



Get Clarity On Generics

Cost-Effective CT & MRI Contrast Agents

 **FRESENIUS
KABI**

[WATCH VIDEO](#)

AJNR

The Association between FLAIR Vascular Hyperintensity and Stroke Outcome Varies with Time from Onset

W.J. Shang, H.B. Chen, L.M. Shu, H.Q. Liao, X.Y. Huang, S. Xiao and H. Hong

This information is current as of August 29, 2025.

AJNR Am J Neuroradiol 2019, 40 (8) 1317-1322

doi: <https://doi.org/10.3174/ajnr.A6142>

<http://www.ajnr.org/content/40/8/1317>

The Association between FLAIR Vascular Hyperintensity and Stroke Outcome Varies with Time from Onset

W.J. Shang, H.B. Chen, L.M. Shu, H.Q. Liao, X.Y. Huang, S. Xiao, and H. Hong



ABSTRACT

BACKGROUND AND PURPOSE: FLAIR vascular hyperintensity has been recognized as a marker of collaterals in ischemic stroke, but the impact on outcome is still controversial. We hypothesized that the association between FLAIR vascular hyperintensity and outcome varies with time.

MATERIALS AND METHODS: We included 459 consecutive patients with middle cerebral artery stroke and divided them into 3 groups by symptom-to-MR imaging time (group 1, ≤ 7 days; group 2, 8–14 days; group 3, ≥ 15 days). The FLAIR vascular hyperintensity score, ranging from 0 to 3 points, was based on territory distributions of different MCA segments. The associations between FLAIR vascular hyperintensity and outcome with time were analyzed qualitatively and quantitatively.

RESULTS: No patients underwent MR imaging within 6 hours of onset. The proportion of FLAIR vascular hyperintensity (+) and severe stenosis or occlusion of MCA was not significantly dependent on time. In groups 1 and 2, FLAIR vascular hyperintensity (+) was significantly associated with larger lesions, the prevalence of flow injury, and unfavorable outcome ($mRS \geq 2$). There were no such associations in group 3. Multiple logistic regressions demonstrated that FLAIR vascular hyperintensity (+) was an independent risk factor for unfavorable outcome in group 2. Infarction volume tended to increase with the increase of the distal FLAIR vascular hyperintensity score in groups 1 and 2, while declining in group 3.

CONCLUSIONS: FLAIR vascular hyperintensity is associated with unfavorable outcome within 6 hours to 14 days of onset, while the wider distribution of distal FLAIR vascular hyperintensity may be favorable beyond 14 days of onset in MCA infarction. Symptom-to-MR imaging time should be considered when assessing the prognostic value of FLAIR vascular hyperintensity.

ABBREVIATION: FVH = FLAIR vascular hyperintensity

Cerebral ischemic damage depends on both the degree and duration of hypoperfusion. However, collaterals neither de-

velop rapidly nor remain invariant after arterial occlusion. Some collaterals such as leptomeningeal anastomoses may be anatomically present, though enhanced capacity for cerebral blood flow likely requires time to develop.^{1–5} Angiogenesis induced by hypoxia and increased fluid shear stress also requires several days to weeks.^{1,2} Furthermore, the incipient development of collaterals does not guarantee their persistence.¹ Therefore, the prognostic significance of an observation of the presence of collaterals after ischemic stroke varies with time.

FLAIR vascular hyperintensity (FVH), seen in some patients with ischemic stroke, is known to be associated with stenosis and occlusion of vessels and the attendant slow flow.^{6–9} FVH is also thought to be a marker of collaterals and a predictor of outcome.^{4,6–23} However, the relationship between these interpreta-

Received March 13, 2019; accepted after revision June 17.

From the Department of Neurology (W.J.S., H.B.C., L.M.S., S.X., H.H.), National Key Clinical Department and Key Discipline of Neurology, The First Affiliated Hospital, Sun Yat-sen University, Guangzhou, China; and Department of Neurology (L.M.S., H.Q.L., X.Y.H.), The Seventh Affiliated Hospital, Sun Yat-sen University, Shenzhen, China.

W.J. Shang and H.B. Chen contributed to the work equally.

The study was supported by grants from: National Natural Science Foundation of China (No. 81671134); Medical Scientific Research Foundation of Guangdong Province of China (No. A2017219); Guangzhou Science and Technology Programme (No. 2014Y2-00502); Guangdong Provincial Key Laboratory for Diagnosis and Treatment of Major Neurological Diseases (No. 2017B030314103); Southern China International Cooperation Base for Early Intervention and Functional Rehabilitation of Neurological Diseases (No. 2015B050501003); Guangdong Provincial Engineering Center for Major Neurological Disease Treatment; and Guangdong Provincial Translational Medicine Innovation Platform for Diagnosis and Treatment of Major Neurological Disease.

Please address correspondence to Hua Hong, Department of Neurology, National Key Clinical Department and Key Discipline of Neurology, The First Affiliated Hospital, Sun Yat-sen University, 58 Zhongshan 2nd Rd, Guangzhou, Guangdong Province, China, 510080; e-mail: hhsuns@126.com

Indicates open access to non-subscribers at www.ajnr.org

Indicates article with supplemental on-line tables.

