

3-25-2020

Associations between Aquaglyceroporin Gene Polymorphisms and Risk of Stroke among Patients with Hypertension

Qingyun Tu
Southeast University, China

Li Yan
Southeast University, China

Chun Wang
Thomas Jefferson University

Aohan Han
Southeast University, China

Yu Qin
Follow this and additional works at: <https://jdc.jefferson.edu/medoncfp>
Jiangsu Provincial Center for Disease Control and Prevention

 Part of the [Oncology Commons](#)

[Let us know how access to this document benefits you](#)

See next page for additional authors

Recommended Citation

Tu, Qingyun; Yan, Li; Wang, Chun; Han, Aohan; Qin, Yu; Cui, Lan; and Xiang, Quanyong, "Associations between Aquaglyceroporin Gene Polymorphisms and Risk of Stroke among Patients with Hypertension" (2020). *Department of Medical Oncology Faculty Papers*. Paper 117.

<https://jdc.jefferson.edu/medoncfp/117>

This Article is brought to you for free and open access by the Jefferson Digital Commons. The Jefferson Digital Commons is a service of Thomas Jefferson University's [Center for Teaching and Learning \(CTL\)](#). The Commons is a showcase for Jefferson books and journals, peer-reviewed scholarly publications, unique historical collections from the University archives, and teaching tools. The Jefferson Digital Commons allows researchers and interested readers anywhere in the world to learn about and keep up to date with Jefferson scholarship. This article has been accepted for inclusion in Department of Medical Oncology Faculty Papers by an authorized administrator of the Jefferson Digital Commons. For more information, please contact: JeffersonDigitalCommons@jefferson.edu.

Authors

Qingyun Tu, Li Yan, Chun Wang, Aohan Han, Yu Qin, Lan Cui, and Quanyong Xiang

Research Article

Associations between Aquaglyceroporin Gene Polymorphisms and Risk of Stroke among Patients with Hypertension

Qingyun Tu,¹ Li Yan,² Chun Wang,³ Aohan Han,¹ Yu Qin,⁴ Lan Cui,⁴
and Quanyong Xiang^{ID}⁴

¹School of Public Health, Southeast University, Nanjing 210009, China

²Department of Infection Prevention and Control, Zhongda Hospital of Southeast University, Nanjing 210009, China

³Department of Medical Oncology, Thomas Jefferson University, Philadelphia, PA 19107, USA

⁴Department of Noncommunicable Chronic Disease Control, Jiangsu Provincial Center for Disease Control and Prevention, Nanjing 210009, China

Correspondence should be addressed to Quanyong Xiang; quanyongxiang@vip.sina.com

Received 14 May 2019; Revised 6 February 2020; Accepted 19 March 2020; Published 26 March 2020

Academic Editor: Gang Liu

Copyright © 2020 Qingyun Tu et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Background. Dysregulations of AQP7 and AQP9 were found to be related to lipid metabolism abnormality, which had been proven to be one of the mechanisms of stroke. However, limited epidemiological studies explore the associations between AQP7 and AQP9 and the risk of stroke among patients with hypertension in China. **Aims.** We aimed to investigate the associations between genetic variants in AQP7 and AQP9 and the risk of stroke among patients with hypertension, as well as to explore gene-gene and gene-environment interactions. **Methods.** Baseline blood samples were drawn from 211 cases with stroke and 633 matched controls. Genomic DNA was extracted by a commercially available kit. Genotyping of 5 single nucleotide polymorphisms (SNPs) in AQP7 (rs2989924, rs3758269, and rs2542743) and AQP9 (rs57139208, rs16939881) was performed by the polymerase chain reaction assay with TaqMan probes. **Results.** Participants with the rs2989924 GG genotype were found to be with a 1.74-fold increased risk of stroke compared to those with the AA+AG genotype, and this association remained significant after adjustment for potential confounders (odds ratio (OR): 1.74, 95% confidence interval (CI): 1.23-2.46). The SNP rs3758269 CC+TT genotype was found to be with a 33% decreased risk of stroke after multivariate adjustment (OR: 0.67, 95% CI: 0.45-0.99) compared to the rs3758269 CC genotype. The significantly increased risk of stroke was prominent among males, patients aged 60 or above, and participants who were overweight and with a harbored genetic variant in SNP rs2989924. After adjusting potential confounders, the SNP rs3758269 CT+TT genotype was found to be significantly associated with a decreased risk of stroke compared to the CC genotype among participants younger than 60 years old or overweight. No statistically significant associations were observed between genotypes of rs2542743, rs57139208, or rs16939881 with the risk of stroke. Neither interactions nor linkage disequilibrium had been observed in this study. **Conclusions.** This study suggests that SNPs rs2989924 and rs3758269 are associated with the risk of stroke among patients with hypertension, while there were no statistically significant associations between rs2542743, rs57139208, and rs16939881 and the risk of stroke being observed.

1. Introduction

Stroke is the second most common cause of death worldwide and is the leading cause of long-term disability in developing and developed countries [1]. In China, there are more than one million people who die from stroke-related diseases per year. Evidence suggests that the pathogenesis of stroke is the result of interactions between genetic predispositions

and environmental factors [2]. A large body of scientific research has indicated that gene polymorphisms modulate the pathophysiological processes of stroke and confer a small to moderate risk [3–5]. Studies in twins, families, and animal models provide substantial evidence for a genetic contribution to stroke [6, 7]. Recently, polymorphisms of genes involved in lipid metabolism have been found to be associated with stroke occurrence [8].

Aquaporins (AQPs) represent a family of transmembrane proteins permeable to water and, in some cases, also to other solutes [9]. Generally, the AQP family can be divided into three major subgroups based on their permeability characteristics and amino acid sequence homology [10]: AQP0, AQP1, AQP2, AQP4, and AQP5 belong to the group of classical aquaporins; AQP6, AQP8, AQP11, and AQP12 belong to the second subgroup called unorthodox aquaporins [11]; and AQP3, AQP7, AQP9, and AQP10 belong to the third group, called aquaglyceroproteins, which are permeable to small uncharged molecules, such as urea, glycerol, or water [12–16].

