Prevalence and outcome of young stroke patients with middle cerebral artery stenosis

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Abstract

Background: Etiologies of acute ischemic stroke in young adults are heterogeneous. Middle cerebral artery (MCA) stenosis is a common finding in Asians which may be an important cause of stroke in young adults. However, studies of stroke in young Asian populations are rare. Our study was to investigate the prevalence and outcome of young stroke patients with MCA stenosis in Chinese populations.

Methods: Young patients with MCA territory infarction between January 2013 and September 2018 were retrospectively recruited. Subjects were defined as stenosis group (MCA stenosis ≥50%) and no-stenosis group (MCA stenosis<50% or no stenosis) by their MCA stenosis. For patients in stenosis group, they were categorized as uni-MCA stenosis subgroup and multiple stenosis subgroup. Demographic data, risk factors, imaging feature and complications were compared between groups. Prevalence of MCA stenosis and risk factor score (score ≥ 2 or 3) in different age groups were investigated. Modified Rankin Scale (mRS) was used for evaluating functional outcome at discharge (unfavorable outcome: 3–6). Binary logistic regression was performed to determine independent risk factors of unfavorable outcome.

Results: Two hundred forty-nine young stroke patients were included in our study and 110 (44.2%) patients were defined as stenosis group. 55 (50%) patients were categorized as uni-MCA stenosis subgroup and 55 (50%) were multiple stenosis subgroup. The most common traditional vascular risk factors included hypertension, hyperlipemia, smoking, hyperhomocysteinemia and alcohol consumption. Prevalence of risk factor score ≥ 2 or 3 increased with age, but not incidence of MCA stenosis. By TOAST classification, the most common etiologies were large-artery atherosclerosis (41.0%) and small vessel disease (33.7%). Compared with no-stenosis group, patients in stenosis group were more likely to have large territorial infarct, develop complications and have unfavorable outcome. No significant difference was found between patients in uni-MCA stenosis and multiple stenosis subgroups except history of stroke/TIA, risk factor score ≥ 3 and silent infarct. By logistic regression, hypertension (OR = 3.561; 95%CI, 1.494 to 8.492; *p* = 0.004), NIHSS scores at admission (OR = 1.438; 95%CI, 1.276 to 1.620; *p* = 0.000) and infarct size (*p* = 0.015) independently predicted unfavorable outcome.

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Background
In contrast with the decreasing incidence of stroke in older adults, an increasing incidence and hospitalization rates for acute ischemic stroke in young adults have been identified [1]. It has been estimated that the incidence of stroke in young adults increased up to 40% worldwide over the past decades, which has great impact because of high hospitalization costs and loss of the ability to work [1]. Etiologies of young stroke are heterogeneous, which leads to different strategies in treatment and secondary prevention. According to Trial of Org 10, 172 in Acute Stroke Treatment (TOAST) classification, other determined (21.6–26.0%) and undetermined etiology (33.0–39.6%) were more common in young stroke patients, while stroke caused by large-artery atherosclerosis (LAA) was only about 7.5–9.3% [2, 3]. However, these data was mainly collected from European populations. In a study involving 123 Chinese young adults, LAA and small vessel disease (SVD) were the most common causes of acute ischemic stroke [4]. Other studies also showed that proportion of LAA was higher in Asians than that in western populations, indicating that cerebral artery stenosis may be an important cause of stroke in Asian populations [5–7].

Prevalence of cerebral artery stenosis varies by their locations. Risk of intracranial arteries stenosis is much higher than that in extracranial arteries in Asian populations. It has been estimated that intracranial arteries stenosis may contribute to 30 to 50% of ischemic strokes, and middle cerebral artery (MCA) was the most commonly affected vessel [8]. Moreover, symptomatic intracranial artery stenosis seems to be relatively unstable because of high frequency of progression [9]. However, studies of young stroke patients with intracranial artery stenosis are rare. Prevalence and causes of intracranial artery stenosis in young stroke patients are also largely unknown. Relationship between intracranial artery stenosis and clinical outcome in young stroke patients are still uncertain. Therefore, our study focused on young Chinese patients with MCA territory infarction. The goal of the paper was to investigate the prevalence and clinical outcome of the MCA stenosis. We investigated clinical characteristics, etiologies and functional outcome of young stroke patients with MCA stenosis. The possible mechanisms of MCA stenosis were also explored in our study.

