



# Hyperhomocysteinemia Independently Associated with Adult Moyamoya Disease: Hospital Based Study of 237 Patients

Ying ZHANG<sup>1</sup>, Xian FU<sup>2</sup>, Xiangfan ZENG<sup>1</sup>, Jie XU<sup>1</sup>, Hongying LIU<sup>1</sup>, Xuelong LI<sup>1</sup>, Qingchun GAO<sup>1</sup>

<sup>1</sup>Guangzhou Medical University, The Second Affiliated Hospital, Department of Neurology and Institute of Neuroscience, Guangzhou, China

<sup>2</sup>Shenzhen Baoan District Songgang People's Hospital, Department of Neurology, Shenzhen, China

Corresponding author: Xian FU ✉ fuxian2011@163.com

## ABSTRACT

**AIM:** To clarify the risk factors for adult moyamoya disease (MMD) in patients from South China.

**MATERIAL and METHODS:** We prospectively studied adult patients who were diagnosed angiographically with MMD. The demographic profiles, medical history and clinical characteristics were compared between adult MMD and non-MMD stroke patients. Logistic regression analysis was used to determine the risk factors associated with adult MMD.

**RESULTS:** A total of 35 adult MMD patients and 202 adults patients with non-MMD stroke were included. Of the 35 MMD patients, bilateral MMD occurred in 48.6% and bypass surgery was performed in 28.6%; these figures were significantly lower than those reported in patients from Korea and the United States ( $p < 0.05$ ). After adjusting for baseline demographics and potential confounders, multivariate logistic regression analysis was conducted, which showed that the plasma homocysteine level (odds ratio [OR]: 1.10; 95% confidence interval [CI]: 1.06–1.14) and occupation as a technological worker (OR: 4.23; 95% CI: 1.65–10.89) were independently associated with adult MMD.

**CONCLUSION:** Hyperhomocysteinemia and type of occupation were found to be independent risk factors for adult MMD in patients from South China. However, there is still a need for further research to clarify the pathogenesis of MMD. Given the lack of understanding about the risk factors and prevention measures for MMD, we suggest bypass surgery be used for MMD treatment in clinical practice in China to achieve more desirable effects in the management of the disease.

**KEYWORDS:** Moyamoya disease, Angiography, Digital subtraction, Hyperhomocysteinemia, Adult

**ABBREVIATIONS:** MMD: Moyamoya disease, DSA: Digital subtraction angiography, HDL: High-density lipoprotein, LDL: Low-density lipoprotein, ApoA1: Apolipoprotein-A1, ApoB: Apolipoprotein-B, MTHFR: Methylene tetrahydrofolate reductase

## INTRODUCTION

Moyamoya disease (MMD) is characterised by compensatory irregular perforating vessels (i.e., 'moyamoya vessels') that are generated in parallel with bilateral angiopathy of intracranial vascular networks, and are located near the occluded or stenotic regions, corresponding to the lenticulostriate and thalamoperforating arteries (28). The outgrowth of small vessels produces a hazy

image described as a 'puff of smoke' on the radiograph, hence the disease's name *moyamoya* in Japanese (28). The intrinsic nature of MMD is to convert the brain's vascular supply from the internal carotid system to the external carotid system. MMD is a rare disease in most parts of the world, except for Japan and other areas in Asia (31). It has recently been revealed that the age of onset of MMD has two peaks (10). The first peak is observed around 10 years of age, and the second peak occurs between 30 and 45 years of age.

