Prevalence of Stroke and Hypoperfusion in Patients With Isolated Vertigo and Vascular Risk Factors

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Background and Purpose: The aim of this study was to determine the prevalence and associated factors of stroke and hypoperfusion among patients with isolated vertigo and vascular risk factors.

Methods: We studied 157 patients with isolated vertigo who had undergone multimodal magnetic resonance imaging. Magnetic resonance angiography (MRA) was used to measure the diameters of vertebrobasilar arteries and to evaluate morphologic changes to vessels. Measurements obtained included length of the basilar artery and curvature index for the vertebral artery (VA). Perfusion-weighted imaging (PWI) was performed to determine relative cerebral blood flow, relative cerebral blood volume (rCBV), time to peak (TTP), and mean transit time for two mirror regions of interest (ROIs) in each map. Regional hypoperfusion of the cerebellum was considered significant when TTP and mean transit time (MTT) were present in ≥2 adjacent slices.

Results: The prevalence of stroke in patients with isolated vertigo and vascular risk factors was 24.8% (n = 39). Visual assessment revealed cerebellar hypoperfusion in 57.6% (68/118) of non-stroke patients. Multivariate logistic regression indicated that diabetes mellitus (P = 0.049, OR = 2.758), VA stenosis or hypoplasia (P = 0.023, OR = 3.486), and relative TTP of cerebellum (P = 0.002, OR = 3.197) were independent risk factors for stroke and LVA curvature index (P = 0.026, OR = 2.049), VA stenosis and hypoplasia (P = 0.009, OR = 2.977) were independent risk factors for hypoperfusion.

Conclusions: The prevalence of stroke and hypoperfusion is higher in patients with isolated vertigo and vascular risk factors, compared with matched controls. Potential risk factors include diabetes mellitus, VA stenosis or hypoplasia, and enlarged VA curvature index.

Keywords: vertigo, stroke, hypoperfusion, vertebral artery, basilar artery, vessel curvature
INTRODUCTION

Patients with acute isolated vertigo have vertigo without focal neurological symptoms (unsteady gait and nystagmus are allowed) (1, 2) involving or preceded by stroke/transient ischemic attack (TIA) (3). Diagnosis of vertigo in patients presenting with posterior circulation stroke/TIA has recently increased markedly (4, 5). A population-based registry showed that 90% of isolated posterior circulation TIAs (half of which were associated with isolated vertigo symptoms), were not recognized when the patient first presented to a health-care practitioner (6). One retrospective recently reported that ∼20% of patients who have had a brainstem stroke report having also had isolated attacks of vertigo (7). Regrettfully, the diagnosis of stroke in patients with acute isolated vertigo remains a challenge when the diagnosis is based on bedside examination findings and routine MRIs (3). Application of HINTS (head impulse, nystagmus patterns, test of skew) has greatly enhanced the diagnosis of stroke in acute isolated vertigo. However, HINTS could not be applied to the majority of patients with isolated vertigo because the vestibular symptoms or signs had already resolved in about 70% of patients (1). Although diffusion-weighted imaging magnetic resonance (DWI-MR) remains the standard modality with which to confirm acute infarction. However, when performed within 24–48 h of vertigo onset, images obtained with DWI-MR are falsely negative for ∼20% of patients with clinically verified vertebrobasilar stroke (1, 8).

An important recent study that included initial and follow-up images indicated that 12% of patients showed unilateral cerebellar hypoperfusion on perfusion-weighted imaging (PWI) without an infarction on DWI (1). The results obtained with CT perfusion showed that unilateral hypoperfusion of the cerebellum and brainstem heralds the occurrence of isolated vertigo (3). Another study found that disequilibrium in the relative mean transit time (rMTT) of the bilateral cerebellum (as demonstrated with PWI) that hypoperfusion could trigger recurrent isolated vertigo (9). The combination of perfusion imaging analysis and clinical assessment may help to distinguish an attack of vertigo from a subtle stroke/TIA, it remains difficult to identify useful predictors of stroke in patients with isolated vertigo (10, 11).

The present study aims to characterize the prevalence of stroke and hypoperfusion and to analyze clinical findings and radiographic images to facilitate early detection of infarction or hypoperfusion in patients with isolated vertigo and vascular risk factors. To achieve this, we sought to identify traditional cerebrovascular risk factors based on clinical presentation, results of bedside examinations, and morphological changes in vertebrobasilar arteries as determined by magnetic resonance angiography (MRA). Using PWI, we characterized perfusion in the internal inferior cerebellum, dorsolateral medulla, occipital and temporal lobes, pons at the level of the vestibular nucleus, and thalamus in the plane of the internal capsule.

METHODS

Ethical Considerations

This prospective, single-center, observational study was approved by the Institutional Ethics Committee of People’s Hospital of Zhengzhou, China. Each participant in the study provided informed consent.

Experimental Design and Patient Population

This was a prospective study that included four groups of patients with suspected vascular isolated vertigo. The stroke group included patients suffering from isolated vertigo caused by vascular disease as demonstrated by neuroimaging (typically DWI-MR). The non-stroke group comprised participants whose isolated vertigo was clinically suspected to have a vascular etiology, despite negative findings on DWI-MR. Patients in the non-stroke group were further classified, based on PWI findings, as hypoperfusion or non-hypoperfusion.

