Microembolic Signals and Diffusion-weighted MR Imaging Abnormalities in Acute Ischemic Stroke

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Microembolic Signals and Diffusion-weighted MR Imaging Abnormalities in Acute Ischemic Stroke

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BACKGROUND AND PURPOSE: The clinical significance of microembolic signals (MESs) detected by transcranial Doppler sonography (TCD) in acute ischemic stroke remains unclear. The purpose of the present study was to assess the findings of diffusion-weighted MR imaging (DWI) and other clinical characteristics in patients with acute ischemic stroke and MESs.

METHODS: We performed TCD and DWI within 48 hours and 7 days, respectively, after stroke onset in 28 patients with acute brain infarction. The relationship between the number of MESs and DWI findings, risk factors for stroke, National Institutes of Health Stroke Scale (NIHSS) score on admission, and arterial disease was examined.

RESULTS: Ten patients had MESs detected by TCD (MES group) and 18 had no MESs (control group). The frequency of hypertension, diabetes mellitus, hyperlipidemia, and smoking; NIHSS score; blood-coagulation parameters; and interval between stroke onset and DWI study did not differ between the two groups. However, arterial disease was more frequent in the MES group than in the control group. Small, multifocal ischemic lesions (<10 mm in diameter) on DWI were more frequent in the MES group than in the control group. Conventional CT and MR imaging often failed to show these lesions.

CONCLUSION: Small, often asymptomatic DWI abnormalities were more frequent in patients with MESs detected by TCD and with large-vessel occlusive diseases than in stroke patients without MESs. TCD and DWI may provide early clues to the mechanism of stroke in the acute phase.
Ten patients had MESs detected by TCD (MES group) and 18 patients had no MES on control group.

The following clinical data were compiled for all patients: the number of MESs detected from the middle cerebral artery (MCA) by TCD, DWI findings, neurologic deficits, major risk factors for stroke, arterial or cardiac diseases that could be a potential source of embolic stroke, blood-coagulation factors, and the interval between stroke onset and MR study.

Within 24 hours after admission, TCD was performed using a DWL Multidop X unit with a 2.5-MHz probe. Patients were placed supine, and bilateral MCAs were insonated at a depth of between 45 and 55 mm for 30 minutes. The TCD probe was held in place with an elastic headband to reduce the occurrence of movement artifacts. The criteria for MESs included random occurrence, brief duration (<0.1 second), intensity greater than 6 dB above background, primarily unidirectional within the Doppler spectrum, a spike in the power/ intensity trace, and an accompanying audible chrip, moan, or snap sound. All MESs were recorded on a hard disk. The MESs were counted during an off-line review of the Doppler recordings.

Brain CT studies were performed on the day of admission and again 3 days after stroke onset to evaluate ischemic lesions. Within 7 days of stroke onset, MR imaging was performed using a 1.5-T system equipped with single-shot echo-planar imaging to obtain rapid diffusion images. MR studies included axial T1-weighted, axial T2-weighted, and DWI sequences (approximately 30 minutes of scanning time). The imaging parameters were 4000/103 (TR/TE), 128 × 128 matrix, 230-mm field of view, and 4-mm slice thickness with a 2-mm gap between slices. Two b values were used: 0 and 1000 s/mm². Diffusion gradients were applied in successive scans in each of the x, y, and z directions, and DWI images were formed from the average of these values. The criterion for the diagnosis of acute infarcts on DWI was focal high intensity, judged not to be due to normal anisotropic diffusion or magnetic susceptibility artifacts. The diameter of hyperintense lesions on DWI was classified as small (<10 mm), medium (10–30 mm), or large (≥30 mm). These lesions were also categorized as cortical, subcortical, or lacunar infarcts according to their location. The number, size, and location of recent ischemic lesions on DWI were evaluated by two neurologists who had no knowledge of the clinical or laboratory data.

Neurologic deficits on admission were determined by using the National Institutes of Health Stroke Scale (NIHSS) score (23). Major risk factors for stroke included hypertension, diabetes mellitus, hyperlipidemia, smoking, NIHSS score, blood-coagulation parameters, including antithrombin III, thrombin-antithrombin III complex, and D-dimer, were measured on admission.

