Comparison of Preperfusion and Postperfusion Magnetic Resonance Angiography in Acute Stroke
Salvador Pedraza, MD; Yolanda Silva, MD; José Méndez, MD; Luis Inaraja, MD, PhD; Joana Vera, MD; Joaquín Serena, MD, PhD; Antoni Dávalos, MD, PhD

Background and Purpose—The multimodal magnetic resonance imaging study in acute stroke includes perfusion-weighted imaging (PWI) after administration of contrast and magnetic resonance angiography (MRA). However, MRA may overestimate the degree of vessel obstruction caused by limitations to detect low flow states. Our aim was to determine the usefulness of a new fast imaging protocol combining classical MRA, PWI, and postperfusion MRA to improve the diagnostic management in acute ischemic stroke.

Methods—We studied 31 patients with a middle cerebral artery (MCA) infarction within the first 12 hours from the onset of symptoms. All patients had an MCA stenosis or occlusion. The study protocol included a preperfusion MRA and a postperfusion MRA. Modified thrombolysis in myocardial infarction (TIMI) classification was used to assess the patency of vessels.

Results—In 17 patients (group A, 55%), preperfusion MRA and postperfusion MRA accorded in the estimation of vascular status, whereas in 14 patients (group B, 45%) postperfusion MRA showed a better vascular flow than preperfusion MRA. The improvement in the depiction of flow was from a complete occlusion (TIMI I) to a partial occlusion (TIMI II) in 9 patients and from TIMI II to normal patency (TIMI III) in 5 patients. Thirty-six percent of the patients with suspected internal carotid artery occlusion in the preperfusion MRA showed flow in the intracranial internal carotid artery in the postperfusion MRA.

Conclusions—Postperfusion contrast-enhanced MRA can demonstrate arterial segments with low flow and avoid overestimation of vascular obstruction. (Stroke. 2004;35:2105-2110.)

Key Words: angiography • contrast media • magnetic resonance • myocardial infarction

The vascular assessment in the acute ischemic stroke can reveal the presence of a vascular occlusion or a normal patency caused by early recanalization of the vessel obstruction. The distinction is important because it has therapeutic and prognostic consequences. It has been reported that patients with complete occlusion of the distal internal carotid artery are unlikely to recanalize with intravenous thrombolytic therapy. In addition, patients with distal occlusion of internal carotid artery have worse prognosis compared with those with proximal middle cerebral artery (MCA) occlusion. The prognosis is especially poor when leptomeningeal collaterals are insufficient.

The optimal magnetic resonance imaging (MRI) protocol in acute stroke includes diffusion-weighted imaging (DWI) to detect acute ischemic lesion, fluid attenuation inversion recovery to assess chronic lesions, magnetic resonance angiography (MRA) to detect vascular occlusion, and perfusion-weighted imaging (PWI) to estimate brain perfusion. The time of flight (TOF) is the most common MRA technique. However, TOF overestimates the severity of the arterial stenosis caused by the saturation of slow flow and therefore can produce a false diagnosis of vascular occlusion. This diagnostic uncertainty impedes the correct assessment of the hemodynamic status and falsely predicts a worse prognosis. Previous studies have shown that the use of MRA with contrast is an approach to clarify this uncertainty. The contrast shortens the T1 of blood, decreasing the saturation effects and improving the depiction of small vessels and blood–tissue differentiation, particularly in situations of slow flow. However, there are only a few studies on the value of MRA after contrast administration in the setting of acute stroke.

Our aim was to determine the usefulness of a new and fast imaging protocol combining preperfusion MRA and postperfusion MRA to improve the diagnostic management of acute ischemic stroke.

Materials and Methods
This is a secondary study of patients with acute hemispheric ischemic stroke admitted consecutively within the first 12 hours after

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the onset of symptoms between July 2001 and December 2002. The primary aim of this still-ongoing prospective study is to investigate whether the molecular factors associated with early neurological deterioration correlate to the evolution of the DWI, PWI, and MRA abnormalities in the acute phase of territorial infarctions. Clinical exclusion criteria were coma on admission, pure lacunar syndromes, transient ischemic attack, previous cerebral infarction impeding the clinical and neuroradiological evaluation, and chronic severe diseases. For the purpose of this investigation, only patients with an acute MCA infarction associated with a vascular stenosis or occlusion of the MCA in the preperfusion MRA were studied. The ethics committee approved the study and written informed consent was obtained from all the patients or relatives before imaging.