A total of 279 patients seen at the Department of Neurology at Zhengzhou People’s Hospital were enrolled during the period from November 2014 to February 2017. Inclusion criteria were as follows: (1) acute onset of vertigo suspected to result from vascular ischemia; (2) no focal neurological signs or symptoms except for unsteady gait, nystagmus, intolerance of head motion, autonomic nervous symptoms or tinnitus; (3) at least one cerebrovascular risk factor such as hypertension, diabetes or elevated blood glucose, dyslipidemia disorder, coronary atherosclerotic heart disease, history of stroke, hyperhomocysteinemia, smoking, and alcoholism; (4) age > 18 years; (5) positional vertigo considered to be independent of benign paroxysmal positional vertigo, Meniere disease, and vestibular neuritis (based on clinical symptoms and signs, if the results of vestibular function tests were non-negative). Patients were excluded if they had migraine (n = 12), sever emotional disease (n = 1), cognitive disorder (n = 1), compression of the eighth cranial nerve (n = 3), trauma(n = 1), hemorrhage (n = 5), taken drugs that might induce vertigo within the past 2 weeks (n = 4), subclavian steal syndrome (n = 2), not undergo DWI-MR (n = 4), contrast-enhanced MRA (n = 3), Meniere’s disease (n = 8), vestibular neuritis (n = 19), benign paroxysmal positional vertigo (n = 42), and other ototrinolaryngology disease (n = 1). PWI was performed to evaluate the remaining 162 patients. However, for 5 patients, the quality of PWI images was poor; these patients were excluded. In the last, there remained 157 patients with isolated vertigo with at least one vascular risk factor and with intact data.

For the purpose of this study, the following factors were evaluated: history of hypertension [previous diagnosis of arterial hypertension: systolic blood pressure > 140 mmHg and/or diastolic > 90 mmHg, or use of antihypertensive agents (past or present)], diabetes mellitus (previous diagnosis of diabetes or past or present use of antidiabetic agents), hyperlipidemia (cholesterol > 5.17 mmol/L and/or triglycerides > 1.71 mmol/L), hyperhomocysteinemia (> 15.0 μmol/L), consumption of alcohol at least once a week (1 standard alcohol consumption is equivalent to 120 mL of wine, 360 mL of beer, or 45 mL of distilled spirits), smoking (continuous or cumulative history of smoking > 6 mo and ≥1 cigarette per day), history of coronary artery disease (CAD), and presence of cerebrovascular stenosis (> 50% stenosis of ≥1 intracranial artery, according to MRA). All patients with
cerebrovascular risk factors had been previously diagnosed as such and/or were already taking medications for these conditions.

**MR EXAMINATION PROTOCOL AND IMAGE PROCESSING**

MRI was performed in a 3.0 T gradient-echo (GE) Signa HDX (Fairfield, USA). The acquisition sequence included T1- and T2-weighted imaging, fluid attenuated inversion recovery (FLAIR) imaging and DWI and apparent on diffusion coefficient (ADC). DWI was acquired using spin-echo planar imaging sequence with TR 50000, TE 77.6, field of view 220 × 220 mm and slice thickness of 5.0 mm. At the same time, cervical CE-MRA was performed with injection of gadopentetate dimeglumine and three-dimensional time-of-flight (3D-TOF) MRA was also performed in the HDX with a repetition time of 24 ms, TR of 4.8, echo time of 6 ms, TE of 1.6/Fr, field of view of 320 × 320 mm and section thickness of 0.8–1.6 mm. A definite infarction was made when the lesion with high signal on DWI and low signal on ADC.

The means of three dimensional maximum intensity projection (3D-MIP) was used in the imaging reconstruction and data acquisition and we done the work on a post-processing GE machine (Siemens, Inc., Munich, Germany) with the AW Volume share 5 software (version 9.4.05). Measurement information was collected by two experienced observers in order to reduce bias. The diameter of all four segments of both sides of vertebral artery (V1–V4) and the diameter of basilar artery were measured in the reformed window with the same setting. Vertebral artery segments were classified as follows: V1, from origin to the transverse foramina of C5 or C6 vertebrae, V2, from the transverse foramina of C5 or C6 to the transverse foramina to C2, V3, from the C2 transverse foramina to dura, V4, from dura to the confluence of 2 vertebral arteries to form the basilar artery (12, 13).

Diameter of the basilar artery was measured at 3 locations: the point of junction with the vertebral artery, the top point, and the middle point. The average of these 3 values was taken as mean diameter of the basilar artery. In addition, the morphological changes of vertebrobasilar artery system measured also included actual and straight vertebral artery length (V1–V2; Figure 1A), basilar artery actual and straight length, basilar artery bending length (Figure 1B), and angle degree of vertebral artery (14). Basilar artery bending length refers to the distance from the basilar artery bending point to the basilar standard line. Vertebral artery hypoplasia (VAH) was defined as a V4 diameter ≤2 mm, slim or absent, or an asymmetry ration for the two sides of vertebral arteries >1:1.7 (15). Ratio of BA length = Actual length of BA/straight length of BA. Curvature index of VA = (actual length—straight length)/(straight length) × 100% (16). Patients were also classified according to the existence vs. lack of a fetal posterior cerebral artery. In accordance with North American Symptomatic Carotid Endarterectomy Trial method, vertebral artery stenosis was calculated as: (normal diameter at the distal end—narrowest diameter)/normal diameter at the distal end × 100%.

**PWI Examination and Obtaining Parameter Values**

PWI were obtained from dynamic susceptibility contrast-enhanced perfusion MR images using a gradient-echo echo planar image technique measuring the variation of signal intensity. The contrast agent used in our study was gadopentetate dimeglumine (Gd-DTPA), which can alter the local magnetic filed and therefore cause the decrease of signal intensity in the surrounding brain tissue. Because scan protocols and post-processing methods vary widely among manufacturers, the following scanning parameters were used: TR 1,500 ms, TE 15.2 ms, field of view 24 × 24 mm, section thickness of 5 mm with a 1.5 mm gap. During scanning, 0.2 ml/kg Gd-DTPA was injected at a rate of 4 mL/s; this was followed with an equivalent dose of saline. In the last, a total of 1,200 images were obtained from 24 slices and 50 repeated consecutive times and the original images were processed by the GE Brainstat software application (Functool 9.4.05). The AIF protocol was used to generate processed maps of relative cerebral blood flow (rCBF), relative cerebral blood volume (rCBV), mean transit time (MTT), and time to peak (TTP) (Figure 2). The AIF protocol is based on artery input functions and vein output functions; all measurements were performed by an experienced neuroradiologist. PWI was obtained within 24 h from the symptom onset in 82%, and within 48 h in 94%. The remaining 3 patients received PWI between 3 and 4 days from the symptom onset. The results of PWI were independently assessed by a neurologist (Y. S.) and neuroradiologist (M.Q.K.) who were blinded to all clinical information. They visually assessed the presence of asymmetrical hypoperfusion in the cerebellum using TTP and MTT maps (Figure 2). A regional hypoperfusion was considered significant when it was present in ≥2 adjacent slices. If there were any disagreements between the 2, we made a final decision based on consensus. They did not visually assess the medulla oblongata, pons and thalamus because of not easy to visually assess by PWI. All patients with cerebellar hypoperfusion on PWI had received follow-up PWI within 3 month after resolution of the symptoms. We considered cerebellar hypoperfusion as a cause of isolated vertigo in patients with vascular risk factors only when the symptoms of vertigo were not recurrent and the initial hypoperfusion was normalized on the follow-up images.