Ying ZHANG  : 0000-0003-4444-8483  
Xian FU  : 0000-0002-7982-1092  
Xiangfan ZENG  : 0000-0002-0680-1859

Jie XU  : 0000-0001-9293-6890  
Hongying LIU  : 0000-0003-0463-9744  
Xuelong LI  : 0000-0002-3624-5160

Qingchun GAO  : 0000-0002-8014-6493

Numerous studies have been conducted to characterise MMD in patient populations from East Asia (7,11,12,15), and Western countries (14,18), but the pathogenesis of MMD remains largely unknown. Inflammation and immune dysfunction have been implicated in its pathogenesis (27-29); however, the history of such events is often subjective and unreliable. Tuberculous meningitis, atherosclerosis, neurofibromatosis and irradiation have all been reported to be causes of MMD (3), and a congenital cause has also been suggested (13,17). Recent studies have shown that concentrations of certain growth factors or cytokines are increased in the cerebrospinal fluid of patients with MMD. However, there is no internationally accepted aetiology of MMD. Therefore, in this study, we aimed to clarify the risk factors and potential pathogenesis of MMD in adult patients from South China.

## ■ MATERIAL and METHODS

### Patient and Public Involvement

This research was conducted without patient involvement. Patients were not invited to comment on the study design and were not consulted to develop patient-relevant outcomes or interpret the results. Patients were not invited to contribute to the writing or editing of this document for readability or accuracy.

### Study Population

We prospectively studied 35 consecutive patients with adult MMD (age of MMD onset >18 years) admitted to the stroke unit at our hospital (serving a population of more than 10,000,000 in Guangzhou, China). All patients were diagnosed with MMD by digital subtraction angiography (DSA) from January 2014 to August 2018. According to the modified Suzuki scoring (mSS), there 3 MMD patients were classified as mSSI, 17 as mSSII, 13 as mSSIII and 2 as mSSIV.

Inclusion criteria included patients with angiographically identified unilateral or bilateral severe stenosis or occlusion in the distal internal carotid artery or proximal middle and anterior cerebral arteries with eminent lenticulostriate arteries known as 'moyamoya collaterals'.

Exclusion criteria were patients with systemic vasculitis, neurofibromatosis, meningitis, sickle-cell disease, Down syndrome, prior radiotherapy on the skull base, or any other diseases that might be responsible for the observed vasculopathy. Consecutive patients with stroke subtype-matched non-MMD stroke admitted during the same period were selected as controls.

Baseline information on demographics and medical history were obtained predominantly by face-to-face interviews and medical records. Detailed data forms on clinical features were completed on admission by stroke physicians. Written informed consent for study participation was obtained from patients or an appropriate family member of the patient (in the case that the patient was disabled). The study was approved by the Ethics Committees of the second affiliated hospital of Guangzhou Medical University (Guangzhou).

In addition to bypass surgery in some MMD patients, other treatments were similar between the two groups. Cerebral perfusion pressure was guaranteed while avoiding hypovolaemia. Regulation of blood pressure, blood sugar, lipid and dehydration was performed as necessary. Acute management was mainly supportive, with a focus on reducing intracerebral pressure and managing seizures. Medical treatments included vasodilators, antiplatelet agents, antifibrotic agents and fibrinolytic agents. Epileptic patients were treated with anti-convulsants.

### DSA and Biochemical Tests

All examinations were performed using a Politron 1000 VR unit (Siemens, Nürnberg, Germany). After selective common carotid artery catheterisation through right transfemoral artery puncture, iohexol (Omnigraf; Juste, Madrid, Spain) was injected at a rate of 10 mL/s, and three different projections (posteroanterior, lateral and 45° oblique views) were taken. The diagnosis of MMD was confirmed according to previously described criteria. Peripheral venous blood samples were collected in the morning. A Beckman CX5 Automated Analyzer was used to determine the levels of fasting blood glucose, total cholesterol, triglycerides, HDL, LDL, ApoA1, ApoB, prothrombin time and high-sensitivity C-reactive protein. Plasma homocysteine level was determined by fluorescence ratio biochemical assay kit.

### Statistical Analysis

To determine the statistical differences in baseline characteristics, continuous variables were compared using *t* tests or Wilcoxon tests, and categorical variables were compared using  $\chi^2$  tests. Analysis of variance was used to study the homocysteine levels in subgroups of patients with adult MMD divided by the mSS. Logistic regression analysis was used to determine the risk factors associated with adult MMD. Univariate analyses were conducted first, followed by multivariate logistic regression analyses after adjusting for age, sex, medical history (hypertension, coronary heart disease and diabetes) and present history of smoking and alcohol use. Results from the univariate analyses showed that the variables were significantly associated with MMD, indicating statistical significance ( $p < 0.05$ ). All analyses were conducted using PASW statistics 18.0 (SPSS Inc., Chicago, Ill, USA).