MRI Protocol
All patients had a MRI study on admission within the first 12 hours after stroke onset. MR images were obtained on a 1.5-T system (Intera, Philips) with echoplanar capabilities of 25-mTeslas/m and 300- to 350-μsec rise times. The MRI protocol included DWI, fluid attenuation inversion recovery, preperfusion MRA, PWI with gradient-echo echo-planar technique, and postperfusion MRA sequence. The diffusion sequence was obtained with a single-shot spin-echo planar pulse with a diffusion gradient b-value of 0 and 1000 s/mm² along 3 axes. The other parameters were 20 slices, 7-mm slice thickness, 0 gap, 134 ms echo time (TE), 6000 ms repetition time (TR), 67 (epi factor), and 36 seconds of duration. Abnormalities were analyzed in the trace image to avoid anisotropy. The PWI sequence was acquired after administration of 0.2 mL per kilogram of gadolinium DTPA with a bolus technique (5 mL/sec). The parameters were 260, 30 (TR/TE), 12 slices of 10-mm slice thickness, 0 gap, 60 dynamic scans, and 1 minute and 4 seconds of duration. The volume of hypoperfused tissue was measured in the mean transit time map using a threshold of 4 seconds. A secondary analysis was performed using a threshold of 6 seconds.

Preperfusion MRA was performed before PWI and postperfusion MRA was performed immediately after the end of the PWI without the administration of a supplementary dose of contrast. That means that postperfusion MRA was performed 1 minute and 4 seconds after the administration of contrast. While the post-perfusion MRA was being performed, the quality of the PWI maps were checked. Preperfusion and postperfusion MRA were performed using conventional parameters when we did not have any time pressure to administer specific therapies, and when patients were still. We used this high-quality TOF sequence to examine the Circle of Willis and cranial and cervical ICA to rule out the presence of ICA occlusion. The imaging parameters were 18/6/9/20 (TR/TE/flip angle), 230 field of vision, 256×256 matrix, 1 slab with 200 slices, 0.6-mm slice thickness, and tilted optimized nonsaturating excitation pulse. The acquisition time was 6 minutes and 14 seconds. In candidates for thrombolytic therapy and in restless patients, preperfusion and postperfusion MRA were performed using a faster TOF sequence. The imaging parameters were: 19/6/9/20 (TR/TE/flip angle), 150 field of view, 256×256 matrix, 1 slab with 70 slices, 0.8-mm slice thickness, and tilted optimized nonsaturating excitation pulse. The acquisition time was 1 minute and 41 seconds. The other parameters were 20 slices, 7-mm slice thickness, and tilted optimized nonsaturating excitation pulse. The acquisition time was 1 minute and 4 seconds. In candidates for thrombolytic therapy and in restless patients, preperfusion and postperfusion MRA were performed using a faster TOF sequence. The imaging parameters were: 19/6/9/20 (TR/TE/flip angle), 150 field of view, 256×256 matrix, 1 slab with 70 slices, 0.8-mm slice thickness, and tilted optimized nonsaturating excitation pulse.
postperfusion MRA than in preperfusion MRA (Figure 2). Group A included 4 patients with unchanged MCA occlusion but with a better depiction of flow in ICA (2 patients) or posterior cerebral artery (PCA) (2 patients) after contrast. Seventeen patients were studied with the shorter MRA technique, 10 in group A and 7 in group B (P=NS). In the group of 10 patients in whom the longer and the shorter preperfusion MRA techniques were sequentially used, we did not find any difference between them on TIMI classification. Postperfusion MRA disclosed prominent venous signal in all patients; however, in 90% of them MIP improved qualitatively the arterial detail over the venous background.

Six patients were treated with intravenous rt-PA before the MRI study, 1 in group A and 5 in group B (P=0.08), and 6 patients were treated with intravenous rt-PA after the MRI study, 3 in group A and 3 in group B. One patient was treated with desmoteplase/placebo after the MRI study in group B. There were no differences in age, sex, time from symptoms onset to MRI, initial NIHSS, PWI lesion volume, and in the frequency of PWI/DWI mismatch between the 2 groups (Table 2). Patients in group B had, however, a nonsignificant lower proportion of MCA occlusion (P=0.06) and smaller DWI lesion volume (P=0.16). Combined MCA and intracranial ICA occlusion in preperfusion MRA was found in 9 patients (53%) of group A and in 5 patients (36%) of group B (P=NS). Duplex sonography showed ICA cervical occlusion in all cases. However, postperfusion MRA showed a normal flow in the intracranial ICA with persistent MCA occlusion in 2 patients of group A and a normal flow in the intracranial ICA and MCA in 3 patients of group B. Consequently, 36% (5/14) of the patients with suspected intracranial ICA occlusion in the preperfusion MRA showed flow in

Figure 1. Post-perfusion MRA demonstrates the same degree of arterial flow. A, Preperfusion MRA demonstrates normal flow in proximal left MCA with absence of flow in intracranial ICA and in distal left MCA. B, Postperfusion MRA confirms the occlusion of intracranial ICA and distal MCA arteries on the left side.