In recent years, researchers pay more attention to the associations between AQPs' genetic polymorphisms and disease susceptibility. AQPs have been implicated in the regulation of both physiological and pathological water homeostasis and thus represent a promising target for alleviating stroke-induced cerebral edema [17]. Specifically, AQP7—the adipose-specific glycerol channel—represents a gateway for the delivery of adipose-derived glycerol into plasma [15, 18, 19] and it was identified as one of the dysregulated adipose tissue genes in obese humans [20]. AQP7 is also known to play a pivotal role in glycerol metabolism in a wide range of tissues with implications for whole-body energy balance as well as the pathophysiology of obesity and development of insulin resistance [10]. Studies suggest that AQP7 dysregulation may lead to an increased supply of glycerol for hepatic gluconeogenesis and to increased glucose levels in type 2 diabetes [11]. Moreover, AQP9 is considered the main facilitator of glycerol uptake in hepatocytes [11, 21, 22]. Ribeiro et al. found that after focal transient ischemia in mice, a profound increase in AQP9 immunolabeling was detected in astrocytes in peri-infarcted areas after 24 h of postocclusion, with a maximum at 7 days postocclusion [23]. Dysregulation of AQP7 and AQP9 may result in lipid metabolism abnormality, which is one of the important mechanisms of stroke.

However, epidemiological studies on the associations between AQP7 and AQP9 and the risk of stroke among patients with hypertension are still largely lacking. Therefore, this study is aimed at investigating the associations between AQP7 and AQP9 genetic variants and the risk of stroke among patients with hypertension.

2. Materials and Methods

2.1. Ethical Approval. This study was approved by the Institutional Review Board of Jiangsu Provincial Center for Disease Control and Prevention, and each participant signed a written informed consent for their participation.

2.2. Study Population. Participants in this study were recruited from the Follow-up Cohort Study of Hypertension (FCSH), which was established in 2010 in Jiangsu Province of China. In FCSH, over 12,000 patients with hypertension from 5 different counties (Jianye district in Nanjing, Peixian county in Xizhou, Haiyan county in Nantong, Lianshui county in Huaian, and Sihong county in Suqian) of Jiangsu Province were involved. Eligible criteria for inclusion were as follows: adults (18–70 years), the essential hypertension (with the

systolic blood pressure (SBP) ≥ 140 mmHg and/or diastolic blood pressure (DBP) ≥ 90 mmHg), and blood pressure does not reach the control target (i.e., SBP ≥ 140 or DBP ≥ 90 mmHg). Patients were excluded if they met one of the following conditions: with secondary hypertension, having comorbidities (peripheral vascular diseases, coronary artery diseases, autoimmune diseases, systemic inflammatory diseases, blood diseases, malignant tumors, acute myocardial infarction, and stroke) in the previous 3 months of the study, or with life expectancy of less than one year. All participants were followed up once a year. At each follow-up visit, blood pressure and body weight were measured and information about lifestyle (e.g., smoking, drinking, and physical activity) and diseases (e.g., cardiocerebrovascular disease, cancers, and diabetes) was updated.

By the end of 2014, there were 301 person-times cases with stroke with confirmed diagnosis. The cumulative incidence of stroke was 2.51%, and the average follow-up time was 43.4 (± 10.2) months. All patients with stroke were identified by at least two neurology specialists according to clinical signs and symptoms and/or imaging data. Among the 301 cases with stroke, 90 were excluded because of recurrent stroke ($N = 38$), no gene polymorphism test ($N = 19$), no qualified blood sample ($N = 9$), and having other severe diseases (such as cancer, severe emphysema, and severe liver and kidney function) ($N = 24$). We finally included 211 cases with stroke in this study.

The control group was composed of followed-up hypertensive patients who did not have stroke, deep vein thrombosis, or myocardial infarction. Cases and controls were matched on age (± 3 years old), gender, baseline blood pressure (± 5 mmHg), and residential area according to the ratio of 1 : 3. At last, 633 controls were included in this study.

2.3. Data Collection. All participants underwent a face-to-face interview by trained interviewers to complete a structured questionnaire, including demographic information, lifestyle information, medical history, and family history. After the interview, anthropometric indices (height, body weight, and waist circumference (WC)) and blood pressure were measured. Body weight and height were measured according to a standard protocol [24]. Body mass index (BMI) was calculated as weight in kilograms divided by height in meters squared (kg/m^2). Blood pressure was measured using an automated device (OMRON Model HEM-7071, Omron Co.) at the nondominant arm 3 times consecutively with a 1–2 minute interval between two measurements while participants were in a seated position.

Blood samples were collected from participants after overnight fasting for at least 10 h. Serum total cholesterol (TC), low-density lipoprotein cholesterol (LDL-C), high-density lipoprotein cholesterol (HDL-C), triglycerides (TG), and fasting blood glucose (FBG) were measured using an autoanalyzer (Abbott Laboratories).

An individual was categorized as “overweight” if their BMI $\geq 25 \text{ kg}/\text{m}^2$, and central obesity was defined according to the waist circumference (WC) of the participant as men > 90 cm or women > 80 cm [25]. A stringent quality

assurance and quality control procedure was implemented to ensure the validity and reliability of study data.

2.4. SNP Selection and Genotyping Assay. We followed the methods of Wang et al. to select five single nucleotide polymorphisms (SNPs) (rs2989924, rs3758269, and rs62542743 in *AQP7* and rs57139208 and rs16939881 in *AQP9*) and assay genotyping [26]. Allelic discrimination was automatically completed using the Sequence Detection Systems Software version 2.3. The genotyping call rates for 5 SNPs were all >97% in this study. For quality control, the DNA samples from cases and controls and two blank controls were run in the same batch, and the operators performing detection and genotyping did not know the case or control status of the samples. Moreover, 10% of genotyping samples were randomly chosen to redo the assay blindly for consistency check.