## ■ RESULTS

### Study Population

A total of 35 patients with adult MMD and 202 patients with non-MMD stroke (180 patients with ischaemic stroke and 22 patients with haemorrhagic stroke) were included in this study. All analyses herein were based on data from these 237 patients.

Among the 35 patients with adult MMD, the average age was  $48.2 \pm 13.5$  years, and the female-to-male ratio was 1.7. Eighteen patients (51.4%) had ischaemic stroke, and 17 (48.6%) had haemorrhagic stroke. The percentage of patients with unilateral MMD was 51.4% (18/35), and bypass surgery was performed 28.6% (10/35) of the patients, both of which

are lower than in Korea (84.4%, 37.7%) and the United States (60.7%, 100%;  $p < 0.05$ ).

As shown in Table I, stroke patients with adult MMD were younger than non-MMD stroke patients. The proportion of patients who were technological workers was higher in those with MMD stroke. Similarly, the mean serum homocysteine level was significantly higher in patients with adult MMD than in non-MMD stroke patients. In contrast, hypertension was a more common comorbidity in non-MMD patients. There was no difference between homocysteine levels in the subgroups of MMD patients divided by the mSS, as shown in Table II.

Multiple logistic regression analysis was performed to further evaluate the independent risk factors for MMD (versus non-MMD stroke). As shown in Table II, after adjusting for baseline demographics, risk factors for stroke, significant variables from univariate analyses, homocysteine level and being a technological worker were still shown to be significantly associated with MMD.

## DISCUSSION

Moyamoya disease is a rare cerebrovascular disorder. Worldwide, the incidence of MMD is higher in Japan and in other East Asian countries. However, the incidence of MMD is estimated to be less than 11 per 100,000, even in Japan. In patients from South China, the detection rate of MMD is 3.2% in patients undergoing cerebral angiography, yet we do not know the exact incidence of adult MMD. In terms of age and sex distributions, the Chinese patient population has similarities to patient populations from other East Asian countries [such as Japan (20), and Korea (25)] as well as Western countries [such as the United States (30)]. Our experiences indicated several clinical differences between Chinese patients and patients from the above-mentioned countries. Among East Asian populations, ischaemic stroke is the primary clinical symptom. Interestingly, bilateral MMD was observed more frequently in patients with adult MMD in Japan (82.5%), Korea (84.4%) and the United States (60.7%), whereas a higher frequency of unilateral MMD was observed

**Table I:** Baseline Demographics and Clinical Characteristics in Patients with Adult MMD and non-MMD Stroke\*

Variable	MMD stroke (n=35)	Non-MMD stroke (n=202)	p
Age (years)	48.2 ± 13.5	55.6 ± 12.3	<b>0.003</b>
Male	22 (62.9)	140 (69.3)	0.45
Technological worker	18 (51.4)	31 (15.3)	<b>&lt;0.001</b>
Current smoker	10 (28.6)	65 (32.2)	0.67
Current alcohol user	6 (17.1)	37 (18.3)	0.87
With family history	4 (11.4)	18 (8.9)	0.64
<b>Medical history</b>			
Hypertension	11 (31.4)	127 (62.9)	<b>&lt;0.001</b>
Coronary heart disease	1 (2.9)	23 (11.4)	0.123
Diabetes mellitus	5 (14.3)	26 (12.9)	0.82
<b>Pathology results</b>			
Fasting glucose (mmol/L)	5.6 (4.8, 6.5)	5.4 (4.6, 7.0)	0.31
Triglyceride (mmol/L)	1.7 (1.0, 2.7)	1.7 (1.2, 2.3)	0.81
Total cholesterol (mmol/L)	4.8 (4.4, 5.8)	4.9 (4.2, 5.7)	0.78
HDL (mmol/L)	1.0 (0.9, 1.2)	1.0 (0.8, 1.2)	0.83
LDL (mmol/L)	3.0 (2.7, 3.8)	3.0 (2.5, 3.7)	0.34
ApoA1 (g/L)	1.2 (1.0, 1.3)	1.2 (1.1, 1.4)	0.24
ApoB (g/L)	0.9 (0.8, 1.1)	0.9 (0.8, 1.0)	0.58
Homocysteine (g/L)	21.0 (16.2, 33.0)	11.6 (9.5, 14.7)	<b>&lt;0.001</b>
HsCRP (mg/L)	3.8 (0.9, 11.3)	3.2 (1.1, 10.8)	0.75
Prothrombin time (seconds)	14.3 (13.0, 15.4)	14.3 (13.1, 15.2)	0.78