Figure 2. Postperfusion MRA demonstrates a better flow. A, Preperfusion MRA shows the absence of flow in left proximal MCA (TIMI I). B, Postperfusion MRA shows correct distal flow in left MCA with moderate proximal stenosis (TIMI II). This patient did not receive reperfusion therapy.
the postperfusion MRA (Figure 3). In patients without intracranial ICA occlusion in the postperfusion MRA, PWI lesion volume was also comparable between the 2 groups (144 mL [5, 174]) (median [lower quartile, upper quartile]) in group A and 111 mL (85, 147) in group B, \( P = 0.81 \). When the PWI lesion volume was calculated using the threshold of 6 seconds, we obtained similar results.

**Discussion**

This study shows that a new fast combined protocol with postperfusion MRA is useful in the diagnosis of vessel patency in acute ischemic stroke. Nearly half of the patients studied within 12 hours with an MCA stenosis or occlusion in the preperfusion MRA showed a better depiction of MCA flow in the postperfusion MRA. To our knowledge, there are no previous references about the use of MRA after PWI. We have not found clinical or neuroimaging factors associated with a lack of agreement between preperfusion and postperfusion MRA, except a nonsignificant higher proportion of rt-PA treatment before the MRI study in group B and a lower initial DWI volume in group B. Furthermore, PWI lesion volume was comparable between groups, either in the total number of patients or in the subgroup without true ICA occlusions. Although we have studied a limited number of patients, our findings support the notion that integrated information from preperfusion MRA and PWI is not suitable to rule out or suspect an overestimation of vascular obstruction in patients with low flow.

Postperfusion MRA may have therapeutic consequences.\(^1\)\(^-\)\(^7\) In this study, 36% of patients showing an intracranial ICA occlusion in the preperfusion MRA had an overestimation of vascular obstruction, because the intracranial segment of the ICA was patent in the postperfusion study. After rt-PA treatment, ICA occlusion has a lower proportion of recanalization (31%) compared with MCA occlusion (88%).\(^5\) In this context, the occlusion of the ICA without MCA patency has been proposed as a relative exclusion criteria for thrombolysis in acute ischemic stroke.\(^1\)\(^-\)\(^2\) Following this criteria, preperfusion MRA findings might erroneously prevent the administration of intravenous rt-PA in more than one third of patients with suspected intracranial ICA occlusion, which, in this study, represented 10% of the studied population. Further studies must address the issue of therapeutic implications of postperfusion MRA.

Previous studies performed in subacute stroke and not using PWI have shown the usefulness of contrast administration to improve the depiction of intracranial vessels.\(^10\) Yano et al found that MRA with contrast improved the detection of distal MCA branches in 61% of the patients, and that 30% of the branches of MCA became visible only after contrast.\(^11\) Other reports have shown that after the administration of contrast, the depiction of flow in MRA changes from apparent occlusion to normal pattern in 17% of patients and in 7% of vessels.\(^5\)\(^,\)\(^11\)\(^,\)\(^13\) The present study further demonstrates the usefulness of a new acute imaging protocol, performing MRA after PWI without the administration of a supplementary does of contrast. The proposed protocol does not increase the cost or the length of the study, because the standard does of contrast is used and the postperfusion MRA is performed while the qualities of the perfusion maps are checked.