2.5. Statistical Analysis. All the data in this study were double entered by a trained personnel using EpiData 3.5 and analyzed by SPSS 21.0. The differences in demographic characteristics and selected variables between cases and controls were compared using a Student *t*-test for continuous variables and a χ^2 test for categorical variables. The Hardy-Weinberg equilibrium was tested by the goodness-of-fit χ^2 test to compare the observed genotype frequencies to the expected ones among controls. The associations between genotypes and risk of stroke were evaluated using univariate and multivariate logistic regression analyses to determine the crude and adjusted odds ratios (OR) and their 95% confidence intervals (CI). All statistical tests were two sided. $P < 0.05$ was considered statistically significant. The genetic linkage balance test of 5 SNPs was performed using SNPstats software. The generalized multifactorial dimensionality reduction method (GMDR) was used to analyze the gene-gene interaction. Using the incidence of stroke at follow-up as the outcome variable, the five loci of genes rs16939881, rs57139208, rs62542743, rs3758269, and rs2989924 were named as SNP1, SNP2, SNP3, SNP4, and SNP5, respectively, in turn and were included in the GMDR model as the analysis factors. Meanwhile, the age, gender, duration of hypertension, family history of hypertension, family history of stroke, SBP, TC, TG, and FBG were included in the model as potential confounders. The haplotypes of SNPs on the same chromosome were analyzed by SNPstats software.

3. Results

3.1. Demographic Characteristics of Participants. No deviation from the Hardy-Weinberg equilibrium for the polymorphisms examined was observed in the genotype distributions between cases and controls. The demographic characteristics of participants are presented in Table 1. A total number of 211 patients with stroke and 633 controls were included in the study. Significant differences were observed between cases and controls in the duration of hypertension, family history of hypertension, or stroke. SBP, TC, TG, and FBG were higher among patients with stroke than those in their matched control group.

3.2. Associations between SNPs and the Risk of Stroke. The frequencies of genotype and allele of the SNP polymorphisms are shown in Table 2. The genotype distributions of AA, GG, and AG in rs2989924 of *AQP7* were significantly different between cases and controls ($\chi^2 = 8.260$, $P = 0.016$). Participants with rs2989924 GG genotypes had a 1.55-fold increased risk of stroke (OR 1.55, 95% CI 1.12-2.15) compared to those with the AA+AG genotype, and this relationship remained significant after adjustment for covariates (age, gender, SBP, FBG, TC, TG, and LDL-C) (OR 1.74, 95% CI 1.23-2.46). When compared with the rs3758269 CC genotype, participants with CT+TT genotypes had a 33% decreased risk of stroke after adjustment for potential confounders (OR 0.67, 95% CI 0.45-0.99). No association was observed between the genotypes of rs62542743, rs57139208, and rs16939881 and the risk of stroke.

3.3. Stratified Analyses of the Associations between Genotypes of rs2989924 and rs3758269 and Risk of Stroke. Table 3 shows the results of stratified analyses. A significantly increased risk of stroke in the rs2989924 GG vs. AA+AG (recessive model) genotype was observed among men (OR 1.80, 95% CI 1.10-2.93) and the elderly (OR 1.81, 95% CI 1.20-2.75), and this association remained significant after adjustment for potential confounders (OR 2.23, 95% CI 1.32-3.77 and OR 2.20, 95% CI 1.41-3.44, respectively). Among the hypertensive population, the rs2989924 GG vs. AA+AG (recessive model) genotype was found to be associated with an increased risk of stroke among participants who were overweight or central obese, with an unadjusted OR of 2.35 (95% CI 1.45-3.81) and 1.81 (95% CI 1.22-2.70), respectively. However, among patients with ≥ 25 kg/m² BMI, the rs3758269 CT genotype was associated with a reduced risk of stroke compared to those harboring the CC genotype (adjusted OR 0.40, 95% CI 0.21-0.75), and a similar result was observed in the CT+TT vs. CC genotype (dominant mode) (adjusted OR 0.43, 95% CI 0.24-0.77). The rs3758269 CT+TT genotype was associated with a reduced risk of stroke compared with the CC genotype among participants younger than 60 years old (adjusted OR 0.48, 95% CI 0.25-0.94).

3.4. Analysis of Interaction. Table 4 shows the gene-gene interactions, and the results suggested that there was no interaction.

3.5. Haplotype Analysis. The linkage disequilibrium (LD) test was conducted on rs62542743, rs3758269, rs2989924, rs57139208, and rs16939881. Results showed that there was a significant linkage imbalance between rs3758269 and rs2989924, and the D' value is 0.9992. $D' < 0.75$ between the other two SNPs indicated that there was no significant linkage imbalance. If only the SNPs with a frequency of more than 5% were included, three SNPs of *AQP7* can build 8 kinds of haplotype and two SNPs of *AQP9* can build 4 kinds of haplotype. Table 5 shows the results after adjustment for age, gender, duration of hypertension, family history of hypertension, family history of stroke, SBP, TC, TG, and FBG. Individuals with the CTA haplotype in *AQP7* have a higher risk of stroke compared to those with

TABLE 1: Demographic and clinical characteristics by cases with stroke and controls.

Variables	Cases with stroke N (%)	Controls N (%)	P
Degree of education			
Primary school and below	157 (74.4%)	436 (68.9%)	0.291
Junior high school	47 (22.3%)	167 (26.4%)	
High school and above	7 (3.3%)	30 (4.7%)	
Duration of hypertension (years)			
≤5	122 (60.7%)	416 (69.6%)	0.020
>5	79 (39.3%)	182 (30.4%)	
Familial history of hypertension	48 (23.1%)	102 (16.3%)	0.029
Familial history of stroke	10 (4.8%)	11 (1.7%)	0.014
Familial history of CHD	7 (3.4%)	11 (1.8%)	0.167
Familial history of diabetes	5 (2.4%)	11 (1.7%)	0.556
Central obesity	158 (74.9%)	480 (75.8%)	0.781
BMI (kg/m ²)			
Normal	78 (37.0%)	243 (38.4%)	0.922
Overweight	89 (42.2%)	264 (41.7%)	
Obese	44 (20.9%)	126 (19.9%)	
SBP (mmHg)	153.64 ± 12.88*	151.16 ± 10.60*	0.012
DBP (mmHg)	94.87 ± 8.16*	93.74 ± 6.92*	0.071
TC (mmol/L)	4.49 ± 0.76*	4.31 ± 0.79*	0.006
TG (mmol/L)	1.90 ± 1.26*	1.68 ± 1.02*	0.019
FBG (mmol/L)	5.45 ± 2.31*	5.07 ± 1.38*	0.023
HDL-C (mmol/L)	1.63 ± 0.41*	1.59 ± 0.39*	0.130
LDL-C (mmol/L)	2.04 ± 0.77*	2.01 ± 0.72*	0.652