Indicates article with supplemental on-line photos.

<http://dx.doi.org/10.3174/ajnr.A6142>

tions is confusing. FVH may represent slow collateral flow effective in maintaining perfusion to penumbral regions, restricting the progress of ischemic lesions and improving outcome.¹⁰⁻¹⁷ However, FVH can also seem to correspond to a perfusion deficit, larger lesions, and poor outcomes.^{6-8,18-23} The exact reason for such a huge discrepancy is still unclear. In addition to the differences in patient populations and FVH classifications, we found that many reported symptom-to-MR imaging times ranged from 3 hours to several days. Additionally, Maeda et al²⁴ showed that FVH decreased with time, indicating that differences in symptom-to-MR imaging time might account for the discrepancy.

Moreover, most studies^{6-8,10-18,21,22} were performed during the superacute or acute stages of ischemic stroke; however, many patients with subacute or even chronic ischemic strokes present to hospitals and undergo MR imaging. FVH has been observed in areas supplied by collateral blood flow in patients with symptom-to-MR imaging times that are several weeks in duration.²⁵ Given that FVH is easily obtained via MR imaging, it could serve as a helpful predictive measure in patients with ischemic stroke.

We hypothesized that the prognostic value of FVH in ischemic stroke might change with the MR imaging observation time relative to onset. Therefore, we here evaluate the dependence of the relationship between FVH and outcome on symptom-to-MR imaging time, in cases of middle cerebral artery ischemic stroke.

MATERIALS AND METHODS

Study Population

We retrospectively analyzed the records of patients with ischemic stroke treated in the First Affiliated Hospital of Sun Yat-sen University from April 2009 to February 2017. The inclusion criteria were the following: 1) patients with responsible lesions within the MCA distribution territory; 2) patients with FLAIR and 3D time-of-flight MRA images; 3) patients with ipsilateral M1-MCA stenosis or occlusion identified by 3D time-of-flight MRA; and 4) patients with a complete evaluation, which included a personal history, vascular risk factors (smoking, hypertension, diabetes mellitus, hyperlipidemia, previous stroke/transient ischemic attack, coronary artery disease, and atrial fibrillation), routine blood tests, and cardiologic work-up. Scores on the NIHSS at admission, mRS at discharge, and hospitalization days were also collected. We excluded the following: 1) patients with concomitant anterior cerebral artery or posterior circulation strokes; 2) patients with severe artifacts on FLAIR images; 3) patients with severe stenosis or occlusion of the ipsilateral internal carotid artery; and 4) patients having undergone endovascular therapy. Patients were divided into 3 groups according to symptom-to-MR imaging time: group 1, ≤ 7 days; group 2, 8–14 days; group 3, ≥ 15 days.

MR Imaging

MR imaging was performed on 3T scanners (Magnetom Trio and Magnetom Verio; Siemens, Erlangen, Germany) with a 12-channel head coil. The neuroimaging protocol comprised T1WI (TR, 500 ms; TE, 8.9 ms), T2WI (TR, 4000 ms; TE, 100 ms), FLAIR (TR, 9000 ms; TE, 111 ms; TI, 2500 ms), and DWI (TR, 5800 ms; TE, 100 ms; matrix number, 384×384 ; two b-values of 0 and 1000 s/mm²). All patients' imaging data were independently reviewed by 2 readers who were blinded to the clinical data and

follow-up. Reader A had 18 years of neuroradiology experience, and reader B had 6 years of neuroradiology experience.

FVH

FVHs were defined as focal, tubular, or serpentine hyperintensities in the subarachnoid space relative to CSF and corresponding to the typical arterial course.¹¹ On the basis of the location and extent, FVHs were classified into proximal and distal FVHs. Proximal FVHs were defined as the presence of FVHs only within the territories of the MCA M1 and/or M2 segments. Distal FVHs were defined as the presence of FVHs in the MCA M3 and/or distal segments, which were further classified into anterior and posterior for the superior and inferior trunks of the MCA, respectively. The M3-segment FVHs, anterior FVHs, and posterior FVHs were all scored as 1 point.

Lesion Quantification

We used the ASPECTS on MR imaging.²⁶ The vascular territories were classified into perforator, pial, border zone, and large-territory infarct.²⁷

Angiography

The signal intensity of the MCA on the intracranial MRA was visually classified into the following 4 grades according to ability to visualize the MCA: All M3 branches of the MCA were visualizable to the cortical surface (grade A); ≥ 1 M3 branch could not be visualized on the cortical surface (grade B); the superior or inferior trunk of the MCA or ≥ 1 M2 branch could not be visualized along its course (grade C); and the M1 could not be visualized along its course (grade D) (On-line Fig 1).²⁸ Posterior cerebral artery laterality was considered to be present if ≥ 1 segmental extent of the ipsilateral posterior cerebral artery was observable on MRA.⁵