To detect potential cardiac sources of emboli, all patients were examined using a 12-lead electrocardiograph (ECG), by 24-hour ECG monitoring and by transthoracic echocardiography. Causes of embolic cardiac disorders included nonvalvular atrial fibrillation, acute myocardial infarction, previous myocardial infarction with intraventricular thrombus, mitral valve disease, prosthetic cardiac valve, and dilated cardiomyopathy. Potential aortic sources of emboli or patency of the foramen ovale were evaluated in 13 patients by transesophageal echocardiography (TEE). Localized raised lesions in the aorta with a maximal intimal-medial thickness of over 40 mm and an obviously irregular surface or broad acoustic shadow were defined as complicated lesions.

All patients underwent color-flow duplex carotid sonography on the day of admission. Conventional cerebral angiography was performed in 17 patients and the remaining 11 patients underwent MR angiography. Stenoses were graded according to the method used in the North American Symptomatic Carotid Endarterectomy Trial (24).

All patients in whom MESs were detected by TCD were treated immediately with an antiplatelet agent (ticlopidine, 200 mg). A follow-up TCD study was performed after antiplatelet therapy to determine whether the frequency of MESs had changed.

The Mann-Whitney U test was applied to detect differences in age, number of lesions on DWI, NIHSS score, and interval between stroke onset and MR assessment. All other findings were assessed by the χ² test between two groups. Differences were assumed to be significant at P = .05.

Results

In the MES and control groups, one and two MCAs, respectively, ipsilateral to the side of hemiparesis could not be studied by TCD because of a lack of an acoustic window. Therefore, Doppler waves from 19 and 34 MCAs, respectively, were obtained.

Table 1 lists the clinical features and background characteristics of the two groups. The frequency of hypertension, diabetes mellitus, hyperlipidemia, smoking, NIHSS score, blood-coagulation param-

<table>
<thead>
<tr>
<th>Blood-coagulation parameters, mean ± SD</th>
<th>MES Group</th>
<th>Control Group</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>AT III (%)</td>
<td>96.9 ± 19.5</td>
<td>91.2 ± 13.7</td>
<td>NS</td>
</tr>
<tr>
<td>TAT (μg/L)</td>
<td>3.92 ± 5.8</td>
<td>5.3 ± 8.2</td>
<td>NS</td>
</tr>
<tr>
<td>D-dimer (μg/mL)</td>
<td>0.76 ± 0.53</td>
<td>2.06 ± 3.5</td>
<td>NS</td>
</tr>
<tr>
<td>NIHSS score, mean ± SD</td>
<td>8.5 ± 4.7</td>
<td>8.9 ± 5.6</td>
<td>NS</td>
</tr>
<tr>
<td>Interval between stroke onset and MR study (d)</td>
<td>3.3 ± 2.2</td>
<td>3.9 ± 2.5</td>
<td>NS</td>
</tr>
</tbody>
</table>

Note.—MES indicates microembolic signal; AT III, antithrombin III; TAT, thrombin-antithrombin III complex; NS, nonsignificant.
eters, and interval between stroke onset and MR examination did not differ between the two groups.

Significant arterial disease contralateral to the side of hemiparesis was observed in 17 patients (60%): MCA occlusion was found in four patients, M2 occlusion in two, internal carotid artery (ICA) occlusion in one, intra- or extracranial ICA stenosis of 50% or more in four and five, respectively, and ICA stenosis of less than 50% with ulceration in one. Arterial disease was more frequent in the MES group than in the control group ($P = .02$).

Emboligenic cardiac diseases were all NVAF and present in five patients, all of whom were in the control group. The frequency of embolicogenic cardiac diseases between the two groups did not differ. Complicated lesions were revealed by TEE in two patients in the MES group and in two patients in the control group. A patent foramen ovale was observed in one patient in the MES group and in one in the control group.