* represents mean ± SD. SD: standard deviation; CHD: coronary heart disease; BMI: body mass index; SBP: systolic blood pressure; DBP: diastolic blood pressure; TC: total cholesterol; TG: triglycerides; FBG: fasting blood glucose; HDL-C: high-density lipoprotein cholesterol; LDL-C: low-density lipoprotein cholesterol.

the highest frequency of single-body CCG ($P < 0.05$). As shown in Table 6, the analysis of AQP9 by SNPstats found that there was no statistically significant difference in the risk of stroke between individuals with the highest frequency of haplotype CC and those with the remaining haplotypes ($P > 0.05$).

3.6. The Logistic Analysis of 5 SNPs and Risk of Stroke. The five loci of genes rs2989924, rs3758269, rs62542743, rs57139208, and rs16939881; duration of hypertension; family history of hypertension; family history of stroke; SBP; TC; TG; FBG; and daily intake of fruits were used as independent variables, and stroke was used as a dependent variable for logistic regression analysis. The results of multivariate analysis are shown in Table 7, which indicated that high TC was a risk factor for stroke (OR 1.31, 95% CI 1.05-1.63) and the rs2889924 AG genotype may be a protective factor for stroke.

4. Discussion

In this population-based case-control study, we investigated 5 SNPs in the AQP7 and AQP9 genes and found that the SNPs of rs2989924 and rs3758269 in AQP7 were associated with the risk of stroke, while no association was observed between AQP9 SNPs and the risk of stroke.

This study found a significantly increased risk of stroke in rs2989924 GG vs. AA+AG (recessive model) genotypes among men. According to some preclinical and basic studies, female hormones like estrogen and progesterone may have the ability to reduce the incidence of stroke, cardiovascular disease, and the tissue damage after an occurrence of ischemia [27, 28]. In this study, we found that people aged 60 or above with the GG vs. AA+AG genotype variation in SNP rs2989924 were with an increased risk of stroke, while the CT+TT genotype variation in rs3758269 was found to be related to a reduced risk of stroke among people under 60 years old. Such findings indicated that aging might cause an adverse reverse genetic variation of the AQP7 genotype and is likely to cause stroke and aggravate stroke symptoms [29]. Kim and Vemuganti's study had also reported that the tolerance of vasculature to blood pressure and the capacity of the brain's self-healing notably declined with increasing age, which contributed to increase the susceptibility of brain stroke [30].

Prudente et al. recruited 685 cases of type 2 diabetes mellitus (345 females/340 males) and 292 controls (185 females/107 males) in Caucasus and found that the risk of the AG+GG genotype was 1.36 times higher compared with the AA genotype (OR 1.36, 95% CI 1.01-1.84), while stratified analysis showed that the risk of the AG+GG

TABLE 2: Genotype frequency distribution in cases and controls.

Variable	Strokes, N (%)	Control, N (%)	χ^2	P	OR	95% CI	OR	95% CI*
<i>rs2989924</i>								
AA	41 (19.5)	117 (19.0)	8.260	0.016	1.00			
AG	86 (41.0)	317 (51.4)			0.774	0.505-1.188	0.645	0.409-1.018
GG	83 (39.5)	183 (29.7)			1.294	0.833-2.010	1.292	0.817-2.043
AA	41 (19.5)	117 (19.0)	0.032	0.858	1.00			
AG+GG	169 (80.5)	500 (81.0)			0.965	0.649-1.433	0.885	0.585-1.340
AA+AG	127 (60.5)	434 (70.3)	6.987	0.008	1.00			
GG	83 (39.5)	183 (29.7)			1.550	1.119-2.148	1.741	1.232-2.461
<i>rs3758269</i>								
CC	164 (77.7)	456 (72.2)	2.958	0.228	1.00			
CT	44 (20.9)	159 (25.2)			0.769	0.527-1.124	0.687	0.456-1.035
TT	3 (1.4)	17 (2.7)			0.491	0.142-1.696	0.500	0.143-1.754
CC	164 (77.7)	456 (72.2)	2.525	0.112	1.00			
CT+TT	47 (22.3)	176 (27.8)			0.743	0.514-1.073	0.669	0.450-0.994
CC+CT	208 (98.6)	615 (97.3)	1.098	0.295	1.00			
TT	3 (1.4)	17 (2.7)			0.522	0.151-1.798	0.545	0.156-1.904
<i>rs62542743</i>								
CC	192 (91.0)	584 (92.3)	0.361	0.835	1.00			
CA	18 (8.5)	46 (7.3)			1.190	0.674-2.102	1.054	0.573-1.939
AA	1 (0.5)	3 (0.5)			1.014	0.105-9.804	0.935	0.093-9.371
CC	192 (91.0)	584 (92.3)	0.341	0.559	1.00			
CA+AA	19 (9.0)	49 (7.7)			1.179	0.678-2.053	1.047	0.579-1.892
CC+CA	210 (99.5)	630 (99.5)	0.000	1.000	1.00			
AA	1 (0.5)	3 (0.5)			1.000	0.103-9.665	0.931	0.093-9.325
<i>rs57139208</i>								
CC	163 (77.3)	491 (77.6)	0.980	0.613	1.00			
CT	46 (21.8)	130 (20.5)			1.066	0.729-1.559	1.068	0.717-1.591
TT	2 (0.9)	12 (1.9)			0.502	0.111-2.267	0.284	0.036-2.254
CC	163 (77.3)	491 (77.6)	0.009	0.924	1.00			
CT+TT	48 (22.7)	142 (22.4)			1.018	0.702-1.477	1.005	0.679-1.488
CC+CT	209 (99.1)	621 (98.1)	0.872	0.351	1.00			
TT	2 (0.9)	12 (1.9)			0.495	0.110-2.231	0.280	0.035-2.218
<i>rs16939881</i>								
CC	184 (87.2)	551 (87.0)	0.442	0.802	1.00			
CG	26 (12.3)	76 (12.0)			1.024	0.637-1.648	1.038	0.632-1.704
GG	1 (0.5)	6 (0.9)			0.499	0.060-4.173	0.554	0.065-4.686
CC	184 (87.2)	551 (87.0)	0.004	0.953	1.00			
CG+GG	27 (12.8)	82 (13.0)			0.986	0.619-1.571	1.002	0.617-1.628
CC+CG	210 (99.5)	627 (99.1)	0.432	0.511	1.00			
GG	1 (0.5)	6 (0.9)			0.498	0.060-4.157	0.551	0.065-4.661

