Magnetic resonance angiography (MRA) allows visualisation of both extra- and intra-cranial vessels in a fast and non-invasive manner and can play an important role in clinical decision-making. Assessment of the intra-cranial vessels has led to multiple improvements of the technique with a significant increase in the accuracy of intra-cranial MRA. Prior research has shown benefit of careful review of the 3D TOF source images.

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along with maximum intensity projections (MIP) and collapsed MIPs.\textsuperscript{1,5} Source images are particularly helpful for the assessment of intra-cranial stenoses. While most previous studies have concentrated on the proximal intracranial vessels, a few studies have reported difficulty in evaluating peripheral vessel branches.\textsuperscript{1,2} Contrast-enhanced MRA\textsuperscript{4,5} improves conspicuity of vessels of small caliber and slow blood flow.

In the setting of acute ischemic stroke clinicians seek information about the status of intracranial vessels. Hyperacute neurovascular imaging is challenging for number of reasons: (i) information needs to be obtained quickly; (ii) patients suffering severe disabling stroke do not tolerate prolonged imaging time and; (iii) relatively short acquisition time limits achievable image quality.

The purpose of this study was to investigate the reliability and accuracy of intracranial MRA of both proximal and distal vessels compared to conventional angiography in the setting of acute ischemic stroke (AIS) and transient ischemic attack (TIA).

METHODS

Patients

We reviewed our research imaging database to identify patients who had undergone MR angiography. All patients with AIS or TIA, who for clinical reasons, underwent both 3 T MRI and conventional angiography within 48 hours from one another were recruited. Written informed consent to the use of the imaging in research was provided by all patients or their surrogate before MR imaging and conventional angiography.

MR Imaging Protocol

Magnetic resonance studies were performed on a 3.0 Tesla MRI system (Signa; GE Medical Systems, Waukesha, WI) equipped with high speed gradients (40-mT/m peak strength, 184-µs rise time) using a standard quadrature head coil. The acute stroke protocol included standard anatomic imaging, diffusion-weighted imaging (DWI), three dimensional time-of-flight (3D TOF) MR angiography, and dynamic susceptibility contrast perfusion-weighted imaging (DSCPWI).\textsuperscript{5} Time-of-flight angiograms were collected before (non-enhanced) and after (enhanced) the perfusion imaging contrast injection (20 ml of MR contrast – Magnevist, Berlex, Wayne, NJ - at 5 ml/s).

The imaging parameters for the conventional non-enhanced two-slab 3D TOF sequence were 24/3.3/15° (TR/TE/flip angle), acquisition bandwidth of 12.5 kHz, and a 240 x 144 x 46-mm acquired volume with a 256 x 192 x 42 acquisition matrix, which was then reconstructed to a 512 x 512 x 84 matrix. Total acquisition time was 2 minutes 48 seconds. Axial slabs were prescribed from skull base to the circle of Willis. Post-contrast 3D TOF MR angiography focused on visualizing the intracranial circulation and therefore the non-enhanced technique was modified. First, an inclined slab was used to image the intracranial vasculature including the distal ICA and the intracranial circulation and therefore the non-enhanced technique was modified. Second, to maximize vessel signal intensity on post-contrast images, the TR and flip angle were increased. Post-contrast MR angiographic parameters were as follows: 32/3.4/35° (TR/TE/flip angle) acquisition bandwidth of 12.5 kHz, and a 240 x 144 x 36-mm acquired volume with a 256 x 192 x 40 acquisition matrix, which was then reconstructed to a 512 x 512 x 80 matrix. Total acquisition time was 4 minutes 9 seconds. Using the non-enhanced MRA source images at the circle of Willis, maximum intensity projection (MIP) images were generated at multiple projection angles every 12.9° about the superior/inferior (S/I) axis. Axial collapsed MIP images of both non-enhanced and enhanced MRA volumes were also generated.

Twenty-four patients had both non-enhanced and enhanced MRA, four patients had only non-enhanced MRA, and one patient had only enhanced MRA. Four enhanced MRA studies were excluded because of incorrect slab position.

