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

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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 ≥ 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 develop 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. Angiogenesis induced by hypoxia and increased fluid shear stress also requires several days to weeks. Furthermore, the incipient development of collaterals does not guarantee their persistence. 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. FVH is also thought to be a marker of collaterals and a predictor of outcome. However, the relationship between these interpreta-

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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.\textsuperscript{10-17} However, FVH can also seem to correspond to a perfusion deficit, larger lesions, and poor outcomes.\textsuperscript{6-8,16-23} 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 al\textsuperscript{24} showed that FVH decreased with time, indicating that differences in symptom-to-MR imaging time might account for the discrepancy.

Moreover, most studies\textsuperscript{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.\textsuperscript{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, \( \leq 7 \) days; group 2, 8–14 days; group 3, \( \geq 15 \) 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 \( \times \) 384; two b-values of 0 and 1000 s/mm\(^2\)). 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.\textsuperscript{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.\textsuperscript{26} The vascular territories were classified into perforator, pial, border zone, and large-territory infarct.\textsuperscript{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); \( \geq 1 \) M3 branch could not be visualized to the cortical surface (grade B); the superior or inferior trunk of the MCA or \( \geq 1 \) 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).\textsuperscript{28} Posterior cerebral artery laterality was considered to be present if \( \geq 1 \) segmental extent of the ipsilateral posterior cerebral artery was observable on MRA.\textsuperscript{5}

**Statistical Analysis**

The \( \kappa \) 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 \( \chi^2 \) test, or \( \chi^2 \) 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 \( \chi^2 \) 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 \( \geq 2 \)), signifying an unfavorable outcome. All covariates with a \( P \) value \( < .05 \) 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...
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 κ = 0.88 (95% CI, 0.68–1.00). The proportion of observed FVH (+) was not significantly dependent on time (61.3%, 61.6%, and 0.88 (95% CI, 0.68 –1.00). The proportion of observed FVH (+) did not appear as an independent predictor of unfavorable outcome (P = .79). Details of the clinical and demographic characteristics of patients by symptom-to-MRI 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 = .001), FVH (+) demonstrated a higher NIHSS score at admission (P = .001), FVH (+) demonstrated a higher NIHSS score at admission (P = .001), and MCA severe stenosis or occlusion (P = .008) and MCA severe stenosis or occlusion (P = .006) were observed in patients with stroke within 24 hours of onset. For those with FVH, in group 2, patients with FVH demonstrated a higher prevalence of unfavorable outcome (mRS ≥ 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-MRI 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.

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

| Characteristics | Group 1 (n = 191) | Group 2 (n = 164) | Group 3 (n = 104) |
|-----------------|------------------|------------------|------------------|
|                 | FVH (n = 74)     | FVH (n = 117)    | FVH (n = 63)     | FVH (n = 101)   | FVH (n = 60)    | FVH (n = 66)    |
|                 | Value            | Value            | Value            | Value            | Value            | Value            |
|                 | P Value          | P Value          | P Value          | P Value          | P Value          | P Value          |
| Women           | 27 (16.5)        | 37 (19.6)        | 20 (31.7)        | 27 (26.7)        | 10 (22.7)        | 20 (33.3)        |
|                 | .53              |                  | .60              |                  | .28              |                  |
| Age (yr)        | 60.7 ± 14.3      | 63.9 ± 12.5      | 66 (46–73)       | 65.5 (54–74)     | 55.1 ± 13.7      | 63.6 ± 11.0      |
|                 | .11              |                  | .88              |                  | .001b            |                  |
| Symptom-to-admission time (days) | 3 (2–4) | 2 (1–4) | .05 | 5 (4–7) | 4 (3–7) | .21 |
| Risk factors    |                  |                  |                  |                  |                  |                  |
| Hypertension    | 58 (78.4)        | 82 (70.1)        | 35 (55.6)        | 70 (69.3)        | 31 (70.5)        | 45 (75.5)        |
|                 | .24              |                  | .09              |                  | .66              |                  |
| Diabetes mellitus | 29 (39.2)     | 28 (23.9)        | 16 (25.4)        | 36 (35.6)        | 12 (27.3)        | 23 (12.7)        |
|                 | .03a             |                  | .23              |                  | .64              |                  |
| Hypercholesterolemia | 24 (32.4)   | 43 (36.8)        | 24 (38.1)        | 30 (29.7)        | 8 (18.2)         | 8 (13.3)         |
|                 | .64              |                  | .31              |                  | .59              |                  |
| Smoker          | 24 (32.4)        | 52 (44.4)        | 25 (39.7)        | 48 (47.5)        | 19 (43.2)        | 26 (43.3)        |
|                 | .13              |                  | .34              |                  | 1.00             |                  |
| Coronary heart disease | 5 (6.8)    | 6 (5.1)          | 11 (6.6)         | 6 (5.9)          | 6 (13.6)         | 5 (8.3)          |
|                 | .88              |                  | .35              |                  | .59              |                  |
| Atrial fibrillation | 4 (5.4)     | 6 (4.8)          | 2 (3.2)          | 1 (1.0)          | 1 (2.3)          | 2 (3.3)          |
|                 | .93              |                  | .88              |                  | 1.00             |                  |
| Previous stroke | 38 (24.3)        | 29 (24.8)        | 9 (14.3)         | 25 (24.8)        | 9 (20.5)         | 19 (31.7)        |
|                 | 1.00             |                  | .12              |                  | .27              |                  |
| NIHSS score at admission | 4 (2–7) | 6 (3–10) | .02b | 4 (2–9) | 5 (2–9) | .32 |
|                 | 4 (2–9)          | 5 (2–9)          | 5 (2–10)         | 4.5 (2–9)        | .56              |                  |
| Hospitalization (days) | 10.5 (8–13) | 13 (10–16) | .00b | 12 (9–15) | 13 (9–15) | .95 |
|                 | 13.5 (8–15)      | 12.5 (9–15–26)   |                  |                  | .94              |                  |
| Outcome at discharge |                  |                  |                  |                  |                  |                  |
| mRS             | 2 (1–3)          | 3 (2–4)          | 2 (1–3)          | 3 (2–4)          | 2.5 (1–4)        | 3 (2.5–4)        |
|                 | .001b            |                  | .10              |                  | .54              |                  |
| mRS ≥ 2         | 54 (73.0)        | 100 (85.5)       | 43 (68.3)        | 85 (84.2)        | 31 (70.5)        | 45 (75.0)        |
|                 | .04a             |                  | .02b             |                  | .66              |                  |
| mRS ≥ 3         | 31 (41.9)        | 70 (59.8)        | 29 (46.0)        | 52 (51.5)        | 22 (50.0)        | 34 (56.7)        |
|                 | .02a             |                  | .52              |                  | .55              |                  |

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

a Data are No. (%), mean ± SD, and median (interquartile range).
b P < .05.

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-MRI 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,1,2,9 which increases to 75.9%–100% when accompanied by severe stenosis or occlusion of the MCA or ICA.11,12 Maeda et al14 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,
unlike Maeda et al.\cite{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.\cite{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.\cite{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,\cite{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.\cite{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\cite{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.\cite{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.\cite{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-
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. 

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 exclusion 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, while few 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.

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