The regions of interest (ROIs) were located in the internal inferior cerebellum, dorsolateral medulla, occipital and temporal lobes, corona radiata, pons at the level of the vestibular nucleus, and thalamus in the plane of the internal capsule. Mirrored ROIs were automatically created on the same plane in the position of symmetry, after the axis of symmetry was created. Each ROI was outlined in the map of CBF firstly in reference to MTT to avoid the large vessels and then automatically copied onto the CBV, MTT, and CBV panels. Considering the physiological differences, the impact of hemodynamic constants and allowing convenient comparisons between individuals, the relative values of the two mirror ROI of each map were calculated including...
rrCBF, rrCBV, rMTT, and rTTP (= the absolute difference of parameter value between left to right ROI), which represent the absolute difference between the two symmetry ROI.

To assess the reliability of PWI, we also performed PWI in 32 patients with peripheral vestibulopathy. Patients were considered to have peripheral vestibulopathy if MRI and DWI scanning images were negative but tests of vestibular function were positive.

**Statistical Analysis**

Statistical analyses were performed using SPSS 19.0 software (IBM, Chicago, IL, USA). Normally distributed continuous variables were expressed as mean ± standard deviation and independent-samples t-test was used for two groups comparison, one-way analysis of variance (ANOVA) for three or more groups comparison. Whereas, non-normal distribution parametric were presented as median (range) if not passed the normal distribution test, and Mann-Whitney U-test were used for the comparison between the two groups, while to compare the differences among three groups or more, Kruskal-Wallis H test were used. Categorical variables were expressed as percentage and the chi square test was used in all the test of rate. According to the sample size and expected value, special chi square formula, the correction formula or the exact probability formula were used, respectively. In all of the tests, P < 0.05 were considered statistically significant. Variables with a P < 0.05 in the univariate analysis were selected and evaluated by multivariate logistic regression models with the forward stepwise selection method.

**RESULTS**

**General Demographic Findings**

In total, 157 eligible patients with isolated vertigo, aged 28–90 years (median, 59.9 ± 12.6 years) were included in the study. Male patients accounted for 49.7% of the total. Among 157 patients, brain DWI-MR revealed recent ischemic stroke in 39 (24.8%) patients. In the remaining 118 patients without infarction lesions, PWI revealed cerebellar hypoperfusion by visually assessed in 68 patients (57.6%, 68/118). Ten of the patients also had a reduced CBF with normal CBV in the corresponding areas. Forty patients had a perfusion defect in the area of the medial posterior inferior cerebellar artery; the remaining 28 perfusion defects were found in the whole posterior inferior cerebellar artery. Among 68 patients, 28 (41.2%) had not recurrent vertigo symptoms and showed a perfusion defect in unilateral or bilateral cerebellum on TTP and MTT maps and were shown to normalize the initial cerebellar hypoperfusion without a complete infarction on the follow-up images. PWI revealed a perfusion defect in 2 of the 32 patients with peripheral vestibulopathy.

**Analysis of Infarcts**

Of the 39 infarcted patients, infratentorial lesions, and supratentorial lesions were both found. The stroke lesions in our study involved were cerebellum (n = 25). Eighteen patients had infarctions restricted to the unilateral (n = 13) or bilateral (n = 5) cerebellum, whereas 7 had additional infarctions in the frontal (n = 1), thalamus (n = 3) or occipital lobe (n = 3),
FIGURE 2 | Images of a case of isolated vertigo suspected to be caused by hypodynamics of brain. (A) Sixty-one-year-old man had isolated vertigo on 4 April 2016 and performed with MRI, MRA, CEMRA, and PWI. (A) DWI-MRI was normal and showed no acute infarctions; (B) outlines of ROIs of bilateral cerebellum and medulla; (C) time-signal intensity curve; (D,E) cerebral blood flow (CBF) and cerebral blood volume: a similar color differences were not observed in the rCBF and rCBV panels. (F) Mean transit time (MTT): values of the right cerebellum varied from the left ones and are not obvious; (G) time to peak (TTP): TTP values obtained for the right cerebellum are longer than those obtained for the left cerebellum; (H,I) CEMRA and MRA showed the tortuous, dominant left vertebral artery, and severe stenosis at the beginning point.

brainstem (3 pons, 2 medulla oblongata, 1 mesencephalon, 1 mesencephalon association with pons), pedunculus cerebellaris medius ($n = 2$), splenium of corpus callosum ($n = 2$), and temporoparietal ($n = 3$) area. Among 39 patients, 16 had acute transient vertigo, and 23 had continuous vertigo (>24 h). In patients with transient vertigo, infarcts usually occurred in the area of the posterior inferior cerebellar artery (cerebellum; $P = 0.04$); in patients with persistent vertigo, infarcts were typically found in the area of the basilar artery (brainstem; $P = 0.01$; Table 1).

Vascular Risk Factors and Clinical Characteristics for Stroke and Hypoperfusion

Stroke was more frequent among patients with history of stroke, hypertension, hyperhomocysteinemia, diabetes mellitus, or cerebrovascular stenosis, compared to patients without
cerebrovascular risk factors. There were significant differences between the stroke group and non-stroke group in the incidence of continuous vertigo and rotary nystagmus; however, the overall incidences of nystagmus and sympathetic symptoms were higher in the non-stroke cohort than in the stroke cohort (Table 2).