Angiography protocol

Conventional angiograms were performed in a standard manner starting with an arch aortogram followed by selective injections of the carotid and/or vertebral arteries. Biplane digital subtraction angiography was performed (LCA/LP Advantx Angiography unit, General Electric). Intracranial vessels were imaged with a minimum field of view of 15 cm in the anteroposterior plane and 11.4 cm in the lateral plane at a magnification factor of 2.6. A high resolution (1024 x 1024) matrix was used.

MRA review

Film copies of axial collapsed MIP images and source images for both non-enhanced and enhanced 3D TOF, as well as non-enhanced MIP images were prepared for review. Five readers – two neuroradiologists, one stroke neurologist, and two stroke fellows blind to conventional angiography results and all clinical information except symptom side scored all MRA films. Each observer was given both non-enhanced and enhanced films at the same time, although the non-enhanced were scored before the enhanced films. The following intra-cranial vessels and vessel segments: internal carotid artery (ICA); M1 middle cerebral artery (MCA); M2 MCA; M3 MCA; anterior cerebral artery (ACA); vertebral arteries; basilar artery and posterior cerebral artery (PCA) were scored as patent or occluded based on signal intensity. A vessel or vessel segment was considered occluded if there was absent flow on all of MRA source images, MIPs and collapsed MIPs, within each modality (with or without MR contrast).

Angiography review

One experienced neuroradiologist blinded to MRA findings reviewed all angiograms. He was given the information about symptom side, and also which procedure was performed first: MRA or conventional angiography. He was asked to score only that part of the conventional angiogram, which was closest in time to the MRA in order to minimize the likelihood of vascular changes that may have occurred in between. Identical to the MRA scoring method, the analogous vessel segments were rated as patent or occluded. This statement was considered a gold standard.

Data analysis

Kappa statistics were used to assess inter-observer reliability. Agreement was considered to be slight (κ=0.0-0.2), fair (κ=0.21-0.4), moderate (κ=0.41-0.60), substantial (κ=0.61-0.80), or
almost perfect (κ = 0.81-1.00). Contingency tables were used to assess the accuracy of non-enhanced and enhanced 3D TOF MRA compared to conventional angiography. Magnetic resonance summary scores used for this comparison were the median of 5 raters which is equivalent to a 3 of 5 consensus. We did not include the posterior circulation data in this analysis because of a paucity of occlusions in this territory.

RESULTS

Twenty-nine patients (7 females, 22 males), aged 18 to 83 years (median 53 years) with acute stroke (26 cases) or TIA (3 cases) were enrolled. National Institutes of Health Stroke Scale (NIHSS) scores were obtained at patient presentation and prior to the MR exam by qualified examiners. Median NIHSS was 14 (range 0 to 25). Fourteen patients received thrombolysis during MRA or angiography. Angiography preceded MRA in 15 cases. Thirteen patients (45%) had their two studies within 3 hours one from another. Median time between non-enhanced and enhanced MRA was 22 min. Twenty-six occlusions were diagnosed during conventional angiography among 252 intra-cranial vessels assessed: 25 in the anterior circulation (ICA, M1-MCA 10, M2-MCA 4, M3-MCA 3) and one in posterior circulation (PCA 1).

Overall agreement among 5 raters was moderate for both non-enhanced (κ = 0.50 CI95 0.47-0.53) and gadolinium-enhanced (κ = 0.46 CI95 0.37-0.58) images. A gradient of agreement existed such that agreement for the proximal vessels was excellent, while for the distal vessels, agreement was poorer (Table 1). The overall accuracy of non-enhanced MRA was 84.5% (CI95 0.79-0.89) and for enhanced MRA 73.2% (CI95 0.65-0.80). The non-enhanced MRA showed sensitivity of 84.2% (CI95 60.4-96.6) and specificity of 84.6% (CI95 78.6-89.4). The enhanced MRA showed sensitivity of 69.2 (CI95 38.6-90.9) and specificity of 73.6 (CI95 65.5-80.7). This was not substantially changed by inclusion of dynamic cases where reperfusion occurred due to therapeutic intervention.

In five cases the vessel status was dynamic as re-perfusion was achieved between angiography and the MR imaging as the result of interventional therapy. Consequently, there was clear discrepancy in the scoring of the vessel status by MRA and conventional angiography. After excluding those cases the accuracy of MRA was very good including distal vessels (Table 2).