Methods
Subject population
Patients with acute ischemic stroke, admitted to Shandong Provincial Qianfoshan Hospital, Shandong University between January 2013 and September 2018, were retrospectively recruited. Patients fulfilled the following inclusion criteria were included: (1) age 18–55 years; (2) acute ischemic stroke in MCA territory confirmed by diffusion weighted imaging (DWI); (3) adequate assessment of cerebral arteries by magnetic resonance angiography (MRA) or computed tomography angiography (CTA) or digital subtraction angiography (DSA); (4) informed consent obtained from all subjects to participate in the study by telephone. As our study focused on young stroke patients with MCA stenosis, patients with ischemic stroke in other vascular territories were not included except watershed infarct between MCA and anterior cerebral artery or posterior cerebral artery. Those young stroke patients without available clinical information were also excluded in our study. In order to observe incidence of MCA stenosis in different age group, we used the upper age limit of 55 years to define young stroke [10, 11]. The study protocol was approved by the Ethics Committee of Shandong Provincial Qianfoshan Hospital, Shandong University. Treatments of acute ischemic stroke were according to stroke early management guidelines.

Clinical information
Medical records and brain imagings of included patients were reviewed by trained neurologists. We extracted data on patient demographics, relevant medical history and neurological severity in a standard form. We recorded and scored traditional vascular risk factors, including hypertension (history/treatment or persistently elevated blood pressure: systolic ≥140 mmHg; diastolic ≥90 mmHg), diabetes mellitus (history/treatment or diagnosed at discharge), coronary artery disease (history/treatment or diagnosed at discharge), atrial fibrillation (history/treatment or diagnosed at discharge), hyperlipidemia (history/treatment or low-density lipoprotein cholesterol ≥2.6 mmol/L on admission), hyperhomocysteinemia (history/treatment or diagnosed at discharge), smoking, alcohol consumption, history of transient ischemic attack (TIA) or stroke and family history of stroke. We also collected “rare” risk factors of stroke in...
young patients, including history of other cardiac disease, peripheral arterial diseases, sleep apnea, autoimmune disease (including Systemic lupus erythematosus (SLE), Rheumatoid arthritis (RA), Sjogren’s syndrome and vasculitis), migraine, abuse of illicit drugs, pregnancy and use of oral contraceptive pills. Etiologies of stroke in young patients were assigned by two reviewers who were unknown to the aim of our study, using TOAST classification [12]. Conflicting results were resolved by consensus.

Clinical outcome and complications
National Institutes of Health Stroke Scale (NIHSS) was used to assess the stroke severity at admission and discharge. Modified Rankin Scale (mRS) was used for evaluating neurologic disability (Score range: 0–6), with mRS score of 0 to 2 representing a favorable outcome and score of 3 to 6 representing an unfavorable functional outcome. Complications of stroke were defined by medical records, including pulmonary infection, urinary tract infection, deep vein thrombosis, stress ulcer, epilepsy, hemorrhagic transformation and myocardial infarction. The diagnosis of complications had been reported in a previous study [13].

Imaging analysis
All patients underwent magnetic resonance (MR) assessment on a 1.5 T MR scanner (Philips; Holland) within 1 week of onset. Imaging sequences included axial T2-weighted imaging, T1-weighted imaging, fluid-attenuated inversion recovery (FLAIR) sequences and DWI. Based on standard templates, infarct size was defined as: (1) small infarct (maximum diameter<1.5 cm); (2) medium infarct (lesion in a cortical superficial branch of MCA, or lesion involving the MCA deep branch, or lesion in border-zone of MCA); (3) large territorial infarct (lesion involving complete territory of MCA or lesion involving 2 cortical superficial branches of MCA or lesion involving a cortical superficial branch of MCA associated to the MCA deep branch) [14]. White matter lesions were assessed by Fazekas scores (mild: 0 to 2; moderate: 3 to 4; severe: 5 to 6). Silent brain infarcts (SBIs) were defined as focal hyperintensities on T2 and FLAIR sequences, 3 mm in diameter, without corresponding neurologic symptoms.