There were significant differences between patients with hypoperfusion and patients without non-hypoperfusion who had a history of smoking, nystagmus, and gait instability (Table 2).

**Radiological Imaging Risk Factors for Stroke and Hypoperfusion**

Stroke and non-stroke groups differed significantly in terms of mean diameter of the basilar artery (P = 0.02) and VA stenosis or hypoplasia (P = 0.00). PWI maps revealed significant differences between stroke and non-stroke groups in terms of rrCBF of pons, rTTP of centrum semiovale, rMTT, and rTTP of cerebellum ROIs (P = 0.01, 0.01, 0.01, 0.00, respectively; Table 3). However, the curvature index of left vertebral artery (P = 0.01) and levels of VA stenosis or hypoplasia (P = 0.00) were similar in the hypoperfusion and non-hypoperfusion groups. As for the PWI parameters, only rTTP of cerebellum ROIs was found to differ significantly between hypoperfusion and non-hypoperfusion groups (P = 0.00; Table 3).

In the current study, 3.2% (5/157) of patients with isolated vertigo had a fetal posterior cerebral artery (2 bilateral, 2 right, 1 left). The prevalence of fetal posterior cerebral artery was similar in the stroke and non-stroke groups. rrCBF, rrCBV, rMTT, and rTTP values for occipital and temporal lobe ROIs were similar in patients with vs. without fetal posterior cerebral arteries.

The PWI maps revealed that rrCBF, rrCBV, rMTT, and rTTP values for the occipital and temporal lobe ROIs were similar in the stroke and non-stroke groups (Table 3). Similarly, the prevalence of fetal posterior cerebral artery was similar in the hypoperfusion and non-hypoperfusion groups. Relative rCBF, rrCBF, rMTT, and rTTP values for occipital lobe ROIs were similar in the hypoperfusion and non-hypoperfusion groups. Though rrCBF and rrCBV of temporal lobe ROIs differed significantly between the hypoperfusion and non-hypoperfusion groups, neither rMTT nor rTTP of temporal lobes ROIs was significantly delayed in the hypoperfusion group (Table 3).

**Multivariate Logistic Regression Analysis for Stroke and Hypoperfusion**

Multivariate logistic regression indicated that diabetes mellitus (P = 0.049, 95%CI: 1.85–10.56, OR = 2.758), VA stenosis or hypoplasia (P = 0.00), and history of vertigo (P = 0.00) were significant risk factors for Stroke and Hypoperfusion.

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### Table 1 | Analysis of infarct lesions.

| Temporary vertigo | Continuous vertigo | *P* |
|-------------------|--------------------|-----|
| Stroke (n = 16)   | Non-stroke (n = 23) | 0.35 |

| ACI, anterior circulation infarction; PCI, posterior circulation infarction; PICA, posterior inferior cerebellar artery; BA, basilar artery; PCA, posterior cerebral artery. |

### Table 2 | Cerebrovascular risk factors and clinical characteristics stroke and hypoperfusion.

|                  | Stroke (n = 39) | Non-stroke (n = 118) | *P*  |
|------------------|----------------|----------------------|------|
| Age (years)      | 66 ± 11.1      | 59.0 ± 12.9          | 0.12 |
| Male gender (%)  | 23 (59.0)      | 55 (46.6)            | 0.18 |
| Hypertension (%) | 32 (82.1)      | 71 (60.1)            | 0.01 |
| Diabetes mellitus (%) | 23 (59.0) | 36 (30.5)           | 0.00 |
| Dyslipidemia (%) | 21 (53.8)      | 82 (69.5)            | 0.08 |
| Coronary heart disease (%) | 6 (15.4) | 28 (23.7) | 0.27 |
| Smoking (%)      | 14 (35.9)      | 27 (22.9)            | 0.11 |
| Alcoholism (%)   | 10 (25.6)      | 18 (15.3)            | 0.14 |
| Stroke history (%) | 13 (33.3) | 13 (11.0)           | 0.00 |
| Hyperhomocysteinemia (%) | 13 (33.3) | 21 (17.8) | 0.04 |
| Hyperuricemia (%) | 0 (0)         | 3 (2.5)              | 0.32 |
| History of vertigo (%) | 0 (0) | 7 (5.9)             | 0.12 |
| Gait instability (%) | 23 (61.5) | 64 (54.2)          | 0.43 |
| Hearing impairment (%) | 1 (2.6) | 6 (5.1)            | 0.51 |
| Trigger factors (%) | 51 (1.5) | 11 (2.0)            | 1.00 |
| Nystagmus (%)    | 23 (59.0)      | 89 (75.4)            | 0.04 |
| Continuous vertigo (%) | 23 (59.0) | 41 (34.7) | 0.01 |
| Sympathetic symptoms (%) | 29 (74.4) | 106 (89.8) | 0.02 |

### Table 3 | Cerebrovascular risk factors and clinical characteristics stroke and hypoperfusion.

|                  | Stroke (n = 68) | Non-stroke (n = 50) | *P*  |
|------------------|----------------|---------------------|------|
| Age (years)      | 60.7 ± 13.7    | 56.6 ± 11.6          | 0.09 |
| Male gender (%)  | 28 (41.2)      | 27 (54.0)            | 0.18 |
| Hypertension (%) | 42 (61.8)      | 29 (58.0)            | 0.68 |
| Diabetes mellitus (%) | 24 (35.3) | 12 (24.0)           | 0.19 |
| Dyslipidemia (%) | 45 (66.2)      | 37 (74.0)            | 0.36 |
| Coronary heart disease (%) | 17 (25.0) | 11 (22.0) | 0.71 |
| Smoking (%)      | 11 (16.2)      | 16 (32.0)            | 0.04 |
| Alcoholism (%)   | 10 (14.7)      | 8 (16.0)             | 0.85 |
| Stroke history (%) | 9 (13.2)     | 4 (8.0)              | 0.37 |
| Hyperhomocysteinemia (%) | 12 (17.6) | 11 (22.0) | 0.31 |
| Hyperuricemia (%) | 0 (0)         | 1 (2.0)              | 0.81 |
| History of vertigo (%) | 3 (4.4) | 2 (4.0)             | 0.71 |
| Gait instability (%) | 24 (45.0) | 24 (48.0)          | 0.00 |
| Hearing impairment (%) | 1 (1.5) | 3 (6.0)            | 0.70 |
| Trigger factors (%) | 55 (80.9) | 3 (6.0)            | 0.41 |
| Nystagmus (%)    | 16 (23.5)      | 34 (68.0)            | 0.00 |
| Continuous vertigo (%) | 21 (30.9) | 20 (40.0) | 0.30 |
| Sympathetic symptoms (%) | 63 (92.6) | 43 (86.0) | 0.24 |