Although agreement among raters was not improved with contrast enhancement, we found a positive impact of contrast administration on vessel assessment. In the ICA, use of contrast non-significantly raised sensitivity to 100%, while in distal MCA (M2/M3 segments) a trend to improved specificity was observed (p=0.087) (Figure 1). Similarly, level of training was relevant. The overall specificity of the neurologists’ assessments of the post-contrast images was 0.77 (CI95 0.58-0.90), while the analogous result of the neuroradiologists’ was 1.00 (CI95 0.88-1.00). Overall, neurologists’ accuracy ranged from 77-81% for pre- or post-contrast MRI with or without the dynamic cases. This compares to 86-91% for neuroradiologists. Accuracy was always equivalent or non-significantly higher for the contrast-enhanced assessment.

DISCUSSION

Thrombolytic therapy remains the only proven therapeutic intervention with a potential to restore perfusion and reverse symptoms in patients suffering AIS. Imaging modalities (MR and CT) that are capable of rapidly identifying vessel occlusion causing symptoms of stroke, are indispensable in this scenario. While occlusion identification substantiates therapeutic efforts, any imaging protocol for AIS or TIA has specific restrictions and represents a trade-off between patient comfort, technical optimization and diagnostic accuracy.

The reliability of MRA has improved with technical innovation. Korogi et al2 noted in 1994 that each of the observers, participating in their study of unenhanced TOF MR angiography, interpreted several ICA occlusions or stenoses of greater than 50% as normal. Consequently the authors found MRA unreliable for the assessment of severe ICA stenoses. The addition of 3D TOF source images resulted in increased sensitivity to 100% for moderately and severely stenosed, as well as occluded vessels.7 Heiserman et al2 achieved good results 3D TOF source images were combined with projections and collapsed MIP images. Thus, the interpretation protocol must utilize all available images including MIP and source images to achieve the best diagnostic result.

Robinson et al8 described application of the multiple overlapping thin slab acquisition (MOTSA) in the evaluation of carotid artery. Multiple overlapping thin slab acquisition combines the advantages of two- and three-dimensional TOF

Table 1: Inter-rater agreement for pre- and post contrast MRA

<table>
<thead>
<tr>
<th>VESSEL</th>
<th>PRE-CONTRAST κ(95% LCI)</th>
<th>POST-CONTRAST κ(95% LCI)</th>
</tr>
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<tbody>
<tr>
<td>ICA</td>
<td>0.92 (0.81)</td>
<td>0.7 (0.58)</td>
</tr>
<tr>
<td>MCA-M1</td>
<td>0.53 (0.44)</td>
<td>0.55 (0.45)</td>
</tr>
<tr>
<td>MCA-M2/M3</td>
<td>0.17 (0.09)</td>
<td>0.18 (0.08)</td>
</tr>
</tbody>
</table>

Table 2: Sensitivity and specificity of pre-contrast MRA after removing “Dynamic” cases

<table>
<thead>
<tr>
<th>VESSEL</th>
<th>Sensitivity</th>
<th>Specificity</th>
</tr>
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<tbody>
<tr>
<td>ICA</td>
<td>0.86 (0.42-1)</td>
<td>0.94 (0.73-1)</td>
</tr>
<tr>
<td>Post-gadolinium</td>
<td>1 (0.75-1)</td>
<td>0.93 (0.66-1)</td>
</tr>
<tr>
<td>M1</td>
<td>0.75 (0.35-0.97)</td>
<td>0.83 (0.69-0.93)</td>
</tr>
<tr>
<td>Post-gadolinium</td>
<td>0.67 (0.22-0.96)</td>
<td>0.79 (0.59-0.92)</td>
</tr>
<tr>
<td>M2/M3</td>
<td>0.80 (0.28-0.99)</td>
<td>0.69 (0.53-0.82)</td>
</tr>
<tr>
<td>Post-gadolinium</td>
<td>0.50 (0.07-0.93)</td>
<td>0.87 (0.69-0.96)</td>
</tr>
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</table>
techniques: noise reduction and thin section detail with improved blood flow signal. However, early MR angiograms obtained using a MOTSA technique and postprocessed by the MIP algorithm displayed signal intensity variation, called venetian blind artifact. This artifact can be overcome by optimizing image acquisition parameters (flip angle, TR, and the slab excitation fraction), as well as post-processing algorithm, which results in a reduction of slab boundary artifact. A similar technique was used in this study.