Statistical Analysis

The κ coefficient was used to assess interobserver agreement for FVH. Continuous variables with a normal distribution were described as mean \pm SD, and non-normally distributed variables were described as median and interquartile range. We compared variables between the patients with and without FVH in each group using a Mann-Whitney *U* test, Student *t* test, Pearson χ^2 test, or χ^2 test with continuity correction depending on the type of variable. We further compared FVH scores and baseline characteristics in all 3 groups using the Kruskal-Wallis-test or Pearson χ^2 test as appropriate. A multiple binary logistic regression analysis was applied to identify independent predictors of unfavorable scores on the modified Rankin Scale at discharge (mRS ≥ 2), signifying an unfavorable outcome. All covariates with a *P* value $\leq .1$ in a univariate analysis were entered into this logistic regression model. The results are presented as odds ratios and the 95% confidence intervals. A 2-tailed value of *P* $< .05$ indicated statistical significance (SPSS for Windows, Version 20.0; IBM, Armonk, New York).

RESULTS

In total, 579 consecutive patients met all the inclusion criteria, and 120 patients were excluded (concomitant anterior cerebral

Table 1: Clinical and demographic patient characteristics by symptom-to-MRI time and the presence of FVH^a

Characteristics	Group 1 (n = 191)			Group 2 (n = 164)			Group 3 (n = 104)		
	FVH (-) (n = 74)	FVH (+) (n = 117)	P Value	FVH (-) (n = 63)	FVH (+) (n = 101)	P Value	FVH (-) (n = 44)	FVH (+) (n = 60)	P Value
Women	27 (36.5)	37 (31.6)	.53	20 (31.7)	27 (26.7)	.60	10 (22.7)	20 (33.3)	.28
Age (yr)	60.7 ± 14.3	63.9 ± 12.5	.11	66 (46–73)	65.5 (54–74)	.88	55.1 ± 13.7	63.6 ± 11.0	.001 ^b
Symptom-to-admission time (days)	3 (2–4)	2 (1–4)	.05	5 (4–7)	4 (3–7)	.21	16.5 (12–31)	19.5 (12–30.75)	.79
Risk factors									
Hypertension	58 (78.4)	82 (70.1)	.24	35 (55.6)	70 (69.3)	.09	31 (70.5)	45 (75)	.66
Diabetes mellitus	29 (39.2)	28 (23.9)	.03 ^b	16 (25.4)	36 (35.6)	.23	12 (27.3)	13 (21.7)	.64
Hypercholesterolemia	24 (32.4)	43 (36.8)	.64	24 (38.1)	30 (29.7)	.31	8 (18.2)	8 (13.3)	.59
Smoker	24 (32.4)	52 (44.4)	.13	25 (39.7)	48 (47.5)	.34	19 (43.2)	26 (43.3)	1.00
Coronary heart disease	5 (6.8)	6 (5.1)	.88	1 (1.6)	6 (5.9)	.35	6 (13.6)	5 (8.3)	.59
Atrial fibrillation	4 (5.4)	8 (6.8)	.93	2 (3.2)	1 (1.0)	.88	1 (2.3)	2 (3.3)	1.00
Previous stroke	18 (24.3)	29 (24.8)	1.00	9 (14.3)	25 (24.8)	.12	9 (20.5)	19 (31.7)	.27
NIHSS score at admission	4 (2–7)	6 (3–10)	.02 ^b	4 (2–9)	5 (2–9)	.32	5 (2–10)	4.5 (2–9)	.56
Hospitalization (days)	10.5 (8–13)	13 (10–16)	.001 ^b	12 (9–15)	13 (9–15)	.95	13.5 (8–15)	12.5 (9.25–16)	.94
Outcome at discharge									
mRS	2 (1–3)	3 (2–4)	<.001 ^b	2 (1–3)	3 (2–4)	.10	2.5 (1–4)	3 (1.25–4)	.54
mRS ≥ 2	54 (73.0)	100 (85.5)	.04 ^b	43 (68.3)	85 (84.2)	.02 ^b	31 (70.5)	45 (75.0)	.66
mRS ≥ 3	31 (41.9)	70 (59.8)	.02 ^b	29 (46.0)	52 (51.5)	.52	22 (50.0)	34 (56.7)	.55

Note:—(–) indicates negative; (+), positive.

^a Data are No. (%), mean ± SD, and median (interquartile range).

^b $P < .05$.

artery or posterior circulation strokes, $n = 27$; severe artifacts on FLAIR, $n = 18$; severe stenosis or occlusion of the ipsilateral internal carotid artery, $n = 47$; endovascular therapy, $n = 28$). We finally included 459 patients, and FVH was observed in 278 patients (60.6%). The interobserver agreement for FVH was $\kappa = 0.88$ (95% CI, 0.68–1.00). The proportion of observed FVH (+) was not significantly dependent on time (61.3%, 61.6%, and 57.7%; $P = .79$). Details of the clinical and demographic characteristics of patients by symptom-to-MR imaging time and the presence of FVH are presented in Table 1. In group 1, patients with FVH demonstrated a higher NIHSS score at admission ($P = .02$), a longer hospitalization ($P = .001$), a higher mRS at discharge ($P = .001$), and a higher prevalence of unfavorable outcome ($mRS \geq 2$; $P = .04$) or poor outcome ($mRS \geq 3$; $P = .02$) than those without FVH. In group 2, patients with FVH demonstrated a higher prevalence of unfavorable outcome ($mRS \geq 2$; $P = .02$). However, in group 3, we found no significant difference in outcome between patients with and without FVH.