DWI, diffusion-weighted imaging; PWI, perfusion-weighted imaging.
## TABLE 3 | Risk factors for stroke and hypoperfusion on radiological imaging.

|                      | DWI                      | p     | PWI (non-stroke) | P     |
|----------------------|--------------------------|-------|------------------|-------|
|                      | Stroke (n = 39)          |       | Non-stroke (n = 118) |       |
|                      |                         |       |                  |       |
| Diameters of RVA     | 2.76 ± 0.68              | 0.77  | 2.76 ± 0.68      | 0.77  |
| Diameters of LVA     | 3.19 ± 0.67              | 0.83  | 3.19 ± 0.67      | 0.83  |
| Actual lengths of RVA| 2.80 ± 0.66              | 0.83  | 2.80 ± 0.66      | 0.83  |
| Actual lengths of LVA| 114.23 ± 12.45           | 0.85  | 114.23 ± 12.45   | 0.85  |
| Straight lengths of RVA| 118.18 ± 13.66          | 0.84  | 118.18 ± 13.66   | 0.84  |
| Straight lengths of LVA| 105.44 ± 9.52          | 0.86  | 105.44 ± 9.52    | 0.86  |
| Straight lengths of BA| 107.08 ± 15.13          | 0.70  | 107.08 ± 15.13   | 0.70  |
| Curvature index of RVA| 8.36 ± 6.98             | 0.69  | 8.36 ± 6.98      | 0.69  |
| Curvature index of LVA| 10.90 ± 7.44            | 0.72  | 10.90 ± 7.44     | 0.72  |
| Diameters of BA      | 3.12 ± 1.54              | 0.02  | 3.12 ± 1.54      | 0.02  |
| Actual length of BA  | 26.07 ± 5.83             | 0.09  | 26.07 ± 5.83     | 0.09  |
| Straight length of BA| 24.45 ± 5.54             | 0.75  | 24.45 ± 5.54     | 0.75  |
| Ratio of BA length   | 1.07 ± 0.50              | 0.45  | 1.07 ± 0.50      | 0.45  |
| BA bending length    | 3.12 ± 3.00              | 0.18  | 3.12 ± 3.00      | 0.18  |
| Angle degree of VAs  | 72.11 ± 38.97            | 0.40  | 72.11 ± 38.97    | 0.40  |
| VA stenosis or hypoplasia | 29 (74.4)              | 0.00  | 29 (74.4)        | 0.00  |
| Fetal PCA            | 3(0.08)                  | 0.10  | 3(0.08)          | 0.10  |
| rCBF of medulla      | 6.79 ± 6.99              | 0.87  | 6.79 ± 6.99      | 0.87  |
| rCBV of medulla      | 0.61 ± 0.52              | 0.99  | 0.61 ± 0.52      | 0.99  |
| rMTT of medulla      | 0.97 ± 1.05              | 0.94  | 0.97 ± 1.05      | 0.94  |
| rTTP of medulla      | 0.77 ± 0.92              | 0.99  | 0.77 ± 0.92      | 0.99  |
| rCBF of cerebellum   | 6.90 ± 7.64              | 0.33  | 6.90 ± 7.64      | 0.33  |
| rCBF of cerebellum   | 0.79 ± 0.97              | 0.11  | 0.79 ± 0.97      | 0.11  |
| rMTT of cerebellum   | 1.47 ± 2.07              | 0.01  | 1.47 ± 2.07      | 0.01  |
| rTTP of cerebellum   | 1.45 ± 1.69              | 0.00  | 1.45 ± 1.69      | 0.00  |
| rCBF of pons         | 4.89 ± 3.75              | 0.89  | 4.89 ± 3.75      | 0.89  |
| rCBV of pons         | 0.58 ± 0.40              | 0.01  | 0.58 ± 0.40      | 0.01  |
| rMTT of pons         | 0.84 ± 1.08              | 0.18  | 0.84 ± 1.08      | 0.18  |
| rTTP of pons         | 1.05 ± 1.29              | 0.14  | 1.05 ± 1.29      | 0.14  |
| rCBF of thalamus     | 3.24 ± 2.49              | 0.91  | 3.24 ± 2.49      | 0.91  |
| rCBF of thalamus     | 0.33 ± 0.27              | 0.71  | 0.33 ± 0.27      | 0.71  |
| rMTT of thalamus     | 1.24 ± 4.00              | 0.19  | 1.24 ± 4.00      | 0.19  |
| rTTP of thalamus     | 0.80 ± 1.15              | 0.02  | 0.80 ± 1.15      | 0.02  |
| rCBF of centrum semiovale | 3.09 ± 3.13           | 0.76  | 3.09 ± 3.13      | 0.76  |
| rCBV of centrum semiovale | 0.26 ± 0.25            | 0.57  | 0.26 ± 0.25      | 0.57  |
| rMTT of centrum semiovale | 0.91 ± 0.88           | 0.12  | 0.91 ± 0.88      | 0.12  |
| rTTP of centrum semiovale | 1.13 ± 1.23           | 0.01  | 1.13 ± 1.23      | 0.01  |
| rCBF of occipital lobe| 2.44 (4.13)             | 0.71  | 2.44 (4.13)      | 0.71  |
| rCBV of occipital lobe| 0.23 (0.25)             | 0.71  | 0.23 (0.25)      | 0.71  |
| rMTT of occipital lobe| 0.57 (0.98)             | 0.42  | 0.57 (0.98)      | 0.42  |
| rTTP of occipital lobe| 0.59 (1.23)             | 0.01  | 0.59 (1.23)      | 0.01  |
| rCBF of temporal lobe| 3.49 (3.76)             | 0.26  | 3.49 (3.76)      | 0.26  |
| rCBV of temporal lobe| 0.32 (0.37)             | 0.80  | 0.32 (0.37)      | 0.80  |
| rMTT of temporal lobe| 0.76 (1.26)             | 0.40  | 0.76 (1.26)      | 0.40  |
| rTTP of temporal lobe| 0.76 (1.39)             | 0.95  | 0.76 (1.39)      | 0.95  |