\*Values are reported as mean ± SD, number of subjects, or median (IQR); **Abbreviations:** **HDL:** high-density lipoprotein, **LDL:** low-density lipoprotein, **ApoA1:** apolipoprotein-A1, **ApoB:** apolipoprotein-B, **HsCRP:** high sensitivity C-reactive protein.

**Table II:** Logistic Regression Analysis of Independent Risk Factors for Adult MMD

Variable	Crude, n=237		Adjusted, n=237 <sup>†</sup>	
	OR (95% CI)	p	OR (95% CI)	p
<b>Sociodemographic</b>				
Age (years)	0.96 (0.93, 0.98)	<b>0.002</b>	-	-
Male	1.33 (0.63, 2.82)	0.45	-	-
Technological worker	5.84 (2.72, 12.56)	<b>&lt;0.001</b>	4.233 (1.65, 10.89)	<b>0.003</b>
Current smoker	0.84 (0.38, 1.86)	0.67	-	-
Current alcohol user	0.92 (0.37, 2.38)	0.87	-	-
With family history	1.32 (0.42, 4.16)	0.64	-	-
<b>Medical history</b>				
Hypertension	0.27 (0.13, 0.58)	<b>0.001</b>	0.26 (0.10, 0.71)	<b>0.009</b>
Coronary heart disease	0.23 (0.03, 1.75)	0.16	-	-
Diabetes mellitus	1.13 (0.40, 3.17)	0.82	-	-
<b>Pathology results</b>				
Fasting glucose (mmol/L)	1.04 (0.93, 1.17)	0.50	-	-
Triglyceride (mmol/L)	1.11 (0.97, 1.27)	0.14	-	-
Total cholesterol (mmol/L)	1.04 (0.77, 1.40)	0.79	-	-
HDL (mmol/L)	1.56 (0.40, 6.14)	0.53	-	-
LDL (mmol/L)	0.97 (0.73, 1.28)	0.81	-	-
ApoA1 (g/L)	0.25 (0.05, 1.24)	0.09	-	-
ApoB (g/L)	1.57 (0.31, 7.98)	0.59	-	-
Homocysteine (g/L)	1.08 (1.04, 1.11)	<b>&lt;0.001</b>	1.100 (1.06, 1.14)	<b>&lt;0.001</b>
HsCRP (mg/L)	0.99 (0.97, 1.01)	0.36	-	-
Prothrombin time (seconds)	1.01 (0.84, 1.22)	0.91	-	-

<sup>†</sup>Adjusted for Sociodemographic, medical history, and significant ( $p < 0.05$ ) variables from the univariate analyses: homocysteine.

in Chinese patients. In addition, we found that the frequency of bypass surgery was significantly lower in China in comparison with other East Asian countries (such as Japan [63.5%] and Korea [(37.7%)] and Western countries (such as the United States [100.0%]). Because bypass surgery is a proven effective therapy for MMD (19,32), this finding suggests room for improvement in the clinical management of MMD patients in China.