Details of the radiologic characteristics of patients by symptom-to-MR imaging time and FVH status are presented in On-line Table 1, Figs 1–3, and On-line Figs 2–4. No patients underwent MR imaging within 6 hours of onset. More patients with FVH demonstrated severe stenosis or occlusion of the MCA than those without it, independent of time (group 1, $P < .001$; group 2, $P < .001$; group 3, $P = .001$). In groups 1 and 2, patients with FVH demonstrated larger cortical extents of MCA and total lesion scores than those without it (group 1, $P < .001$, $P < .001$; group 2, $P = .003$, $P = .003$), and poorer MCA signal intensities (grades B–D; group 1, $P < .001$; group 2, $P < .001$). However, these differences were absent in group 3 ($P = .73$, $P = .45$, $P = .18$). We observed no significant differences in FVH scores ($P = 0.25$) across the 3 groups. The MCA cortical area and total lesion volume tended to increase with the distal FVH score in groups 1 and 2 and decrease in group 3 (Fig 3). Details of the baseline characteristics of the 3 groups are presented in On-line Table 2. There were no differences among the 3 groups, apart from hypercholesterolemia.

Variables associated with an unfavorable clinical outcome are shown in On-line Table 3. In group 1, sex ($P = .01$), NIHSS score on admission ($P < .001$), infarct volume ($P = .001$), and FVH ($P = .04$) were included in our analysis. After we adjusted for representative variables, FVH (+) did not appear as an independent predictor of unfavorable outcome ($P = .14$). In group 2, age ($P = .003$), NIHSS score on admission ($P < .001$), infarct volume ($P = .001$), FVH ($P = .002$), posterior cerebral artery laterality ($P = .008$), and MCA severe stenosis or occlusion ($P = .006$) were included in our analysis. After we adjusted for representative variables, FVH (+) appeared as an independent predictor of unfavorable outcome (odds ratio, 2.95; 95% CI, 1.06–8.18; $P = .04$). In group 3, FVH (+) was not included in our analysis ($P = .66$).

DISCUSSION

Our novel results demonstrate that the clinical value of FVH varies with time. Within 6 hours to 14 days of onset, FVH (+) is associated with greater lesion volume, poorer MCA signal intensity, and unfavorable outcome. These associations disappear beyond 14 days of onset. In addition, the lesion volume tended to increase with the distal FVH score within 6 hours to 14 days of onset and to decrease beyond 14 days of onset. Therefore, the symptom-to-MR imaging time, not just the presence or distribution of FVH, played an important role in predicting stroke outcomes in this study.

FVH is recognized as a marker of slow flow induced by severe stenosis or occlusion of vessels.^{6–9} The presence of FVH ranges from 44.1% to 100% in patients with stroke within 24 hours of onset,^{11,29} which increases to 75.9%–100% when accompanied by severe stenosis or occlusion of the MCA or ICA.^{11,12} Maeda et al²⁴ showed that the presence of FVH decreases from 100% within 24 hours of onset to 50% within 10–13 days of onset. In the present study, the proportion of severe stenosis or occlusion of the MCA was as high as 80.6%, and most of these patients did not undergo MR imaging a short time after onset, which could explain the median prevalence of FVH (60.6%). In addition, we saw no significant differences in the prevalence of FVH in our 3 groups,

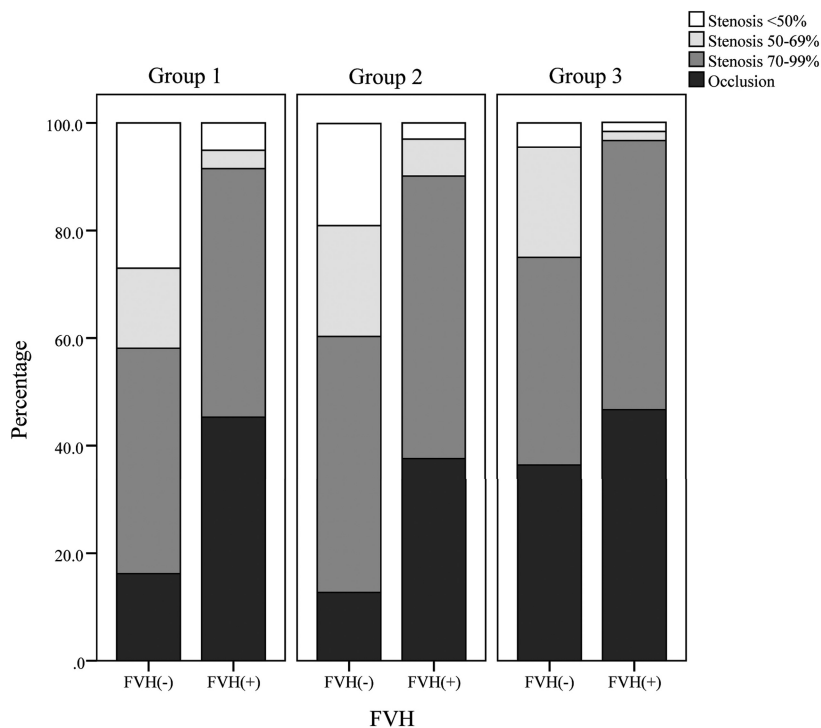


FIG 1. Detailed middle cerebral artery vascular status by symptom-to-MR imaging time, which increases with group number and the presence of FLAIR vascular hyperintensity.