OR: odds ratio; CI: confidence interval. *Adjusted for age, gender, systolic blood pressure, fasting glucose, total cholesterol, triglycerides, and low-density lipoprotein cholesterol.

genotype was 1.66 times higher in females who were obese than that of the AA genotype (OR 1.66, 95% CI 1.01-2.74) [31]. In this study, the finding that the *AQP7* rs2989924 genotype variation was a risk factor for obesity-related diseases is consistent with findings of Prudente et al. However, the rs3758269 CT genotype in patients with BMI (kg/m²)

≥ 25 can reduce the risk of stroke compared with the CC genotype and the CT+TT vs. CC genotype (dominant model) after adjustment for potential confounders (OR 0.40, 95% CI 0.21-0.75 and OR 0.43, 95% CI 0.24-0.77, respectively). According to the stratification analysis of WC, no association between the rs3758269 genotype and stroke was found.

TABLE 3: Stratified analysis between the *AQP7* gene and the risk of stroke.

Variable	(Control/stroke)			OR (95% CI)			OR* (95% CI)			OR (95% CI)			OR* (95% CI)			OR (95% CI)			OR* (95% CI)			
	AA	AG	GG	AA	AG	GG	AA	AG	GG	AA+AG	AA+AG	GG	AA+AG	AA+AG	GG	AA+AG	GG	AA+AG	GG	AA+AG	GG	
rs2989924																						
Males	58/18	122/41	68/40	1.00	1.08 (0.57-2.05)	0.71 (0.36-1.43)	1.90 (0.98-3.66)	1.79 (0.90-3.56)	1.37 (0.76-2.48)	1.11 (0.60-2.07)	1.80 (1.10-2.93)	1.40 (0.90-2.17)	1.46 (0.91-2.32)	1.19 (0.66-2.13)	2.20 (1.41-3.44)	1.23 (0.73-2.07)	1.39 (0.68-2.85)	1.43 (0.74-2.75)	1.59 (1.09-2.32)	1.81 (1.22-2.70)	1.47 (0.37-5.78)	1.47 (0.37-5.90)
Females	59/23	195/45	115/43	1.00	0.59 (0.33-1.06)	0.57 (0.31-1.07)	0.96 (0.53-1.74)	0.99 (0.53-1.85)	0.73 (0.43-1.25)	0.73 (0.41-1.30)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.19 (0.66-2.13)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)
Age ≤ 60 (years)	52/19	131/40	69/27	1.00	0.84 (0.44-1.58)	0.61 (0.30-1.24)	1.07 (0.54-2.13)	0.86 (0.41-1.80)	0.92 (0.51-1.66)	0.71 (0.37-1.35)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.19 (0.66-2.13)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)
>60 (years)	65/22	186/46	114/56	1.00	0.73 (0.41-1.31)	0.64 (0.35-1.18)	1.45 (0.81-2.59)	1.62 (0.89-2.95)	1.01 (0.59-1.71)	0.10 (0.58-1.73)	1.81 (1.20-2.75)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)	1.21 (0.71-2.07)
BMI (kg/m ²) < 25	54/18	168/47	103/33	1.00	0.84 (0.45-1.57)	0.79 (0.40-1.56)	0.96 (0.50-1.86)	1.04 (0.51-2.11)	—	0.89 (0.47-1.67)	—	—	1.23 (0.73-2.07)	—	—	—	—	—	—	—	—	—
≥25	63/23	149/39	80/50	1.00	0.72 (0.40-1.30)	0.58 (0.31-1.11)	1.71 (0.95-3.10)	1.67 (0.89-3.11)	1.07 (0.62-1.82)	0.97 (0.55-1.71)	2.14 (1.36-3.36)	1.43 (0.74-2.75)	1.39 (0.68-2.85)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)
Without central obesity	33/10	73/23	45/20	1.00	1.04 (0.45-2.43)	0.84 (0.34-2.08)	1.47 (0.61-3.54)	1.23 (0.48-3.15)	1.20 (0.55-2.65)	0.99 (0.43-2.29)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)
With central obesity	84/31	244/63	138/63	1.00	0.70 (0.43-1.15)	0.62 (0.37-1.05)	1.24 (0.74-2.06)	1.30 (0.77-2.21)	0.89 (0.57-1.41)	0.87 (0.54-1.41)	1.59 (1.09-2.32)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)	1.43 (0.74-2.75)
rs3758269																						
Males	178/76	63/23	10/0	1.00	0.86 (0.49-1.48)	0.80 (0.44-1.46)	—	—	0.74 (0.43-1.27)	0.70 (0.39-1.26)	—	—	—	—	—	—	—	—	—	—	—	—
Females	278/88	96/21	7/3	1.00	0.69 (0.41-1.17)	0.59 (0.33-1.07)	1.35 (0.34-5.35)	1.32 (0.33-5.32)	0.74 (0.44-1.22)	0.65 (0.37-1.13)	1.47 (0.37-5.78)	1.47 (0.37-5.78)	1.47 (0.37-5.90)	1.47 (0.37-5.90)	1.47 (0.37-5.90)	1.47 (0.37-5.90)	1.47 (0.37-5.90)	1.47 (0.37-5.90)	1.47 (0.37-5.90)	1.47 (0.37-5.90)	1.47 (0.37-5.90)	1.47 (0.37-5.90)
Age ≤ 60 (years)	177/68	68/19	10/0	1.00	0.73 (0.41-1.30)	0.54 (0.27-1.07)	—	—	0.63 (0.36-1.13)	0.48 (0.25-0.94)	—	—	—	—	—	—	—	—	—	—	—	—
>60 (years)	279/96	91/25	7/3	1.00	0.80 (0.48-1.32)	0.78 (0.46-1.33)	1.25 (0.32-4.91)	1.25 (0.31-5.03)	0.83 (0.51-1.34)	0.82 (0.49-1.36)	1.31 (0.33-5.15)	1.31 (0.33-5.15)	1.31 (0.33-5.15)	1.31 (0.33-5.15)	1.31 (0.33-5.15)	1.31 (0.33-5.15)	1.31 (0.33-5.15)	1.31 (0.33-5.15)	1.31 (0.33-5.15)	1.31 (0.33-5.15)	1.31 (0.33-5.15)	1.31 (0.33-5.15)
BMI (kg/m ²) < 25	249/71	79/28	6/0	1.00	1.24 (0.75-2.06)	1.06 (0.61-1.85)	—	—	—	0.98 (0.57-1.70)	—	—	—	—	—	—	—	—	—	—	—	—
≥25	207/93	80/16	11/3	1.00	0.45 (0.25-0.80)	0.40 (0.21-0.75)	0.61 (0.17-2.23)	0.65 (0.17-2.45)	0.47 (0.27-0.81)	0.43 (0.24-0.77)	0.72 (0.20-2.62)	0.72 (0.20-2.62)	0.72 (0.21-2.99)	0.72 (0.21-2.99)	0.72 (0.21-2.99)	0.72 (0.21-2.99)	0.72 (0.21-2.99)	0.72 (0.21-2.99)	0.72 (0.21-2.99)	0.72 (0.21-2.99)	0.72 (0.21-2.99)	0.72 (0.21-2.99)
Without central obesity	112/42	36/10	5/1	1.00	0.74 (0.34-1.62)	0.75 (0.32-1.77)	0.53 (0.06-4.70)	0.68 (0.07-6.19)	0.72 (0.34-1.52)	0.74 (0.33-1.68)	0.57 (0.07-4.99)	0.57 (0.07-4.99)	0.57 (0.08-6.51)	0.57 (0.08-6.51)	0.57 (0.08-6.51)	0.57 (0.08-6.51)	0.57 (0.08-6.51)	0.57 (0.08-6.51)	0.57 (0.08-6.51)	0.57 (0.08-6.51)	0.57 (0.08-6.51)	0.57 (0.08-6.51)
With central obesity	344/122	123/34	12/2	1.00	0.78 (0.51-1.20)	0.67 (0.42-1.07)	0.47 (0.10-2.13)	0.46 (0.10-2.13)	0.75 (0.49-1.45)	0.65 (0.41-1.03)	0.50 (0.11-2.25)	0.50 (0.11-2.25)	0.51 (0.11-2.34)	0.51 (0.11-2.34)	0.51 (0.11-2.34)	0.51 (0.11-2.34)	0.51 (0.11-2.34)	0.51 (0.11-2.34)	0.51 (0.11-2.34)	0.51 (0.11-2.34)	0.51 (0.11-2.34)	0.51 (0.11-2.34)