Assessment of cerebral arteries was conducted by MRA, CTA or DSA. Stenosis of cerebral artery was calculated using previous published method in the Warfarin Aspirin Symptomatic Intracranial Disease Study (WASID) [15]. Stenosis was defined as narrowing of the lumen ≥50%. Cerebral artery stenosis was measured separately by two experienced neurologists who were blind to clinical information. Conflicting results were resolved by consensus. According to MCA stenosis, included subjects were defined as stenosis group (stenosis ≥50%) and no-stenosis group (stenosis<50% or no stenosis). For patients in stenosis group, they were categorized as uni-MCA stenosis subgroup and multiple stenosis subgroup (stenosis in MCA and other cerebral artery). Patients suspected of cardioembolism would perform 24 h-electrocardiogram (ECG) monitoring, transthoracic or transoesophageal echocardiography. In our study, 54 (21.7%) patients conducted assessment of transthoracic or transoesophageal echocardiography.

Statistic analysis
SPSS for windows 17.0 was used for data storage and statistical analysis. All statistics were presented as mean and SD for continuous variables with normal distribution, median and interquartile range for continuous variables with non-normal distribution, frequency and percentages for categorical variables. Univariate analysis was performed between stenosis group and no-stenosis group. And univariate analysis was also performed between stenosis subgroups. Continuous variables with normal distribution were compared with Student t test with significance set at p < 0.05, while Wilcoxon rank sum test for continuous variables with non-normal distribution. Categorical variables were compared by means of χ² test or fisher’s exact, Wilcoxon rank sum test. Binary logistic regression was performed to determine risk factors of unfavorable functional outcome.

Results
During the study period, 562 young stroke patients were identified. Sixty-two patients without adequate assessment of cerebral arteries and 251 patients with stroke which was not located in MCA territory were excluded. Finally, 249 young stroke patients were included in our study, with a mean age of 48 ± 6 years. In those included patients, 79 (31.7%) were female. Demographic and clinical characteristics of included patients were shown in Table 1. The most frequent traditional vascular risk factors were hypertension (58.6%), hyperlipemia (55.0%), smoking (41.0%), hyperhomocysteinemia (36.1%) and alcohol consumption (33.7%). In rare risk factors, autoimmune disease (3.6%) was the most commonly detected. Only 16 (6.4%) patients presented with wake-up stroke and 56 (22.5%) patients had large territorial infarct. Cerebral arteries were assessed by MRA in 206 patients, CTA in 26 patients and DSA in 55 patients. Thirty-three patients had both MRA and DSA and 5 patients had both CTA and DSA. One hundred and ten (44.2%) patients had MCA stenosis and defined as stenosis group. For those in stenosis group, 55 (50%) patients were classified as uni-MCA stenosis subgroup and 55 (50%) were multiple stenosis subgroup. By age groups, prevalence of patients with risk factor score ≥2 or 3
Table 1  Demographic and characteristics of young adults with MCA territory infarction