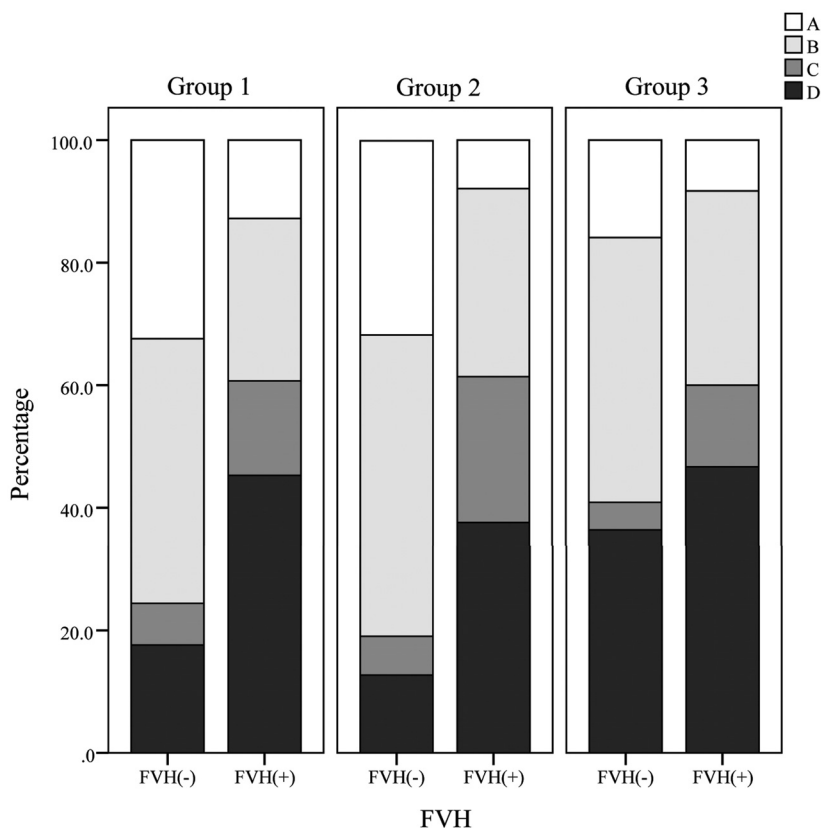


FIG 2. Detailed middle cerebral artery signal intensity by symptom-to-MR imaging time, which increases with group number and the presence of FLAIR vascular hyperintensity. A, All M3 branches of the MCA are visualized on the cortical surface. B, One or more M3 branches are not visualized on the cortical surface. C, The superior or inferior trunk of the MCA or ≥ 1 M2 branch is not visualized along its course. D, M1 branches are not visualized along the course of the MCA.

unlike Maeda et al.²⁴ One reason could be that the proportion of severe stenosis or occlusion and the signal intensity of MCA were not significantly different in our 3 groups, and the Maeda et al.²⁴ study did not measure the degree of MCA lesion. Another possible reason is the much greater sample size in our study. The sample size in Maeda et al.²⁴ was 27 patients, with only 2 patients imaged within 10–13 days of onset, which would have led to serious errors.

The prognostic value of FVH has been widely investigated,^{4,6-23} with greatly divergent results. Besides populations, end points, and FVH classifications, we think that the symptom-to-MR imaging time plays an important role in explaining the discrepancies in the literature. In our study, FVH (+) was associated with unfavorable clinical and radiographic outcomes within 6 hours to 14 days of onset, but these associations disappeared beyond 14 days of onset. On reviewing previous studies, we found that the symptom-to-MR imaging time was within 6 hours of onset or within the time window of reperfusion therapy in most of the studies that showed an association between FVH and good outcome.^{10,11,13-17} On the other hand, the symptom-to-MR imaging time was between 12 and 24 hours and several days in most of the studies showing an association between FVH and a poor outcome^{6-8,18-20}; these results are similar to ours for the group within 6 hours to 14 days of onset.

We assumed that the role of FVH would vary with time. Within 6 hours of onset or within the time window of reperfusion therapy, FVH might be a marker of leptomeningeal anastomoses that are anatomically present and develop dramatically and rapidly after the onset of acute ischemic stroke.^{2,5} The areas with FVH, especially those with FVH–DWI mismatch, represent the brain tissue at risk of infarction, which could be saved by reperfusion therapy to reduce the final lesion volume and improve outcome.^{6,9,16,29} This may explain why patients with FVH have better outcomes than those without FVH in this period.