OR: odds ratio; CI: confidence interval. * Adjusted for age, gender, systolic blood pressure, fasting glucose, total cholesterol, triglycerides, low-density lipoprotein cholesterol, course of hypertension, family history of hypertension, and family history of stroke.

TABLE 4: Gene interaction analysis by GMDR method.

Model combination	Training sample accuracy	Verify sample accuracy	Cross-validation consistency	<i>P</i>
SNP3 SNP5	0.567	0.527	4/10	0.1719
SNP1 SNP4 SNP5	0.578	0.542	4/10	0.1719
SNP1 SNP2 SNP4 SNP5	0.592	0.536	6/10	0.1719
SNP1 SNP2 SNP3 SNP4 SNP5	0.608	0.542	10/10	0.1719

TABLE 5: Association between haplotypes of three SNPs of the *AQP7* gene and the risk of stroke.

Haplotype	rs62542743	rs3758269	rs2989924	Frequency	OR (95% CI)	<i>P</i>
Haplo.base	C	C	G	0.5520	1.00	—
Haplo.2	C	C	A	0.2804	1.12 (0.86-1.47)	0.430
Haplo.3	C	T	A	0.1243	1.56 (1.05-2.32)	0.029

TABLE 6: Association between haplotypes of two SNPs of the *AQP9* gene and the risk of stroke.

Haplotype	rs62542743	rs16939881	Frequency	OR (95% CI)	<i>P</i>
Haplo.base	C	C	0.8308	1.00	—
Haplo.2	T	C	0.1004	1.05 (0.72-1.53)	0.80
Haplo.base	C	C	0.8308	1.00	—

TABLE 7: The logistic analysis of 5 SNPs and the risk of stroke.

Variables	OR	95% CI	<i>P</i>
rs2889924AG vs. GG	0.513	0.353-0.746	<0.001
AA vs.GG	0.782	0.496-1.232	0.782
Higher TC	1.307	1.047-1.631	0.018
Fasting blood glucose	1.144	1.037-1.263	0.007
Systolic blood pressure	1.015	1.001-1.029	0.041

However, Asian women have a higher prevalence of abdominal and visceral adiposity than Caucasian women with the same BMI, indicating a significant correlation between WC and visceral adiposity volume [32]. Oikonomou et al. reported that the *AQP7* messenger RNA (mRNA) increased in younger obese prepubertal (YOP) children but decreased in the obese adolescents (OA) who also had increased insulin and homeostatic model assessment-insulin resistance (HOMA-IR) [33]. This may be due to the expression of *AQP7* in adipose tissue where it facilitates the efflux of glycerol, and *AQP7* deficiency has been linked to increased glycerol kinase activity and triglyceride accumulation in adipose tissue, leading to obesity and secondary development of insulin resistance [10]. No association between *AQP9* and the risk of stroke was found in this study, but *AQP9* was found to be the only aquaglyceroporin expressed in the brain and was detected in tanocytes [34]. In addition, *AQP9* and *AQP4* can act in synergy contributing to the facilitation of water movements between the CSF and brain parenchyma [35].