We found that hyperhomocysteinemia was an independent risk factor for MMD. Serum homocysteine level was significantly higher in patients with adult MMD than that in patients with non-MMD stroke, even after adjusting for extraneous variables. This finding is consistent with previous findings from case reports of MMD (bilateral or unilateral), in which homocysteine level was also noted to be a risk factor for MMD (5,23). Many retrospective and prospective studies have confirmed that hyperhomocysteinemia is a potential independent

risk factor for atherosclerosis (22), and an increased homocysteine level has been reported to be an independent risk factor for cerebral atherosclerosis (2). Hyperhomocysteinemia is a risk factor for middle cerebral artery stenosis (16), and is recognised to be associated with the presence and severity of systemic atherosclerosis (6). The potential mechanisms of action of homocysteine have been summarised as follows (1,26): damage to endothelial cells; reduction in NO generation; stimulation of smooth muscle cell proliferation; increasing foam cell formation; promotion of platelet aggregation, which in turn promotes thrombus formation; and reduction in the flexibility of vessels. Homocysteine can also enhance the negative impacts from risk factors such as smoking and abnormal lipid metabolism, resulting in the development of inflammation (4). These results suggest that homocysteine may have the same effects on adult MMD as it has on atherosclerosis. As a result, atherosclerosis secondary to increased homocysteine was the likely cause of moyamoya (5). Our hypothesis is sup-

ported by the presence of peripheral atherosclerotic disease, ischaemic heart disease, the relevant risk factors for atherosclerosis and the absence of other predisposing conditions of MMD in the patient samples in our study. On the other hand, one case report of homocystinuria with moyamoya was also described (9), and the possible role of the enzyme activity of methylenetetrahydrofolate reductase (MTHFR), which is associated with hyperhomocysteinemia, in MMD was discussed in the literature (24). A recent study indicated an association between two novel single-nucleotide polymorphisms in the gene regulating homocysteine metabolism (rs9651118 in MTHFR and rs117353193 in TCN2), resulting in increased homocysteine levels in patients with MMD (7,8). We believe that the genetic metabolic factor for homocysteinemia in the release of nitric oxide is the proposed mechanism (21), resulting in the development of moyamoya vessels and stroke (23).

Interestingly, we also found that the patient's occupation (ie, technological worker) plays a significant role in the development adult MMD. Previous studies have shown that radiation is one cause of MMD. Results from our study may implicate the role of radiation in adult MMD. However, we excluded patients with a history of radiotherapy on the skull base. In addition, we did not divide technological workers into subgroups (eg, exposure vs. nonexposure to radiation) because of insufficient data. Therefore, it is difficult to confirm the exact effects of radiation on adult MMD in this study.

There are several limitations of our study that are worth attention and further discussion. First, this study was a single-city hospital-based study in China, and as such, concerns about its external validity in other regions of China are raised. Moreover, only MMD patients with stroke were assessed, and the risk factors for MMD patients who have not yet experienced end-organ diseases may be different from those reported herein. Second, several important potential risk factors for MMD, such as concentrations of certain growth factors or cytokines, were not studied because they were out of scope for this study. Nonetheless, the definite similarity of results from this study to those from previous case reports reassures us of the validity of our data.

## CONCLUSION

In conclusion, results from our study indicate an independent association between elevated homocysteine levels and MMD, and also suggest that environmental factors such as type of occupation may play an important role in the development of MMD in China. Furthermore, given the practical blank of application and proven efficacy, bypass surgery should be more widely promoted in clinical practice.

## ACKNOWLEDGEMENTS

This study was supported by grants from Guangzhou Municipal Science & Technology Commission (No. 201704020043) and Guangzhou health science and technology general guidance project(No.20201A011088).