**TABLE 2. Clinical and MRI Findings in the 2 Groups at Baseline**

<table>
<thead>
<tr>
<th></th>
<th>Group A (n=17)</th>
<th>Group B (n=14)</th>
<th>P</th>
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<tbody>
<tr>
<td>Age</td>
<td>67.6±12.0</td>
<td>67.2±12.0</td>
<td>0.924</td>
</tr>
<tr>
<td>Sex, male</td>
<td>47%</td>
<td>57%</td>
<td>0.72</td>
</tr>
<tr>
<td>Time from symptom onset</td>
<td>240 (186, 310)</td>
<td>238 (165, 344)</td>
<td>0.95</td>
</tr>
<tr>
<td>Initial NIHSS score</td>
<td>17 (10, 20)</td>
<td>15 (7, 19)</td>
<td>0.40</td>
</tr>
<tr>
<td>Initial volume in DWI, mL</td>
<td>22 (5, 124)</td>
<td>8 (6, 25)</td>
<td>0.16</td>
</tr>
<tr>
<td>Initial volume in PWI, mL</td>
<td>164 (23, 180)</td>
<td>100 (33, 135)</td>
<td>0.26</td>
</tr>
<tr>
<td>Thrombolysis before MRI</td>
<td>1 (7.7%)</td>
<td>5 (35.7%)</td>
<td>0.08</td>
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</table>

**MCA Occlusion**

<table>
<thead>
<tr>
<th></th>
<th>Group A</th>
<th>Group B</th>
<th>P</th>
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<tbody>
<tr>
<td>Preperfusion MRA</td>
<td>16</td>
<td>9</td>
<td></td>
</tr>
<tr>
<td>Postperfusion MRA</td>
<td>16</td>
<td>0</td>
<td></td>
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**MCA Stenosis**

<table>
<thead>
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<th>Group A</th>
<th>Group B</th>
<th>P</th>
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<tbody>
<tr>
<td>Preperfusion MRA</td>
<td>1</td>
<td>5</td>
<td></td>
</tr>
<tr>
<td>Postperfusion MRA</td>
<td>1</td>
<td>9</td>
<td></td>
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**ICA Occlusion**

<table>
<thead>
<tr>
<th></th>
<th>Group A</th>
<th>Group B</th>
<th>P</th>
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<tbody>
<tr>
<td>Preperfusion MRA</td>
<td>9</td>
<td>5</td>
<td></td>
</tr>
<tr>
<td>Postperfusion MRA</td>
<td>7</td>
<td>2</td>
<td></td>
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than preperfusion MRA, so postperfusion MRA should substitute preperfusion MRA in future studies to reduce the time consumption of the MR study.

Postperfusion MRA has, however, some limitations. The overlap of the venous enhanced vessels (cavernous sinus or transverse sinus) over the arterial vessels make the assessment of the arterial branches difficult. The venous enhancement is related to the short time (8 seconds) between the arterial and venous phases in the brain, and it depends on the contrast dose. In this context, the use of lower doses (5 to 10 mL) of contrast and the suppression of enhanced structures in the MIP of postperfusion MRA may help to obtain better images. The 3-step strategy we used is rather complex, so preperfusion MRA and the analysis of source images might be avoided to simplify the process. Although contrast-enhanced vessels should be cautiously interpreted, our results agree with those from other authors who found that venous enhancement has no effect on the assessment of most vessels. A further limitation of postperfusion MRA is related to the difficult differentiation between 2 situations: the presence of distal flow beyond a proximal vascular stenosis or the presence of a vascular occlusion with distal flow caused by the collateral circulation. Nevertheless, whatever the mechanism responsible for the distal flow, postperfusion MRA can show the patency of distal vessels not seen in the preperfusion study.

The present study has other limitations. First, we used 2 different types of TOF sequences depending on the collaboration of the patient at the acute setting and on the time pressure. However, the comparison of both techniques sequentially performed in a small group of patients did not show differences in the estimation of TIMI degree, so we think this fact did not influence our findings. Second, we cannot completely rule out an overestimation of false MCA occlusions in this series because of the more frequent use of intravenous thrombolysis before MRA in group B. Although the time between preperfusion and postperfusion MRA was too short to suspect a recanalization between the 2 examinations, we cannot rule out in some patients with overestimated MCA occlusions in the preperfusion MRA that the better flow in the postperfusion MRA could be related to the rt-PA treatment. A further point of interest is that postperfusion MRA results cannot be fully translated into meaningful information on blood supply of the tissue, because PWI lesion volume was comparable in patients with and without overestimation of ICA and/or MCA obstructions. This finding indicates that most overestimated obstructions reflect an extremely low flow (in many instances caused by ICA proximal occlusions) insufficient to provide a significantly better tissue perfusion status in comparison with true arterial obstructions.

The data presented show that precontrast MRA overestimates the degree of vascular obstruction. Postperfusion contrast-enhanced MRA can demonstrate arterial segments with low flow and avoid overestimation of vascular occlusion. Although this information may be crucial for patients eligible for reperfusion therapies, a PWI study is necessary to obtain comprehensive information on tissue perfusion.

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