LD is the nonrandom association of alleles at different loci and plays an important role in diverse aspects of human

genetics. In this study, we found that there was a strong linkage disequilibrium between rs3758269 and rs2989924 of the *AQP7* gene ($D' = 0.9992$), while the LD relationships between the other loci were weak. By analyzing the LD association strength, the appropriate tagSNP is selected, which can help to reduce the number of SNPs in the study and find the SNP associated with the disease as a marker. A set of SNP combinations on the same chromosome that are interrelated and tend to be passed on to offspring as a whole is called haplotype. Haplotype association studies are more useful than single SNP analysis. In this study, *AQP7* and *AQP9* loci constitute haplotypes, and more than 5% of the haplotypes were included in the study. This study found that individuals carrying CTA haplotypes in *AQP7* had a higher risk of stroke than those with the highest frequency of CCG haplotypes (OR 1.56, 95% CI 1.05-2.32).

Multivariate logistic regression analysis was conducted to investigate the possible genetic and environmental factors associated with stroke, and results showed that higher TC was a risk factor of stroke (OR 1.31, 95% CI 1.05-1.63) and rs2889924 AG vs. GG (A>G) was a protective one. However, no interaction between *AQP7* and *AQP9* was found in this study.

This is the first study in China, which investigated the association between genetic variants in *AQP7* and *AQP9* and the risk of stroke among hypertensive patients. Nested with a large cohort study including more than 12,000 hypertensive cases with a >5-year follow-up, this case-control study included strictly matched cases and controls, so that results from this study are more likely to be reliable. In the future, further studies using large and independent samples are warranted to confirm the findings of this study. Moreover, future

functional studies on associated SNPs will help to better understand the underlying biological mechanisms. To summarize, this study suggests that the genetic variants of *AQP7* SNPs are associated with the risk of stroke among patients with hypertension in China.

Data Availability

The data used to support the findings of this study are available from the corresponding author upon reasonable request.

Conflicts of Interest

The authors declared no conflict of interests.

Authors' Contributions

Qingyun Tu and Li Yan contributed equally to this work.

Acknowledgments

This study was supported by the Natural Science Foundations of China (81573199) and the Natural Science Foundation of Jiangsu Province (BK20151593).

References

- [1] V. L. Feigin, B. Norrving, and G. A. Mensah, "Global burden of stroke," *Circulation Research*, vol. 120, no. 3, pp. 439–448, 2017.
- [2] Y. Li, L. Chen, S. Yao et al., "Association of polymorphisms of the matrix metalloproteinase 9 gene with ischaemic stroke in a southern Chinese population," *Cellular Physiology and Biochemistry*, vol. 49, no. 6, pp. 2188–2199, 2018.
- [3] D. Liu, L. Liu, Z. Song, Z. Hu, J. Liu, and D. Hou, "Genetic variations of oxidative stress related genes *ALOX5*, *ALOX5AP* and *MPO* modulate ischemic stroke susceptibility through main effects and epistatic interactions in a Chinese population," *Cellular Physiology and Biochemistry*, vol. 43, no. 4, pp. 1588–1602, 2017.
- [4] Y. H. Yue, X. D. Bai, H. J. Zhang et al., "Gene polymorphisms affect the effectiveness of atorvastatin in treating ischemic stroke patients," *Cellular Physiology and Biochemistry*, vol. 39, no. 2, pp. 630–638, 2016.
- [5] W. He, M. Lu, G. Li, Z. Sun, D. Liu, and L. Gu, "Methylene tetrahydrofolate reductase (*MTHFR*) rs868014 polymorphism regulated by miR-1203 associates with risk and short term outcome of ischemic stroke," *Cellular Physiology and Biochemistry*, vol. 41, no. 2, pp. 701–710, 2017.
- [6] E. Floßmann, U. G. R. Schulz, and P. M. Rothwell, "Systematic review of methods and results of studies of the genetic epidemiology of ischemic stroke," *Stroke*, vol. 35, no. 1, pp. 212–227, 2004.
- [7] B. Jeffs, J. S. Clark, N. H. Anderson et al., "Sensitivity to cerebral ischaemic insult in a rat model of stroke is determined by a single genetic locus," *Nature Genetics*, vol. 16, no. 4, pp. 364–367, 1997.
- [8] L. C. Hsu, L. S. Hsu, and T. H. Lee, "Association of apolipoprotein A1 and A5 polymorphisms with stroke subtypes in Han Chinese people in Taiwan," *Gene*, vol. 684, pp. 76–81, 2019.
- [9] F. Gregoire, V. Lucidi, A. Zerrad-Saadi et al., "Analysis of aquaporin expression in liver with a focus on hepatocytes," *Histochemistry and Cell Biology*, vol. 144, no. 4, pp. 347–363, 2015.
- [10] F. M. Iena and J. Lebeck, "Implications of aquaglyceroporin 7 in energy metabolism," *International Journal of Molecular Sciences*, vol. 19, no. 1, p. 154, 2018.
- [11] A. Rojek, J. Praetorius, J. Frokiaer, S. Nielsen, and R. A. Fenton, "A current view of the mammalian aquaglyceroporins," *Annual Review of Physiology*, vol. 70, no. 1, pp. 301–327, 2008.
- [12] A. S. Verkman, "Novel roles of aquaporins revealed by phenotype analysis of knockout mice," *Reviews of Physiology, Biochemistry and Pharmacology*, vol. 155, pp. 31–55, 2005.
- [13] T. Hibuse, N. Maeda, A. Nagasawa, and T. Funahashi, "Aquaporins and glycerol metabolism," *Biochimica et Biophysica Acta (BBA) - Biomembranes*, vol. 1758, no. 8, pp. 1004–1011, 2006.
- [14] M. Hara-Chikuma and A. S. Verkman, "Physiological roles of glycerol-transporting aquaporins: the aquaglyceroporins," *Cellular and Molecular Life Sciences*, vol. 63, no. 12, pp. 1386–1392, 2006.
- [15] A. Rodriguez, V. Catalan, J. Gomez-Ambrosi, and G. Fruhbeck, "Role of aquaporin-7 in the pathophysiological control of fat accumulation in mice," *FEBS Letters*, vol. 580, no. 20, pp. 4771–4776, 2006.
- [16] N. Maeda, "Implications of aquaglyceroporins 7 and 9 in glycerol metabolism and metabolic syndrome," *Molecular Aspects of Medicine*, vol. 33, no. 5-6, pp. 665–675, 2012.
- [17] L. E. Prelich, L. Ma, J. C. Wright, S. Singh, X. Geng, and Y. Ding, "Progress in AQP research and new developments in therapeutic approaches to ischemic and hemorrhagic stroke," *International Journal of Molecular Sciences*, vol. 17, no. 7, article 1146, 2016.
- [18] H. Kuriyama, I. Shimomura, K. Kishida et al., "Coordinated regulation of fat-specific and liver-specific glycerol channels, aquaporin adipose and aquaporin 9," *Diabetes*, vol. 51, no. 10, pp. 2915–2921, 2002.
- [19] J. Lebeck, "Metabolic impact of the glycerol channels AQP7 and AQP9 in adipose tissue and liver," *Journal of Molecular Endocrinology*, vol. 52, no. 2, pp. R165–R178, 2014.
- [20] E. Berger, S. Héraud, A. Mojallal et al., "Pathways commonly dysregulated in mouse and human obese adipose tissue: FAT/CD36 modulates differentiation and lipogenesis," *Adipocyte*, vol. 4, no. 3, pp. 161–180, 2015.
- [21] G. Calamita, P. Gena, D. Ferri et al., "Biophysical assessment of aquaporin-9 as principal facilitative pathway in mouse liver import of glucogenetic glycerol," *Biology of the Cell*, vol. 104, no. 6, pp. 342–351, 2012.
- [22] S. Jelen, S. Wacker, C. Aponte-Santamaría et al., "Aquaporin-9 protein is the primary route of hepatocyte glycerol uptake for glycerol gluconeogenesis in mice," *The Journal of Biological Chemistry*, vol. 286, no. 52, pp. 44319–44325, 2011.
- [23] M. de Castro Ribeiro, L. Hirt, J. Bogousslavsky, L. Regli, and J. Badaut, "Time course of aquaporin expression after transient focal cerebral ischemia in mice," *Journal of Neuroscience Research*, vol. 83, no. 7, pp. 1231–1240, 2006.
- [24] Y. Tian, C. Jiang, M. Wang et al., "BMI, leisure-time physical activity, and physical fitness in adults in China: results from a series of national surveys, 2000–14," *The Lancet Diabetes and Endocrinology*, vol. 4, no. 6, pp. 487–497, 2016.