Beyond the time window of reperfusion therapy, however, the observation of persistent FVH may represent persis-

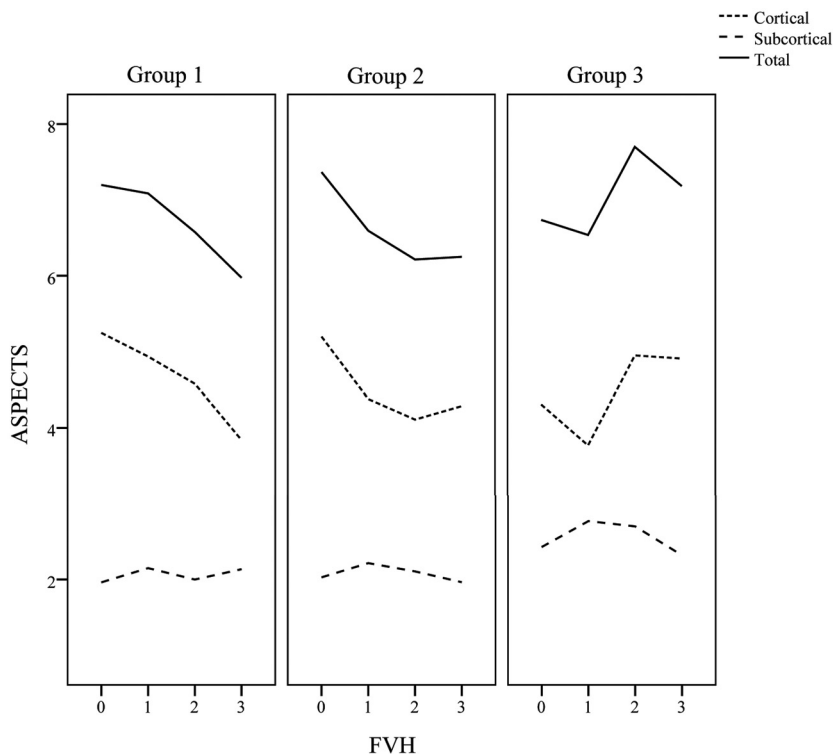


FIG 3. The ASPECTS lesion volume against the distal FLAIR vascular hyperintensity score.

tent vessel occlusion and impaired hemodynamics.^{4,14,29,30} Therefore, patients with FVH could be influenced much more by hemodynamic instability than patients without it.¹⁹ This difference might be correlated with a poor outcome in patients with FVH during this period. Some studies that re-examined MR imaging after the superacute period support this assumption.^{4,14,29} Sanossian et al¹⁴ found that a persistence of FVH on follow-up MR imaging was associated with a decreased or absent flow signal intensity on MRA, whereas the absence of FVH was seen in the setting of a return of the MR imaging flow signal intensity. This result is similar to our finding that patients with FVH had poorer MCA signal intensity. However, the observation of a decrease in FVH after reperfusion therapy predicted successful recanalization, smaller infarct volume, and good outcomes.^{4,29}

Here, FVH was not significantly associated with clinical or radiologic outcome in group 3, though we continued to see a preponderance of patients with FVH who had severe stenosis or occlusion of the MCA. In addition, the infarct volume tended to decrease with increases in the distal FVH score in group 3, which is the opposite of the results in groups 1 and 2. This outcome indicates that the wider distribution of distal FVH may be favorable when the symptom-to-MR imaging time extends beyond 14 days. We think that this possibility could be explained in the following 2 ways: First, new collaterals require time to develop significant capacity for cerebral blood flow.^{1,5} Second, achieving functional vessels through arteriogenesis requires several days to weeks.^{1,2} Therefore, these collaterals likely contribute more to alleviating the subacute or chronic stages of stroke than to mitigating acute ischemic events. The observation of no or little FVH in the subacute or chronic stages of stroke may represent an absence of effective collaterals. Future research should further refine

our knowledge of the importance of the symptom-to-MR imaging time beyond 14 days.

Unlike most studies that focused on the superacute stage of ischemic stroke, we used a much longer time span after ischemic stroke. Consequently, our study conditions more closely resembled “real world” clinical scenarios. Patients can undergo MR imaging any time after stroke onset, and using only FVH to evaluate the superacute ischemic stroke stage is insufficient for clinical work. However, FVH is relatively easy to detect on MR imaging and may therefore reflect collateral status and potentially act as a substitute for conventional angiography.³¹

This study has some limitations. First, this is a retrospective clinical study. However, it shows, by analysis, a clear time dependency of the prognostic value of FVH. Further prospective studies are needed to confirm our findings. Second, we excluded patients with endovascular therapy. Although this ex-

clusion ensures data homogeneity and avoids the influence of a diversity of treatments on outcome, it leads to no patients receiving MR imaging within 6 hours of onset in this study. However, many studies have focused on the superacute stage of stroke,^{10,11,13-17,21,22} while few²⁵ have focused on the subacute and chronic stages.

CONCLUSIONS

For patients with MCA infarctions, FVH (+) is associated with unfavorable outcomes within 6 hours to 14 days of onset, while the wider distribution of distal FVH may be favorable beyond 14 days of onset. Therefore, the symptom-to-MR imaging time should be taken into account when assessing the prognostic value of FVH.