DWI showed not only hyperintense abnormalities corresponding to the lesions revealed by CT but also other small lesions that were not shown by CT. The small lesions on DWI were detected in the cerebral cortex or subcortex in 18 patients (Figs 1 and 2). T1- and T2-weighted MR images showed the abnormalities in the subcortex corresponding to DWI alterations, but not the small lesions in the cortex. Small hyperintense lesions on DWI were more frequent in the MES group than in the control group ($P < .004$); however, the frequency of medium and large hyperintense lesions did not differ between the two groups (Fig 3).

Table 2 shows the clinical characteristics of all patients in the MES group. The MES counts were recorded not only from 10 MCAs contralateral to the side of hemiparesis but also from three MCAs ipsilateral to the side of hemiparesis. The number of MESs per 30-minute isonation period varied from 1 to 303. Patient 1 had ICA stenosis of less than 50% with ulceration and a patent foramen ovale. Because MESs were documented bilaterally, the source of MESs was more likely to be the patient foramen ovale. Further examinations showed deep venous thrombosis in the lower legs; therefore, we determined that this patient had a paradoxical brain embolism through a patent foramen ovale. Patient 6 had MESs detected from bilateral MCAs and no significant cerebral arterial disease, but had complicated lesions in the aorta, suggesting an aortogenic brain embolism. Patient 9 had ICA stenosis contralateral to the side of hemiparesis and ICA occlusion ipsilateral to the side of hemiparesis. Spotty DWI lesions were observed in both hemispheres. Spotty brain infarctions ipsilateral to the side of hemiparesis may be caused by small emboli from the right ICA to the left MCA through the anterior communicating artery.

In the MES group, five patients had 10 or more MESs per 30 minutes and the remaining five had less than 10 MESs per 30 minutes. The frequency of arterial disease and small or medium or large hyperintense lesions on DWI was not significantly different between these two groups. On the other hand, the NIHSS score was higher in the group with 10 or more MESs per 30 minutes than in the group with less than 10 MESs per 30 minutes ($P = .007$) (Table 3).

In the MES group, no patient was started on antithrombotic therapy before TCD examination and all were treated with an antiplatelet agent immediately after the detection of MESs. Follow-up TCD studies revealed that the frequency of MESs decreased in nine patients 4 to 10 days after antiplatelet therapy was begun. However, transient dysarthria occurred in one patient (patient 3). In the control group, five patients with embolicogenic cardiac disease were treated with heparin (10,000 U/day) before TCD examination, but the remaining 13 patients were started on antithrombotic therapy after TCD.

**Discussion**

In our study, patients with acute ischemic stroke and MESs had a greater prevalence of arterial disease and spotty brain lesions on DWI than did patients with acute ischemic stroke and no MESs. It
Fig 2. Patient 6.

A–D. Axial diffusion-weighted images 5 days after stroke onset show several spotty hyperintense abnormalities (arrows) in both hemispheres. Thirty-eight and thirty-three MESs were recorded with TCD for 30 minutes from the right and left MCAs, respectively. This patient had neither arterial nor cardiac disease but had complicated lesions in the aorta.

Fig 3. Small lesions on DWI are more frequent in the MES group than in the control group ($P < .004$). However, the frequency of medium and large lesions does not differ between the two groups.
TABLE 2: Clinical characteristic of patients with MESs

<table>
<thead>
<tr>
<th>Case No.</th>
<th>Age (y)/Sex</th>
<th>Side of Hemiparesis</th>
<th>NIHSS Score</th>
<th>No. of MESs</th>
<th>Follow-up TCD Examination</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>23/F</td>
<td>L</td>
<td>5</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>2</td>
<td>45/M</td>
<td>R</td>
<td>2</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>3</td>
<td>62/M</td>
<td>R</td>
<td>1</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>4</td>
<td>71/M</td>
<td>R</td>
<td>1</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>5</td>
<td>74/M</td>
<td>R</td>
<td>1</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>6</td>
<td>69/M</td>
<td>R</td>
<td>1</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>7</td>
<td>69/M</td>
<td>R</td>
<td>1</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>8</td>
<td>65/M</td>
<td>R</td>
<td>1</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>9</td>
<td>59/M</td>
<td>R</td>
<td>1</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>10</td>
<td>50/F</td>
<td>L</td>
<td>5</td>
<td>0</td>
<td>0</td>
</tr>
</tbody>
</table>

Note: MES indicates microembolic signal; NIHSS, National Institutes of Health Stroke Scale.