|                      | ALL (n = 249) | No-stenosis group (n = 139) | Stenosis group (n = 110) | P  | Uni-MCA stenosis subgroup (n = 55) | Multiple stenosis subgroup (n = 55) | P   |
|----------------------|---------------|-----------------------------|--------------------------|----|-----------------------------------|-------------------------------------|-----|
| Age (years)          | 48.0 ± 6.0    | 47.8 ± 5.7                  | 48.3 ± 6.4               | 0.151 | 47.5 ± 7.7                       | 49.1 ± 4.7                        | 0.723 |
| Sex (male, %)        | 170 (68.3%)   | 96 (69.1%)                  | 74 (67.3%)               | 0.763 | 38 (69.1%)                       | 36 (65.5%)                        | 0.684 |
| Traditional risk factors |             |                             |                          |     |                                   |                                     |     |
| Hypertension         | 146 (58.6%)   | 86 (61.9%)                  | 60 (54.5%)               | 0.244 | 25 (45.5%)                       | 35 (63.6%)                        | 0.056 |
| Diabetes mellitus    | 73 (29.3%)    | 34 (24.5%)                  | 39 (35.5%)               | 0.508 | 17 (30.9%)                       | 22 (40.0%)                        | 0.319 |
| CAD                  | 22 (8.8%)     | 14 (10.1%)                  | 8 (7.3%)                 | 0.440 | 3 (5.5%)                         | 5 (9.1%)                          | 0.714 |
| AF                   | 3 (1.2%)      | 1 (0.7%)                    | 2 (1.8%)                 | 0.838 | 1 (1.8%)                         | 1 (1.8%)                          | 1.000 |
| Hyperlipidemia       | 137 (55.0%)   | 83 (59.7%)                  | 54 (49.1%)               | 0.094 | 27 (49.1%)                       | 27 (49.1%)                        | 1.000 |
| Hyperhomocysteinemia | 90 (36.1%)    | 53 (38.1%)                  | 37 (33.6%)               | 0.464 | 19 (34.5%)                       | 18 (32.7%)                        | 0.840 |
| Alcohol consumption  | 84 (33.7%)    | 47 (33.8%)                  | 37 (33.6%)               | 0.927 | 14 (25.5%)                       | 23 (41.8%)                        | 0.069 |
| Smoking              | 102 (41%)     | 55 (39.6%)                  | 47 (42.7%)               | 0.615 | 25 (45.5%)                       | 22 (40.0%)                        | 0.563 |
| TIA/stroke           | 54 (21.7%)    | 26 (18.7%)                  | 28 (25.5%)               | 0.199 | 6 (10.9%)                        | 22 (40.0%)                        | 0.000 |
| Family history of stroke | 48 (19.3%)   | 25 (18.0%)                  | 23 (20.9%)               | 0.561 | 12 (21.8%)                       | 11 (20.0%)                        | 0.815 |
| Score of traditional risk factors |           |                             |                          |     |                                   |                                     |     |
| ≥ 2                  | 208 (83.5%)   | 115 (82.7%)                 | 93 (84.5%)               | 0.702 | 43 (78.2%)                       | 50 (90.9%)                        | 0.065 |
| ≥ 3                  | 152 (61.0%)   | 87 (62.6%)                  | 65 (59.1%)               | 0.574 | 28 (50.9%)                       | 47 (85.5%)                        | 0.000 |
| Rare risk factors    |              |                             |                          |     |                                   |                                     |     |
| Other cardiac disease| 1 (0.4%)      | 0                            | 1 (0.4%)                 | 0    | –                                 | –                                   | –    |
| PAD                  | 2 (0.8%)      | 2 (0.8%)                    | 0 (0.0%)                 | 0    | –                                 | –                                   | –    |
| Sleep apnea          | 1 (0.4%)      | 1 (0.4%)                    | 0 (0.0%)                 | 0    | –                                 | –                                   | –    |
| Migraine             | 1 (0.4%)      | 0                            | 1 (0.4%)                 | 0    | 1 (0.4%)                         | –                                   | –    |
| Autoimmune disease   | 9 (3.6%)      | 6 (4.3%)                    | 3 (2.7%)                 | 0    | 3 (5.5)                          | –                                   | –    |
| Wake-up stroke       | 16 (6.4%)     | 7 (5.0%)                    | 9 (8.2%)                 | 0.315 | 4 (7.3%)                         | 5 (9.1%)                          | 1.000 |
| NIHSS at admission   | 4 (1, 8)      | 3 (1, 6)                    | 6 (2, 11)                | 0.000 | 6 (2, 12)                        | 5 (2, 11)                         | 0.827 |
| Infarct size         |              |                             |                          |     |                                   |                                     |     |
| small infarct        | 111 (44.6%)   | 89 (64.0%)                  | 22 (20.0%)               | 0.000 | 8 (14.5%)                        | 14 (25.5%)                        | 0.221 |
| medium infarct       | 82 (32.9%)    | 44 (31.7%)                  | 38 (34.5%)               | 0.385 | 18 (32.7%)                       | 20 (36.4%)                        | 0.000 |
| large territorial infarct | 56 (22.5%) | 6 (4.3%)                    | 50 (45.5%)               | 0.000 | 29 (52.7%)                       | 21 (38.2%)                        | 0.000 |
| Silent brain infarcts|              |                             |                          |     |                                   |                                     |     |
| 0                    | 133 (53.4%)   | 71 (51.5%)                  | 62 (56.4%)               | 0.385 | 43 (78.2%)                       | 19 (34.5%)                        | 0.000 |
| 1                    | 30 (12.1%)    | 15 (10.8%)                  | 15 (13.6%)               | 5    | 5 (9.1%)                         | 10 (18.2%)                        | 0.000 |
| ≥ 2                  | 86 (34.5%)    | 53 (38.1%)                  | 33 (30.0%)               | 7    | 7 (12.7%)                        | 26 (47.3%)                        | 0.000 |
increased with age and mild decreased at age of 50 to 55 years, while incidence of MCA stenosis remained stable and declined at age of 45 to 49 years. (Fig. 1).