- [25] Y. Xu, L. Wang, J. He et al., "Prevalence and control of diabetes in Chinese adults," *JAMA*, vol. 310, no. 9, pp. 948–959, 2013.
- [26] Y. Wang, G. Chen, Q. Tu et al., "Associations between aquaglyceroporin gene polymorphisms and risk of type 2 diabetes mellitus," *BioMed Research International*, vol. 2018, Article ID 8167538, 7 pages, 2018.
- [27] A. Paganini-Hill, "Hormone replacement therapy and stroke: risk, protection or no effect?," *Maturitas*, vol. 38, no. 3, pp. 243–261, 2001.
- [28] J. W. Simpkins, S. H. Yang, Y. Wen, and M. Singh, "Estrogens, progestins, menopause and neurodegeneration: basic and clinical studies," *Cellular and Molecular Life Sciences*, vol. 62, no. 3, pp. 271–280, 2005.
- [29] L. Holmegaard, C. Jensen, P. Redfors, C. Blomstrand, C. Jern, and K. Jood, "Long-term progression of white matter hyperintensities in ischemic stroke," *Acta Neurologica Scandinavica*, vol. 138, no. 6, pp. 548–556, 2018.
- [30] T. H. Kim and R. Vemuganti, "Effect of sex and age interactions on functional outcome after stroke," *CNS Neuroscience & Therapeutics*, vol. 21, no. 4, pp. 327–336, 2015.
- [31] S. Prudente, E. Flex, E. Morini et al., "A functional variant of the adipocyte glycerol channel aquaporin 7 gene is associated with obesity and related metabolic abnormalities," *Diabetes*, vol. 56, no. 5, pp. 1468–1474, 2007.
- [32] U. Lim, T. Ernst, S. D. Buchthal et al., "Asian women have greater abdominal and visceral adiposity than Caucasian women with similar body mass index," *Nutrition & Diabetes*, vol. 1, no. 5, article e6, 2011.
- [33] E. Oikonomou, E. Kostopoulou, A. P. Rojas-Gil, G. Georgiou, and B. E. Spiliotis, "Adipocyte aquaporin 7 (AQP7) expression in lean children and children with obesity. Possible involvement in molecular mechanisms of childhood obesity," *Journal of Pediatric Endocrinology & Metabolism*, vol. 31, no. 10, pp. 1081–1089, 2018.
- [34] J. Badaut and L. Regli, "Distribution and possible roles of aquaporin 9 in the brain," *Neuroscience*, vol. 129, no. 4, pp. 971–981, 2004.
- [35] M. Amiry-Moghaddam, H. Lindland, S. Zelenin et al., "Brain mitochondria contain aquaporin water channels: evidence for the expression of a short AQP9 isoform in the inner mitochondrial membrane," *The FASEB Journal*, vol. 19, no. 11, pp. 1459–1467, 2005.