Disclosures: Wenjin Shang—RELATED: Grant: National Natural Science Foundation of China (No. 81671134); Medical Scientific Research Foundation of Guangdong Province of China (No. A2017219); Guangzhou Science and Technology Programme (No. 2014Y2-00502). Comments: My institution is supported by the grants from Guangdong Provincial Key Laboratory for Diagnosis and Treatment of Major Neurological Diseases (No. 2017B030314103); Southern China International Cooperation Base for Early Intervention and Functional Rehabilitation of Neurological Diseases (No. 2015B050501003); Guangdong Provincial Engineering Center for Major Neurological Disease Treatment; Guangdong Provincial Translational Medicine Innovation Platform for Diagnosis and Treatment of Major Neurological Disease. Liming M. Shu—RELATED: Grant: National Natural Science Foundation of China (No. 81671134); Medical Scientific Research Foundation of Guangdong Province of China (No. A2017219); Guangzhou Science and Technology Programme (No. 2014Y2-00502).* Su Xiao—RELATED: Grant: Guangdong Provincial Key Laboratory for Diagnosis and Treatment of Major Neurological Diseases (No. 2017B030314103); Southern China International Cooperation Base for Early Intervention and Functional Rehabilitation of Neurological Diseases (No. 2015B050501003); Guangdong Provincial Engineering Center for Major Neurological Disease Treatment; Guangdong Provincial Translational Medicine Innovation Platform for Diagnosis and Treatment of Major Neurological Disease.* Hua Hong—RELATED: Grant: National Natural Science Foundation of China (No. 81671134); Medical Scientific Research Foundation of Guangdong Province of

China (No. A2017219); Guangzhou Science and Technology Programme (No. 2014Y2-00502). *Comments:* Our institution is supported by the grants from Guangdong Provincial Key Laboratory for Diagnosis and Treatment of Major Neurological Diseases (No. 2017B030314103); Southern China International Cooperation Base for Early Intervention and Functional Rehabilitation of Neurological Diseases (No. 2015B050501003); Guangdong Provincial Engineering Center for Major Neurological Disease Treatment; Guangdong Provincial Translational Medicine Innovation Platform for Diagnosis and Treatment of Major Neurological Disease; *Others:* We paid money to Editage, a division of Cactus Communications, for English editing to ensure that the language is clear and free of errors. The intent of our message was not altered in any way during the editing process. *Money paid to institution.