DWI, diffusion-weighted imaging; PWI, perfusion-weighted imaging; RVA, right vertebral artery; LVA, left vertebral artery; BA, basilar artery; PCA, posterior cerebral artery; rCBF, relative cerebral blood flow; rCBV, relative cerebral blood volume; rMTT, relative mean transit time; rTTP, relative time to peak.

hypoperfusion (P = 0.023, 95%CI: 1.55–6.62, OR = 3.486), and rTTP of cerebellum (P = 0.002, 95%CI: 1.67–4.05, OR = 3.197) were independent risk factors for stroke (Table 4) and curvature index of LVA (P = 0.026, 95%CI: 1.00–1.10, OR = 2.049) and VA stenosis or hypoplasia (P = 0.009, 95%CI: 1.31–6.74, OR = 2.977) were independent risk factor for hypoperfusion (Table 5).
were independent risk factors for hypoperfusion. Accurate stroke and curvature index of LV A, V A stenosis or hypoplasia and rTTP of cerebellum were independent risk factors for interestingly, diabetes mellitus, V A stenosis or hypoplasia and recent infarctions, hypoperfusion was found in 57.6%. More was 41.2%. Among all patients with isolated vertigo without initial hypoperfusion was normalized on follow-up images 24.8%. The prevalence of cerebellar hypoperfusion for which n

TABLE 4 | Multivariate logistic regression analysis for stroke in patients with isolated vertigo (n = 157).

| Variants                | B      | S.E.  | Wald  | Sig. (p)     | OR (95% C.I.) |
|-------------------------|--------|-------|-------|--------------|---------------|
| Hypertension            | 0.770  | 0.581 | 1.760 | 0.185        | 2.161 (1.01–7.57) |
| Diabetes mellitus       | 1.015  | 0.515 | 3.884 | 0.049        | 2.758 (1.85–10.58) |
| Stroke history          | 1.096  | 0.643 | 2.904 | 0.088        | 2.993 (0.57–4.78) |
| Hyperhomocysteinemia    | 0.499  | 0.544 | 0.843 | 0.358        | 1.647 (0.15–13.4) |
| Nystagmus               | -0.804 | 0.561 | 2.054 | 0.152        | 0.447 (0.28–2.28) |
| Continuous vertigo      | -0.219 | 0.533 | 0.169 | 0.681        | 0.803 (0.47–1.33) |
| Sympathetic symptoms    | 0.539  | 0.666 | 0.656 | 0.418        | 1.715 (0.76–1.86) |
| Mean BA diameter        | 0.170  | 0.230 | 0.547 | 0.460        | 1.186 (1.19–10.20) |
| VA stenosis or hypoplasia| 1.249  | 0.548 | 5.195 | 0.023        | 3.486 (1.55–6.62) |
| rTTP of cerebellum      | 1.162  | 0.371 | 9.820 | 0.002        | 3.197 (1.67–4.05) |
| rCBV of pons            | 0.495  | 0.461 | 1.156 | 0.282        | 1.641 (1.19–10.20) |
| rTTP of centrum semiovale| 0.325  | 0.257 | 1.603 | 0.205        | 1.384 (0.84–2.29) |
| Constant                | -4.682 | 1.275 | 13.494| 0.000        | 0.000 |

VA, vertebral artery; rCBV, relative cerebral blood volume; rTTP, relative time to peak.

TABLE 5 | Multivariate logistic regression analysis for hypoperfusion in patients with isolated vertigo (n = 118).

| Variants                | B      | S.E.  | Wald  | Sig. (p)     | Exp (B)          |
|-------------------------|--------|-------|-------|--------------|------------------|
| Smoking                 | -0.595 | 0.485 | 1.508 | 0.219        | 0.551 (0.21–1.43) |
| Curvature index of LVA  | 0.048  | 0.026 | 3.379 | 0.026        | 2.049 (1.00–1.10) |
| Line length of BA       | -0.005 | 0.060 | 3.625 | 0.057        | 0.909 (0.83–1.00) |
| VA stenosis or hypoplasia| 1.091  | 0.417 | 6.841 | 0.009        | 2.977 (1.31–6.74) |
| Constant                | 1.775  | 1.240 | 2.049 | 0.152        | 1.000 |

LVA, left vertebral artery; BA, basilar artery.

DISCUSSION

In this study of 157 patients with isolated vertigo accompanied by cerebrovascular risk factors, the prevalence of stroke was 24.8%. The prevalence of cerebellar hypoperfusion for which initial hypoperfusion was normalized on follow-up images was 41.2%. Among all patients with isolated vertigo without recent infarctions, hypoperfusion was found in 57.6%. More interestingly, diabetes mellitus, VA stenosis or hypoplasia and rTTP of cerebellum were independent risk factors for stroke and curvature index of LVA, VA stenosis or hypoplasia were independent risk factor for hypoperfusion. Accurate identification of vertigo/dizziness secondary to ischemia and risk factors may lead to appropriate timely intervention, minimizing stroke-related damage, and help in tailoring the most appropriate therapy for patients.