According to TOAST classification, 102 (41.0%) patients were defined as LAA, 84 (33.7%) patients were SVD, 6 (2.4%) patients were cardioembolism, 23 (9.2%) patients were other determined and 34 (13.7%) patients were cryptogenic stroke. Details of cardioembolism and other determined stroke were presented in Table 2. Patent foramen ovale was the most common cause in patients with cardioembolism. Moya-moya disease was the most common etiology of other determined stroke, which was followed by dissection. (Fig. 2) For patients in stenosis group (n = 110), 80 (72.7%) patients were LAA, 7 (6.4%) patients were SVD, 1 (0.9%) patient was cardioembolism, 13 (11.8%) patients were other determined and 9 (8.2%) patients were cryptogenic stroke. For patients in no-stenosis group (n = 139), 22 (15.8%) patients were LAA, 77 (55.4%) patients were SVD, 5 (3.6%) patients were cardioembolism, 10 (7.2%) patients were other determined and 25 (18.0%) patients were cryptogenic stroke. (Table 2).

Compared with patients in no-stenosis group, patients in stenosis group had higher incidence of diabetes mellitus but without significance (35.5% vs 24.5%, p = 0.058). (Table 1) Patients in stenosis group were more likely to have large territorial infarct (45.5% vs 4.3%, p = 0.000) and presented with higher NIHSS score at admission (mean: 6 vs 3, p = 0.000). Regarding the clinical outcome, patients in stenosis group had higher risk of developing complications (15.5% vs 5.0%, p = 0.006) and presenting with poor functional outcome at discharge (mRS > 2: 38.2% vs 15.8%, p = 0.000).

### Table 1 Demographic and characteristics of young adults with MCA territory infarction (Continued)

|                  | ALL (n = 249) | No-stenosis group (n = 139) | Stenosis group (n = 110) | P     | Uni-MCA stenosis subgroup (n = 55) | Multiple stenosis subgroup (n = 55) | P     |
|------------------|--------------|----------------------------|--------------------------|-------|-----------------------------------|-------------------------------------|-------|
|                   |              |                            |                          |       |                                   |                                     |       |
| White matter lesions |              |                            |                          |       |                                   |                                     |       |
| 0–2 | 168 (67.5%) | 92 (66.2%) | 76 (69.1%) | 0.626 | 42 (76.4%) | 34 (60.7%) | 0.182 |
| 3–4 | 63 (25.3%)  | 35 (25.2%) | 28 (25.5%) | 11 (20.0%) | 17 (30.4%) |
| 5–6 | 18 (7.2%)   | 12 (8.6%)  | 6 (5.5%)    | 2 (3.6%)   | 5 (8.9%)   |
| Complications |                   |                            |                          | 0.006 | 8 (14.5%) | 9 (16.4%) | 0.792 |
| NIHSS at discharge |              |                            |                          | 0.000 | 4 (0, 8)   | 4 (1, 7)   | 0.904 |
| mRS at discharge |                   |                            |                          |       | 3 (1, 8)    | 5 (2, 10)     | 0.164 |
| 0–2 | 185 (74.3%) | 117 (84.2%) | 68 (61.8%) | 0.000 | 34 (61.8%) | 34 (61.8%) | 1.000 |
| 3–6 | 64 (25.7%)  | 22 (15.8%) | 42 (38.2%) | 21 (38.2%) | 21 (38.2%) |

CAD indicates coronary artery disease, AF indicates atrial fibrillation, TIA indicates transient ischemic attack, PAD indicates peripheral arterial diseases, NIHSS indicates National Institutes of Health Stroke Scale, mRS indicates modified Rankin Scale.