REFERENCES

1. Liebeskind DS. **Collateral circulation.** *Stroke* 2003;34:2279–84 CrossRef Medline
2. Iwasawa E, Ichijo M, Ishibashi S, et al. **Acute development of collateral circulation and therapeutic prospects in ischemic stroke.** *Neural Regen Res* 2016;11:368–71 CrossRef Medline
3. Bang OY, Goyal M, Liebeskind DS. **Collateral circulation in ischemic stroke: assessment tools and therapeutic strategies.** *Stroke* 2015;46:3302–09 CrossRef Medline
4. Ichijo M, Iwasawa E, Numasawa Y, et al. **Significance of development and reversion of collaterals on MRI in early neurologic improvement and long-term functional outcome after intravenous thrombolysis for ischemic stroke.** *AJNR Am J Neuroradiol* 2015;36:1839–45 CrossRef Medline
5. Uemura A, O'Uchi T, Kikuchi Y, et al. **Prominent laterality of the posterior cerebral artery at three-dimensional time-of-flight MR angiography in M1-segment middle cerebral artery occlusion.** *AJNR Am J Neuroradiol* 2004;25:88–91 Medline
6. Kamran S, Bates V, Bakshi R, et al. **Significance of hyperintense vessels on FLAIR MRI in acute stroke.** *Neurology* 2000;55:265–69 CrossRef Medline
7. Kono T, Naka H, Nomura E, et al. **The association between hyperintense vessel sign and final ischemic lesion differ in its location.** *J Stroke Cerebrovasc Dis* 2014;23:1337–43 CrossRef Medline
8. Cheng B, Ebinger M, Kufner A, et al; Stroke Imaging Repository (STIR) Investigators. **Hyperintense vessels on acute stroke fluid-attenuated inversion recovery imaging: associations with clinical and other MRI findings.** *Stroke* 2012;43:2957–61 CrossRef Medline
9. Toyoda K, Ida M, Fukuda K. **Fluid-attenuated inversion recovery intraarterial signal: an early sign of hyperacute cerebral ischemia.** *AJNR Am J Neuroradiol* 2001;22:1021–29 Medline
10. Mahdjoub E, Turc G, Legrand L, et al. **Do fluid-attenuated inversion recovery vascular hyperintensities represent good collaterals before reperfusion therapy?** *AJNR Am J Neuroradiol* 2018;39:77–83 CrossRef Medline
11. Liu D, Scalzo F, Rao NM, et al. **Fluid-attenuated inversion recovery vascular hyperintensity topography: novel imaging marker for revascularization in middle cerebral artery occlusion.** *Stroke* 2016;47:2763–69 CrossRef Medline
12. Huang X, Liu W, Zhu W, et al. **Distal hyperintense vessels on FLAIR: a prognostic indicator of acute ischemic stroke.** *Eur Neurol* 2012;68:214–20 CrossRef Medline
13. Olindo S, Chausson N, Joux J, et al. **Fluid-attenuated inversion recovery vascular hyperintensity: an early predictor of clinical outcome in proximal middle cerebral artery occlusion.** *Arch Neurol* 2012;69:1462–68 CrossRef Medline
14. Sanossian N, Saver JL, Alger JR, et al. **Angiography reveals that fluid-attenuated inversion recovery vascular hyperintensities are due to slow flow, not thrombus.** *AJNR Am J Neuroradiol* 2009;30:564–68 CrossRef Medline
15. Lee KY, Latour LL, Luby M, et al. **Distal hyperintense vessels on FLAIR: an MRI marker for collateral circulation in acute stroke?** *Neurology* 2009;72:1134–39 CrossRef Medline
16. Legrand L, Tisserand M, Turc G, et al. **Do FLAIR vascular hyperintensities beyond the DWI lesion represent the ischemic penumbra?** *AJNR Am J Neuroradiol* 2015;36:269–74 CrossRef Medline
17. de la Ossa N, Hernandez- Pérez M, Domènech S, et al. **Hyperintensity of distal vessels on FLAIR is associated with slow progression of the infarction in acute ischemic stroke.** *Cerebrovasc Dis* 2012;34:376–84 CrossRef Medline
18. Nam KW, Kwon HM, Park SW, et al. **Distal hyperintense vessel sign is associated with neurological deterioration in acute ischaemic stroke.** *Eur J Neurol* 2017;24:617–23 CrossRef Medline
19. Kim SE, Lee BI, Kim SE, et al. **Clinical significance of fluid-attenuated inversion recovery vascular hyperintensities in borderzone infarcts.** *Stroke* 2016;47:1548–54 CrossRef Medline
20. Gawlitza M, Gragert J, Quäsching U, et al. **FLAIR-hyperintense vessel sign, diffusion-perfusion mismatch and infarct growth in acute ischemic stroke without vascular recanalisation therapy.** *J Neuroradiol* 2014;41:227–33 CrossRef Medline
21. Nave AH, Kufner A, Bücke P, et al. **Hyperintense vessels, collateralization, and functional outcome in patients with stroke receiving endovascular treatment.** *Stroke* 2018;49:675–81 CrossRef Medline
22. Kufner A, Galinovic I, Ambrosi V, et al. **Hyperintense vessels on FLAIR: hemodynamic correlates and response to thrombolysis.** *AJNR Am J Neuroradiol* 2015;36:1426–30 CrossRef Medline
23. Gawlitza M, Bohme J, Maros M, et al. **FLAIR vascular hyperintensities and 4D MR angiograms for the estimation of collateral blood flow in anterior cerebral artery ischemia.** *PLoS One* 2017;12:e0172570 CrossRef Medline
24. Maeda M, Koshimoto Y, Uematsu H, et al. **Time course of arterial hyperintensity with fast fluid-attenuated inversion-recovery imaging in acute and subacute middle cerebral arterial infarction.** *J Magn Reson Imaging* 2001;13:987–90 CrossRef Medline
25. Essig M, von Kummer R, Egelhof T, et al. **Vascular MR contrast enhancement in cerebrovascular disease.** *AJNR Am J Neuroradiol* 1996;17:887–94 Medline
26. Barber PA, Demchuk AM, Zhang J, et al. **Validity and reliability of a quantitative computed tomography score in predicting outcome of hyperacute stroke before thrombolytic therapy: ASPECTS study group—Alberta Stroke Programme Early CT Score.** *Lancet* 2000;355:1670–74 CrossRef Medline
27. Lee DK, Kim JS, Kwon SU, et al. **Lesion patterns and stroke mechanism in atherosclerotic middle cerebral artery disease: early diffusion-weighted imaging study.** *Stroke* 2005;36:2583–88 CrossRef Medline
28. Hirooka R, Ogasawara K, Inoue T, et al. **Simple assessment of cerebral hemodynamics using single-slab 3D time-of-flight MR angiography in patients with cervical internal carotid artery stenocclusive diseases: comparison with quantitative perfusion single-photon emission CT.** *AJNR Am J Neuroradiol* 2009;30:559–63 CrossRef Medline
29. Sakuta K, Saji N, Aoki J, et al. **Decrease of hyperintense vessels on fluid-attenuated inversion recovery predicts good outcome in t-PA patients.** *Cerebrovasc Dis* 2016;41:211–18 CrossRef Medline
30. Liu W, Yin Q, Yao L, et al. **Decreased hyperintense vessels on FLAIR images after endovascular recanalization of symptomatic internal carotid artery occlusion.** *Eur J Radiol* 2012;81:1595–600 CrossRef Medline
31. Kim SJ, Ha YS, Ryoo S, et al. **Sulcal effacement on fluid attenuation inversion recovery magnetic resonance imaging in hyperacute stroke: association with collateral flow and clinical outcomes.** *Stroke* 2012;43:386–92 CrossRef Medline