A previous study showed that the prevalence of stroke among patients with vertigo ranges from 3.2–25% (17–19). One recent study discovered that stroke was diagnosed in 27% of patients with acute transient vestibular syndrome (1). The present study used 3.0T MRI and recruited patients with isolated vertigo with cerebrovascular risk factor load may contribute to the higher detection rate of small infarct lesions and hypoperfusion. Also, considering the high incidence of stroke/hypoperfusion in patients with isolated vertigo and it is of great importance to find predictors for stroke/hypoperfusion for these vertiginous patients in order to perform timely appropriate treatment (6).

Previous studies have shown that central isolated vertigo is caused by lesions restricted to vestibular nuclei, cerebellar flocculus, tonsil, nodulus, inferior cerebellar peduncle, or nucleus prepositus hypoglossi (3, 17). The present study interestingly found that the infarction patients with transient vertigo usually occurred in the cerebellum, while the infarction patients with persistent vertigo usually were in the brain-stem. A disequilibrium of perfusion in bilateral cerebellum implies that hypoperfusion in the posterior circulation may lead to recurrent transient isolated vertigo (9). Using a rat model of global hypoperfusion, the researchers found that the vestibular structures were vulnerable to ischemia more than any other structures in the brainstem and cerebellum, and the medial vestibular nucleus was the most vulnerable (20). These two findings confirmed our hypothesis, but future studies to confirm these findings should include a larger sample size than that achieved for this study.

Previous studies have found VAH may cause hypoperfusion in the area of the posterior inferior cerebellar artery (13, 15). Ahn et al. (21) describes two patients with recurrent isolated vertigo and subsequent cerebellar infarction, probably because of unilateral hypoplastic vertebral artery. In a double-blind retrospective cohort study, results show 85.7% of the patients complaining of isolated positional vertigo with at least three stroke risk factors, had a vertebral artery abnormality as stenotic or hypoplastic (22). A study revealed that the risk of vascular events or long-time risk of stroke was higher in patients with vestibular symptom associated with more vascular risk factors and silent infarctions in CT imaging (8). A 4-year follow-up study showed that patients with vertigo are at higher risk (3.01-fold) for stroke than the general population. Furthermore, vertigo patients with 3 risk factors had a 5.51-fold increase in risk for stroke, compared to patients with vertigo but without risk factors (23). Furthermore, diabetes mellitus can accelerate intracranial atherosclerosis, especially in the posterior cerebral circulation regions, with a tendency to invade the deep perforating branches of the basilar artery (24), which constitutes the main blood supply of the brainstem. In addition, the autonomic nervous enuresis of the posterior circulation is less extensive than for the anterior circulation, which also contributes to the increased risk of PCI in patients with diabetic mellitus (25).

Interestingly, LVA curvature index was associated with lower perfusion, and cerebellar rTTP was an independent risk factor for stroke. It has been widely accepted that TTP and MTT are sensitive indices for hypoperfusion of the brainstem unrelated to CBF or CBV (9). To our knowledge, this is the first study to investigate the effects of VA curvature on hemodynamics captured by PWI. Additional studies will be necessary to confirm the findings presented above. Previous research from our laboratory showed that the combination of cerebrovascular risk
factors and tortuosity of the vertebrobasilar artery may lead to vascular vertigo (26). One study showed that increased vertebrobasilar artery tortuosity was associated with earlier age at dissection and death in children and young adults with connective tissue disorders (16). The findings presented in this study demonstrate the importance of vertebrobasilar radiological evaluations to assess hypoplasia, tortuosity and PWI in patients complaining of isolated positional vertigo/dizziness of unexplained etiology who also present with increased risk for stroke. The findings presented above also show that presence of a fetal posterior cerebral artery was not a significant risk factor for stroke and/or hypoperfusion in the posterior circulation. This effect may be related to the limited number of patients with fetal posterior cerebral arteries included in the study.

This study had several limitations. First, it was a prospective, cross-sectional study that was performed at a single center. A relatively small sample may have contributed to selection bias. Second, the proportion of stroke correlated with isolated vertigo may have been underestimated. The study population did not include patients with history of mild stroke, who were often unwilling to visit the hospital when symptoms had spontaneously resolved. Third, brain MRI may have yielded false-negative results in patients with small infarction lesions; some patients classified as “non-stroke” may actually have had strokes that passed without notice. Fourth, no systematic evaluation was undertaken to assess degree of vertigo or to identify factors that may exacerbate or alleviate symptoms, truncal ataxia, or headache. These important clinical features were therefore not compared between groups. Finally, we were not able to exclude patients with conditions such as basilar migraine, which is also associated with deficits in perfusion.

CONCLUSION

In summary, clinicians should consider the high prevalence of stroke and hypoperfusion in patients with isolated vertigo and vascular risk factors. Some patients presenting with isolated vertigo without lesions in DWI may even have changes in cerebral hemodynamics that can be detected by PWI. Such changes in cerebral hemodynamics may be associated with certain risk factors (e.g., VA stenosis or hypoplasia, VA curvature index). In the future, the link between hypoperfusion and VA hypoplasia or curvature index should be further studied in order to elucidate the etiology of isolated vertigo.

AUTHOR CONTRIBUTIONS

DPZ and HRL designed and conceptualized study, analyzed the data, drafted the manuscript for intellectual content. MZ interpreted the data, revised the manuscript for intellectual content. QKM, SY, YFP, HLZ, and SLZ were major role in the acquisition of data.