![Prevalence of risk factors score and MCA stenosis](image_url)
In comparison between subgroups of patients with MCA stenosis, patients in multiple stenosis subgroup were more likely to have history of stroke/TIA (40.0% vs 10.9%, \( p = 0.000 \)) and higher incidence of traditional risk factor score \( \geq 3 \) (85.5% vs 50.9%, \( p = 0.000 \)). (Table 1) They also had more silent infarcts on brain MR imaging (65.5% vs 21.8%, \( p = 0.000 \)). No significant difference was found between two subgroups in stroke severity, complications and functional outcome.

Sixty-four (25.7%) patients had unfavorable functional outcome at discharge. In order to investigate independent risk factors for unfavorable functional outcome, binary logistic regression was performed, with age, sex, hypertension, risk factor score, wake-up stroke, NIHSS score at admission, infarct size, MCA stenosis and complications in the model. Hypertension (OR = 3.561; 95% CI, 1.494 to 8.492; \( p = 0.004 \)), NIHSS scores at admission (OR = 1.438; 95% CI, 1.276 to 1.620; \( p = 0.000 \)) and infarct size (\( p = 0.015 \)) independently predicted unfavorable functional outcome. (Table 3).

**Discussion**

Intracranial artery stenosis is an important cause of stroke in young adults. In our study, we found that traditional vascular risk factors were common in young stroke patients and the most frequent traditional etiologies were LAA and SVD. 44.2% young patients with MCA territory infarction had MCA stenosis, while incidence of MCA stenosis did not increase with age. Patients with MCA stenosis had worse clinical outcome, however, only hypertension, NIHSS scores at admission and infarct size were independent predictors.

The role of traditional vascular risk factors in young stroke has received more attention in recent years. Prevalence of those factors in young patients has increased over the past 20 y. In a German nationwide case-control study involving 2125 young stroke patients, eight cardiovascular risk factors combined could explain 78.9% of all strokes [16]. In our study, we found that 83.5% patients has two or more traditional vascular risk factors and the most frequent traditional vascular risk factors include hypertension, hyperlipemia, smoking and alcohol consumption, which is consistent with previous studies [4, 17]. We also found that hyperhomocysteinemia, an independent risk factor of stroke, is prevalent in young stroke patients, which has rarely been reported. For rare factors, there are differences between young European and Chinese populations. The most prevalent “rare” risk factors for stroke in young European populations include migraine, illicit drug use, patent foramen ovale, oral contraceptives and pregnancy or puerperium [17]. In our study, autoimmune disease was more common in recruited subjects. Conflicting results may be attributed to different included patients. In our study, only one patient with migraine was included because of positive presentation on DWI and patients with patent foramen ovale were not included if infarction was not located in MCA territory.

Identification of underlying causal etiology is essential for treatment and secondary prevention in young stroke patients. However, studies in Chinese populations are

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**Table 2** Etiologies of young stroke patients according to TOAST classification

