al. Increased aortic pulse wave velocity is associated with silent cerebral small-vessel disease in hypertensive patients. *Hypertension* 2008; 52:1120-1126.

[21] Poels MM, Zaccai K, Verwoert GC, Vernooij MW, Hofman A, van der Lugt A, et al. Arterial stiffness and cerebral small vessel disease: the rotterdam scan study. *Stroke* 2012; 43:2637–2642.

[22] Ochi N, Tabara Y, Igase M, Nagai T, Kido T, Miki T, et al. Silent cerebral microbleeds associated with arterial stiffness in an apparently healthy subject. *Hypertens Res* 2009; 32:255-260.