Improved MRA techniques have been reported using higher matrix, overlapping slabs, a variable flip angle excitation (tilted optimized nonsaturating excitation – TONE), and magnetization transfer suppression (MTS), but this technique comes at the expense of a 60 – 90 min examination time; such times are untenable in acute stroke. Even these significant technical improvements did not eliminate artifacts such as flow saturation, that result in relatively common false positive diagnoses of vessel stenoses, especially at bifurcations.

The advantages of a 3D TOF technique include signal-to-noise reduction and better delineation of small vessels. However, 3D-TOF is more sensitive to saturation resulting in poorer vessel definition. Contrast enhanced intracranial angiography has been explored and may be superior to non-enhanced studies. Loss of signal due to saturation of slowly flowing blood can be compensated by the T1 reduction achieved with paramagnetic extracellular contrast agents, such as gadopentetate dimeglumine. Other techniques suppressing soft tissue background such as magnetization transfer, optimization of imaging parameters, including TR, TE, and flip angle, have been explored and have been shown to improve imaging of the intracranial vessels.

Not all of the above techniques can be implemented at higher fields. In order to optimize our MRA protocol we have modified imaging parameters by using variable flip angle, increased TR and TE for post-contrast MRA (see Methods). For acquisition of non-enhanced 3D-TOF we have used 2 overlapping slabs, while for the enhanced MRA one inclined slab. Some techniques, like magnetization transfer are more difficult at higher fields in order to supress the soft tissue background.

We have determined the performance of MRA in evaluation of the intracranial vasculature. Exact grading of the possible vessel stenosis was not of paramount importance in acute stroke; therefore, we focused on recognising occlusion. Although image resolution remains better with conventional angiography, the raters were able to discern smaller vessel branches using MRA with satisfactory accuracy.

Similar to the other groups we have achieved moderate-to-high sensitivity and specificity in the assessment of the ICA. Analogous results were achieved in evaluation of proximal MCA branches (M1 proximal and distal). Our group has previously explored the issue of contrast influence on evaluation of distal intracranial branches. A small proportion (3.3%) of assessed vessels were less visible after contrast administration. This predominantly occurred in the P1 and P2 segments, which were

Figure: 3D TOF images of a 33-year-old male patient with left M2-MCA occlusion showing the MRA source image, maximum intensity projection (MIP), pre-contrast collapsed MIP, post-contrast collapsed MIP.
not in scope of analysis of the present study, but was also observed in the anterior circulation. Explanation of this phenomenon remains unclear.

We have noted, that less experienced raters tended to overcall the occlusions, while experts (neuroradiologists) were more specific in their readings. Other authors have similarly raised the importance of training and experience in order to evaluate MRA accurately. Non-radiologists may not always be familiar with the technical limitations such as overestimation of the vessel stenosis using MIP algorithm. In our study it was important to pay attention to the boundaries of slab position: vessels not included in the slab appeared occluded on both source and postprocessed images.

There were limitations in our study. The major one was use of thrombolytic in 48% of cases, however, not using tPA for eligible patients would be unethical. In at least five cases, vessel status was clearly different between conventional angiography and MRA. Analysis has been conducted with and without these cases. It is possible that in a few cases vessel status changed even between pre- and post-contrast MRA, since the average time between the two was 20 min. An additional drawback of the study was the lack or exclusion of contrast enhanced studies in eight cases. This obstructs full analysis of the impact of contrast administration on MRA.

The MRA is a reliable screening tool, when reviewed by expert readers familiar with MRI technique and possible artifacts, for the evaluation of intracranial vessels. However, angiography remains the gold standard for assessment of intracranial vessels and allows direct intra-arterial intervention. In the setting of acute stroke MRA is not evaluated in isolation. Other MRI sequences like DWI, FLAIR, and particularly PWI provide valuable information. Thus MRI provides comprehensive, non-invasive assessment of both the vessel and the status of ischemic tissue. Such information is invaluable in diagnosis and planning patient management.

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REFERENCES