| TOAST subtypes                  | ALL (n = 249) | Stenosis group (n = 110) | No-stenosis group (n = 139) |
|--------------------------------|---------------|-------------------------|----------------------------|
| Large artery atherosclerosis   | 102 (41.0%)   | 80 (72.7%)              | 22 (15.8%)                 |
| Small vessel disease           | 84 (33.7%)    | 7 (6.4%)                | 77 (55.4%)                 |
| Cryptogenic etiology           | 34 (13.7%)    | 9 (8.2%)                | 25 (18.0%)                 |
| Other determined etiology      | 23 (9.2%)     | 13 (11.8%)              | 10 (7.2%)                  |
| Moyamoya disease               | 11 (47.8%)    | 9 (69.2%)               | 2 (20.0%)                  |
| Dissection                     | 4 (17.4%)     | 2 (15.4%)               | 2 (20.0%)                  |
| Radiation vasculopathy         | 2 (8.7%)      | 1 (7.7%)                | 1 (10.0%)                  |
| Cerebral amyloid angiopathy    | 2 (8.7%)      | 0                       | 2 (20.0%)                  |
| Other vasculitis               | 2 (8.7%)      | 0                       | 2 (20.0%)                  |
| Infectious vasculitis          | 1 (4.3%)      | 1 (7.7%)                | 0                          |
| Metabolic Encephalopathy       | 1 (4.3%)      | 0                       | 1 (10.0%)                  |
| Cardioembolism                 | 6 (2.4%)      | 1 (0.9%)                | 5 (3.6%)                   |
| Patent foramen ovale           | 3 (50%)       | 0                       | 3 (60.0%)                  |
| Atrial fibrillation            | 1 (16.7%)     | 0                       | 1 (20.0%)                  |
| Myocardial infarction          | 1 (16.7%)     | 0                       | 1 (20.0%)                  |
| Dilated cardiomyopathy         | 1 (16.7%)     | 1 (100%)                | 0                          |

TOAST indicates Trial of Org 10,172 in Acute Stroke Treatment
artery stenosis in young patients is challenging. High-epidemiological studies are still needed in China. mainly come from Korean and Japanese. Larger national Asian countries [18]. These epidemiologic datas were finding that moyamoya disease is more prevalent in of dissection in our study, which may be related with the incidence of moyamoya disease was much higher than that patients in previous studies [2, 3, 17]. However, the incidence of MCA stenosis and our study revealed that 44.2% patients had MCA stenosis [4]. The prevalence of SVD varies among studies, which has not been completely understood. More epidemiologic studies should be conducted. In other determined etiologies, dissection was the most common reported cause for young stroke patients with MCA territory infarction were LAA and SVD. These results are consistent with another Chinese study in which 35.8% patients were LAA and 43.9% patients were SVD [4]. An explanation may be that high frequency of LAA is related with high rate of intracranial artery stenosis in Chinese populations. As the above-mentioned Chinese study showed that 43.1% young stroke patients had multiple intracranial artery stenosis and our study revealed that 44.2% patients had MCA stenosis [4]. The prevalence of SVD varies among studies, which has not been completely understood. More epidemiologic studies should be conducted. In other determined etiologies, dissection was the most common reported cause for young stroke patients in previous studies [2, 3, 17]. However, the incidence of moyamoya disease was much higher than that of dissection in our study, which may be related with the finding that moyamoya disease is more prevalent in Asian countries [18]. These epidemiologic datas were mainly come from Korean and Japanese. Larger national epidemiological studies are still needed in China. Identification of the exact mechanism of intracranial artery stenosis in young patients is challenging. High-resolution magnetic resonance imaging (HR-MRI) may be helpful in distinguishing characters of stenosis by offering arterial wall imaging. HR vessel wall MRI (HR-VW MR) imaging of intracranial atherosclerotic plaque typically demonstrates arterial wall thickening, which eccentrically (not uniformly) involves the circumference of the arterial wall and central nervous system (CNS) vasculitis usually show homogeneous concentric arterial wall thickening and enhancement [19]. In a prospective study involving 95 young Korean patients with unilateral MCA stenosis and no or minimal (≤1) atherosclerotic risk factors, 26 (27.4%) patients were categorized as atherosclerotic disease, 29 (30.5%) were moyamoya disease, 22 (23.2%) were dissection and 18 (18.9%) were vasculitis [20]. However, another Chinese HR-MRI study showed that 98 (80.3%) young patients with unilateral MCA stenosis had eccentric stenosis, showing that atherosclerosis is a common cause of intracranial stenosis in young adults [21]. These results should be interpreted with caution, for patients in Korean study had fewer atherosclerotic risk factors and patients diagnosed as vasculitis or dissection were not excluded in Chinese study [20, 21]. In our study, we found that prevalence of patients with score of risk factors > 2 or 3 increased with age, but incidence of MCA stenosis remained stable and declined at 45 to 49 years. The conflicting tendency between traditional vascular risk factor and MCA stenosis may indicate that non-atherosclerotic factors are still important causes of MCA stenosis in younger stroke patients (< 45 years). Prospective cohort studies by HR-MRI should be conducted to investigate the effect of traditional vascular risk factors on intracranial artery stenosis. Young stroke patients usually have favorable clinical outcome. A previous study reported that 81% of individuals with young stroke had good outcome (mRS < 3) at hospital discharge, which is consistent with our result. It has been estimated that proportions of individuals with good functional outcome among young stroke survivors range from 80 to 94% after a mean follow-up duration of 3–12 years [17]. In our study, we found that patients with MCA stenosis had poor clinical outcome, which may be related to their larger infact size. Hypertension, NIHSS scores at admission and lesion size independently predicted poor clinical outcome, which is consistent with