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REFERENCES

1. Choi JH, Park MG, Choi SY, Park KP, Baik SK, Kim JS, et al. Acute transient vestibular syndrome: prevalence of stroke and efficacy of bedside evaluation. Stroke (2017) 48:556–62. doi: 10.1161/STROKEAHA.116.015507
2. Li W, Feng Y, Lu W, Xie X, Xiong Z, Jing Z, et al. Evaluating the morphological changes of intracranial arteries and whole-brain perfusion in undetermined isolated vertigo. J Neurosci. (2016) 370:70–7. doi: 10.1016/j.jns.2016.09.024
3. Saber Tehrani AS, Kattah JC, Kerber KA, Gold DR, Zee DS, Urrutia VC, et al. Diagnosing stroke in acute dizziness and vertigo: pitfalls and pearls. Stroke (2018) 49:788–95. doi: 10.1161/STROKEAHA.117.016979
4. Lee H. Isolated vascular vertigo. J Stroke (2014) 16:124–30. doi: 10.5853/jos.2014.16.3.124
5. Choi KD, Lee H, Kim JS. Vertigo in brainstem and cerebellar strokes. Curr Opin Neurol. (2013) 26:90–5. doi: 10.1097/WCO.0b013e32835c5edd
6. Paul NL, Simoni M, Rothwell PM. Oxford Vascular Study. Transient isolated brainstem symptoms preceding posterior circulation stroke: a population-based study. Lancet Neurol. (2013) 12:65–71. doi: 10.1016/S1474-4422(12)70299-5
7. Halmagyi GM. Brainstem stroke preceded by transient isolated vertigo attacks. J Neurol (2017) 264: 2170–7. doi: 10.1007/s00415-017-8610-0
8. Felgueiras R, Magalhães R, Correia M, Silva MC. Long-term prognosis of patients presenting first-ever vestibular symptoms in a community-based study. J Stroke Cerebrovasc Dis. (2014) 23:2190–8. doi: 10.1016/j.jstrokecerebrovasdis.2014.04.033
9. Xu X, Jiang L, Luo M, Li J, Li W, Sheng W. Perfusion-weighted magnetic resonance imaging detects recurrent isolated vertigo caused by cerebral hypoperfusion. Int J Neurosci. (2015) 125:449–55. doi: 10.3109/00207454.2014.947370
10. Leker RR, Hur TB, Gomori JM, Paniri R, Eichel R, Cohen JE. Incidence of DWI-positive stroke in patients with vertigo of unclear etiology, preliminary results. Neurol Res. (2013) 35:123–6. doi: 10.1179/1743132812Y.0000000114
11. Blasberg TF, Wolf L, Henke C, Lorenz MW. Isolated transient vertigo: posterior circulation ischemia or benign origin? BMC Neurol. (2017) 17:111. doi: 10.1186/s12883-017-0894-2
12. Zhang DP, Lu GF, Zhang JW, Zhang SL, Ma QK, Yin S. Vertebral artery hypoplasia and posterior circulation infarction in patients with isolated vertigo with stroke risk factors. J Stroke Cerebrovasc Dis. (2017) 26:295–300. doi: 10.1016/j.jstrokecerebrovasdis.2016.09.020
13. Zhang DP, Ma QK, Zhang JW, Zhang SL, Lu GF, Yin S. Vertebral artery hypoplasia, posterior circulation infarction and relative hypoperfusion detected by perfusion magnetic resonance imaging semiquantitatively. J Neurol Sci. (2016) 368:41–6. doi: 10.1016/j.jns.2016.06.043
14. Nishikata M, Hirashima Y, Tomita T, Futatsuya R, Horige Y, Endo S. Measurement of basilar artery bending and elongation by magnetic resonance cerebral angiography: relationship to age, sex and vertebral artery dominance. Arch Gerontol Geriatr. (2004) 38:251–59. doi: 10.1016/j.archger.2003.10.006
15. Thierfelder KM, Baumann AB, Sommer WH, Armbruster M, Opheark C, Janssen H, et al. Vertebral artery hypoplasia: frequency and effect on cerebellar blood flow characteristics. Stroke (2014) 45:1363–68. doi: 10.1161/STROKEAHA.113.004188
16. Norris SA, Orbach DB, Geva T, Singh MN, Gauvreau K, Lacrovir. Increased vertebral artery tortuosity index is associated with adverse outcomes in children and young adults with connective tissue disorders. Circulation (2011) 124:388–96. doi: 10.1161/CIRCULATIONAHA.110.990549
17. Kim SH, Kim HJ, Kim JS. Isolated vestibular syndromes due to brainstem and cerebellar lesions. J Neurol. (2017) 264:83–9. doi: 10.1007/s00415-017-8455-6
18. Kerber KA, Brown DL, Lisabeth LD, Smith MA, Morgenstern LB. Stroke among patients with dizziness, vertigo, and imbalance in the emergency department: a population-based study. Stroke (2006) 37:2484–7. doi: 10.1161/01.STR.0000240329.48263.0d
19. Fernández L, Breinbauer HA, Delano PH. Vertigo and dizziness in the elderly. Front Neurol. (2015) 6:144. doi: 10.3389/fneur.2015.00144
20. Lee JO, Park SH, Kim HJ, Kim MS, Park BR, Kim JS. Vulnerability of the vestibular organs to transient ischemia: implications for isolated vascular vertigo. Neurosci Lett. (2014) 558:180–5. doi: 10.1016/j.neulet.2013.11.016
21. Lee CC, Su YC, Ho HC, Hung SK, Lee MS, Chou P, et al. Risk of stroke in patients hospitalized for isolated vertigo: a four-year follow-up study. Stroke (2011) 42:48–52. doi: 10.1161/STROKEAHA.110.597070
22. Palacio S, McClure LA, Benavente OR, Bazan C III, Pergola P, Hart RG. Lacunar strokes in patients with diabetes mellitus: risk factors, infarct location, and prognosis: the secondary prevention of small subcortical strokes study. Stroke (2014) 45:2689–94. doi: 10.1161/STROKEAHA.114.005018
25. Ichikawa H, Mukai M, Hieda S, Kamiya Y, Akizawa T, Kawamura M. Involvement of the basilar artery in diabetes mellitus: an MRI study of brainstem infarctions. Eur Neurol. (2010) 64:230–5. doi: 10.1159/000319924
26. Zhang HT, Zhang SL, Zhang DP. Two case reports of bilateral vertebral artery tortuosity and spiral twisting in vascular vertigo. BMC Neurol. (2014) 14:14. doi: 10.1186/1471-2377-14-14

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