TABLE 3: Demographic features of patients in MES group

<table>
<thead>
<tr>
<th>No. of lesions on DWI, mean ± SD</th>
<th>Patients with Less than 10 MESs (n = 5)</th>
<th>Patients with 10 or More MESs (n = 5)</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Small (&lt;10 mm)</td>
<td>3.4 ± 2.7</td>
<td>5.8 ± 3.6</td>
<td>NS</td>
</tr>
<tr>
<td>Medium (10–30 mm)</td>
<td>0.2 ± 0.4</td>
<td>0.8 ± 0.4</td>
<td>NS</td>
</tr>
<tr>
<td>Large (&gt;30 mm)</td>
<td>0.4 ± 0.5</td>
<td>0.2 ± 0.4</td>
<td>NS</td>
</tr>
<tr>
<td>Arterial diseases, n (%)</td>
<td>5 (100)</td>
<td>4 (80)</td>
<td>NS</td>
</tr>
<tr>
<td>NIHSS score, mean ± SD</td>
<td>5.0 ± 3.6</td>
<td>12.0 ± 2.5</td>
<td>.007</td>
</tr>
</tbody>
</table>

MES indicates microembolic signal; DWI, diffusion-weighted imaging; NS, nonsignificant; NIHSS, National Institutes of Health Stroke Scale.

has been reported that the main sources of microemboli in ICA stenosis are ulceration and thrombus formation of the atheromatous plaque (25). Small emboli may occlude peripheral brain arteries and generate spotty cortical or subcortical infarcts that can be detected by DWI. However, microemboli detected by TCD are asymptomatic and not evident on DWI. Because spotty cortical lesions on DWI are often asymptomatic (26, 27), emboli larger than microemboli detected by TCD may produce asymptomatic spotty lesions on DWI. MESs may indicate the presence of an unstable plaque or platelet aggregation in arterial disease that promotes an artery-to-artery embolism.

In our study, most patients (82%) with acute stroke had no MESs or relatively few; however, five patients (18%) had 10 or more MESs per 30 minutes. We arbitrarily divided the MES group into patients with infrequent MESs (less than 10 per 30 minutes; patients 1–5) and those with frequent MESs (10 or more per 30 minutes; patients 6–10). No significant difference in the frequency of arterial disease or the number of spotty lesions on DWI was observed between these two groups. However, patients with 10 or more MESs per 30 minutes appeared to have more severe neurologic deficits than did those with less than 10 MESs per 30 minutes. We believe that a large number of microembolisms may diffusely damage the brain and cause these neurologic deficits. Therefore, it is worthwhile to distinguish patients with many MESs from those with no or just a few MESs.

Tegeler et al (14) found that patients with MESs had a higher rate of recurrent stroke. In the present study, MESs were associated with small lesions on DWI. All patients with MESs were treated with an antiplatelet agent soon after detection of MESs with TCD. Follow-up TCD studies showed that the frequency of MESs decreased in all patients who had been treated with antiplatelets. Certainly, it remains unclear whether treatment actually reduced MESs or if this was the natural history. Nevertheless, we think that treatment with antiplatelets should be started as soon as possible to prevent
further ischemic stroke after MES is detected by TCD.

Although atherosclerotic lesions of the aorta are recognized as a potential source of emboli (28, 29), a diagnosis of aortogenic brain embolism cannot be readily established. Of the patients whose MESs were recorded from bilateral MCAs, the source was perhaps aortic lesions or cardiac disease. Therefore, if MESs are detected bilaterally in patients with acute stroke and complicated aortic lesions but without another potential source of emboli, a diagnosis of aortogenic embolic stroke should be considered.

Conclusion

Small, often asymptomatic, DWI abnormalities are more frequently seen in patients with acute stroke who have microemboli and large-vessel occlusive disease than in such patients without microemboli. TCD and DWI may provide early clues to the mechanism of stroke during the acute phase.

Acknowledgments

We thank Naoki Yamada (Department of Diagnostic Radiology) for his valuable comments on the MR studies.

References