Table 3 Binary logistic regression for unfavorable functional outcome at discharge (mRS: 3–6)

| Variables                  | n   | Unfavorable functional outcome | β    | p value  | Exp(B) 95% CI |
|----------------------------|-----|--------------------------------|------|----------|------------|
| Hypertension               | 249 |                                | 1.270| 0.004    | 3.561 (1.494, 8.492) |
| NIHSS scores at admission  | 249 |                                | 0.363| 0.000    | 1.438 (1.276, 1.620) |
| small infarct              | 249 |                                | 0.015| 0.015    | 1.101 (0.992, 1.225) |
| medium infarct             | 249 |                                | 0.005| 0.005    | 1.015 (1.001, 1.030) |
| large territorial infarct  | 249 |                                | 0.044| 0.044    | 1.048 (1.001, 1.098) |

NIHSS indicates National Institutes of Health Stroke Scale, mRS indicates modified Rankin Scale.
previous studies. The relationship between hypertension and poor clinical outcome may be induced by blood pressure variability which is closely related with early neurological deterioration [22].

The strength of our study included the homogeneity of our recruited young patients. Several limitations should be addressed in our study. First, our study was a retrospective study and conducted in single center with inevitable selected bias. Second, only patients with stroke in MCA territory were included in our study which may reduce the external validity of our study. Third, number of patients with adequate assessment of cardioembolism in our study was relatively small, which may underestimate cardioembolism in young stroke patients. However, patients with cardioembolism usually present with multi-infarctions in multi-vascular territories. As our study only focus on MCA stroke, patients with those MR features were excluded which may eliminate the impact of inadequate assessment of cardioembolism. In addition, the majority of our patients were assessed by MRA which may have exaggerated effect on intracranial stenosis. As our study was not a HR-MRI study, we could only provide limited information about the exact mechanisms of MCA stenosis.

Conclusions

44.2% young Chinese patients with MCA territory infarction had MCA stenosis. Prevalence of MCA stenosis did not increase with age. Patients with MCA stenosis had worse clinical outcome, however, only hypertension, NIHSS scores at admission and infarct size were independent predictors.

Abbreviations

MCA: Middle cerebral artery; mRS: Modified Rankin Scale; TOAST: Trial of Org 10,172 in Acute Stroke Treatment; NIHSS: National Institutes of Health Stroke Scale; TIA: Transient ischemic attack; LAA: Large-artery atherosclerosis; SVD: Small vessel disease; DWI: Diffusion weighted imaging; MRA: Magnetic resonance angiography; CTA: Computed tomography angiography; DSA: Digital subtraction angiography; MR: Magnetic resonance; FLAIR: Fluid-attenuated inversion recovery; SBI: Silent brain infarcts; WASID: Warfarin Aspirin Symptomatic Intracranial Disease Study; ECG: Electrocardiogram; HR-MRI: High-resolution magnetic resonance imaging; HR-VW MR: HR vessel wall MRI; CNS: Central nervous system

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Statement

All methods were carried out in accordance with relevant guidelines.

Authors’ contributions

XWJ and ZXY collected data and wrote the manuscript. ZMJ designed the study and revised the manuscript. CH and ZZN participated in analysis of data. All authors read and approved the final manuscript.

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Availability of data and materials

All data generated or analysed during this study are included in this published article and have consent from all patients to publish this data.

Declarations

Ethics approval and consent to participate

The study protocol was approved by the Ethics Committee of Shandong Provincial Qianfoshan Hospital, Shandong University. Informed consent obtained from all subjects to participate in the study.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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