**Clinical Trial Registration:** <http://www.chictr.org.cn/hvshow-project.aspx?id=20390> Unique identifier: ChiCTR1900027368

## REFERENCES

1. An SA, Lee HB, Kim Y, Kim J, Kim HS, Kim WC, Kim OJ, Oh SH: Plasma total homocysteine level is associated with the pulsatility index of cerebral arteries in lacunar infarction. *Yonsei Med J* 54:819-824, 2013
2. Banecka-Majkutewicz Z, Sawula W, Kadzinski L, Węgrzyn A, Banecki B: Homocysteine, heat shock proteins, genistein and vitamins in ischemic stroke-pathogenic and therapeutic implications. *Acta Biochim Pol* 59:495-499, 2012
3. Bao XY, Wang QN, Zhang Y, Zhang Q, Li DS, Yang WZ, Zhang ZS, Zong R, Han C, Duan L: Epidemiology of moyamoya disease in China: Single-center, population-based study. *World Neurosurg* 122:e917-e923, 2019
4. Baszczuk A, Kopczyński Z: Hyperhomocysteinemia in patients with cardiovascular disease. *Postepy Hig Med Dosw (Online)* 68:579-589, 2014
5. Cerrato P, Grasso M, Lentini A, Destefanis E, Bosco G, Caprioli M, Bradac GB, Bergui M: Atherosclerotic adult moyamoya disease in a patient with hyperhomocysteinemia. *Neurol Sci* 28:45-47, 2007
6. Cioni G, Marcucci R, Gori AM, Valente S, Giglioli C, Gensini GF, Abbate R, Boddì M: Increased homocysteine and lipoprotein(a) levels highlight systemic atherosclerotic burden in patients with a history of acute coronary syndromes. *J Vasc Surg* 64:163-170, 2016
7. Duan L, Bao XY, Yang WZ, Shi WC, Li DS, Zhang ZS, Zong R, Han C, Zhao F, Feng J: Moyamoya disease in China: Its clinical features and outcomes. *Stroke* 43:56-60, 2012
8. Duan L, Wei L, Tian Y, Zhang Z, Hu P, Wei Q, Liu S, Zhang J, Wang Y, Li D, Yang W, Zong R, Xian P, Han C, Bao X, Zhao F, Feng J, Liu W, Cao W, Zhou G, Zhu C, Yu F, Yang W, Meng Y, Wang J, Chen X, Wang Y, Shen B, Zhao B, Wan J, Zhang F, Zhao G, Xu A, Zhang X, Liu J, Zuo X, Wang K: Novel susceptibility loci for Moyamoya Disease revealed by a genome-wide association study. *Stroke* 49:11-18, 2017
9. Erol M, Gayret OB, Yigit O, Cabuk KS, Toksoz M, Tiras M: A case of homocystinuria misdiagnosed as moyamoya disease: A case report. *Iran Red Crescent Med J* 18:e30332, 2016
10. Fujimura M, Bang OY, Kim JS: Moyamoya Disease. *Front Neurol Neurosci* 40:204-220, 2016
11. Fujimura M, Fujimura T, Kakizaki A, Sato-Maeda M, Niizuma K, Tomata Y, Aiba S, Tominaga T: Increased serum production of soluble CD163 and CXCL5 in patients with moyamoya disease: Involvement of intrinsic immune reaction in its pathogenesis. *Brain Res* 1679:39-44, 2018
12. Garg AK, Suri A, Sharma BS: Ten-year experience of 44 patients with moyamoya disease from a single institution. *J Clin Neurosci* 17:460-463, 2010
13. Guey S, Kraemer M, Hervé D, Ludwig T, Kossorotoff M, Bergametti F, Schwitalla JC, Choi S, Broseus L, Callebaut I, Genin E, Tournier-Lasserre E, FREX Consortium: Rare RNF213 variants in the C-terminal region encompassing the RING-finger domain are associated with moyamoya angiopathy in Caucasians. *Eur J Hum Gene* 25:995-1003, 2017
14. Hallemeier CL, Rich KM, Grubb RL Jr, Chicoine MR, Moran CJ, Cross DT 3rd, Zipfel GJ, Dacey RG Jr, Derdeyn CP: Clinical features and outcome in north american adults with moyamoya phenomenon. *Stroke* 37:1490-1496, 2006

15. Hishikawa T, Sugiu K, Date I: Moyamoya disease: A review of clinical research. *Acta Med Okayama* 70:229-236, 2016
16. Huang HW, Guo MH, Lin RJ, Chen YL, Luo Q, Zhang Y, Wong KS: Hyperhomocysteinemia is a risk factor of middle cerebral artery stenosis. *J Neurol* 254:364-367, 2007
17. Kim EH, Yum MS, Ra YS, Park JB, Ahn JS, Kim GH, Goo HW, Ko TS, Yoo HW: Importance of RNF213 polymorphism on clinical features and long-term outcome in moyamoya disease. *J Neurosurg* 124:1221-1227, 2016
18. Kraemer M, Heienbrok W, Berlit P: Moyamoya disease in Europeans. *Stroke* 39:3193-3200, 2008
19. Kraemer M, Karakaya R, Matsushige T, Graf J, Albrecht P, Hartung HP, Berlit P, Laumer R, Diesner F: Efficacy of STA-MCA bypass surgery in moyamoya angiopathy: Long-term follow-up of the Caucasian Krupp Hospital cohort with 81 procedures. *J Neurol* 265:2425-2433, 2018
20. Kuroda S, Ishikawa T, Houkin K, Nanba R, Hokari M, Iwasaki Y: Incidence and clinical features of disease progression in adult moyamoya disease. *Stroke* 36:2148-2153, 2005
21. McCully KS: Homocysteine and vascular disease. *Nat Med* 2:386-389, 1996
22. McCully KS: Hyperhomocysteinemia and arteriosclerosis: Historical perspectives. *Clin Chem Lab Med* 43:980-986, 2005
23. Meena DS, Bohra GK, Meena M, Maheshwari BK: Hyperhomocysteinemia in a patient with moyamoya disease. *Case Rep Neurol Med* 2018:7806873, 2018
24. Park YS, Jeon YJ, Kim HS, Han IB, Choi JU, Kim DS, Kim NK: The roles of methylenetetrahydrofolate reductase 677C>T and 1298A>C polymorphisms in moyamoya disease patients. *Childs Nerv Syst* 30:1687-1695, 2014
25. Rhim JK, Cho YD, Jeon JP, Yoo DH, Cho WS, Kang HS, Kim JE, Han MH: Ruptured aneurysms of collateral vessels in adult onset moyamoya disease with hemorrhagic presentation. *Clin Neuroradiol* 28:191-199, 2018
26. Rueda-Clausen C F, Cordoba-Porras A, Bedoya G, Silva FA, Zarruk JG, López-Jaramillo P, Villa LA: Increased plasma levels of total homocysteine but not asymmetric dimethylarginine in Hispanic subjects with ischemic stroke FREC-VI sub-study. *Eur J Neurol* 19:417-425, 2012
27. Shen W, Liao Y, Garcia R, Kesavabhotla K, Xu B, Li H: Association of CD40 SNPs with moyamoya in a Chinese children population. *Br J Neurosurg* 33:398-401, 2019
28. Suzuki J, Takaku A: Cerebrovascular 'moyamoya' disease. Disease showing abnormal net-like vessels in base of brain. *Arch Neurol* 20:288-299, 1969
29. Takahashi Y, Mikami T, Suzuki H, Komatsu K, Yamamoto D, Shimohama S, Houkin K, Sugita S, Hasegawa T, Mikuni N: Development of moyamoya disease after non-herpetic acute limbic encephalitis: A case report. *J Clin Neurosci* 53:250-253, 2018
30. Zeifert PD, Karzmark P, Bell-Stephens TE, Steinberg GK, Dorfman LJ: Neurocognitive performance after cerebral revascularization in adult moyamoya disease. *Stroke* 48:1514-1517, 2017
31. Zhang H, Zheng L, Feng L: Epidemiology, diagnosis and treatment of moyamoya disease. *Exp Ther Med* 17:1977-1984, 2019
32. Zheng J, Yu LB, Dai KF, Zhang Y, Wang R, Zhang D: Clinical features, surgical treatment, and long-term outcome of a multicenter cohort of pediatric moyamoya. *Front Neurol* 22:10-14, 2019