

Generic Contrast Agents

Our portfolio is growing to serve you better. Now you have a *choice*.



[VIEW CATALOG](#)

AJNR

Contrast-Enhanced 3D MR Angiography of the Carotid Artery: Comparison with Conventional Digital Subtraction Angiography

Luca Remonda, Pascal Senn, Alain Barth, Marcel Arnold, Karl-Olof Lövblad and Gerhard Schroth

This information is current as of May 12, 2025.

AJNR Am J Neuroradiol 2002, 23 (2) 213-219
<http://www.ajnr.org/content/23/2/213>

Contrast-Enhanced 3D MR Angiography of the Carotid Artery: Comparison with Conventional Digital Subtraction Angiography

Luca Remonda, Pascal Senn, Alain Barth, Marcel Arnold, Karl-Olof Lövblad, and
Gerhard Schroth

BACKGROUND AND PURPOSE: Since 1996, several preliminary studies have shown the usefulness of contrast material-enhanced MR angiography for imaging supraaortic vessels. The aim of this study was to compare the accuracy of contrast-enhanced 3D MR angiography with that of digital subtraction angiography (DSA) in the evaluation of carotid artery stenosis.

METHODS: A blinded comparison of first-pass contrast-enhanced MR angiography with conventional DSA was performed in 120 patients (240 arteries). MR angiography was performed with a 1.5-T magnet with gradient overdrive equipment, by using a coronal radiofrequency-spoiled 3D fast low-angle-shot sequence after the intravenous injection of gadodiamide. The guidelines of the North American Symptomatic Carotid Endarterectomy Trial for measuring stenosis of the internal carotid artery were applied on maximum intensity projection (MIP) images and conventional catheter angiograms.

RESULTS: Grading of stenoses on MR angiograms agreed with grading of stenoses on DSA images in 89% of arteries. In the severe stenosis group (70–99%), agreement was 93%. All internal carotid occlusions ($n = 28$) and seven of nine pseudo-occlusions were accurately detected with contrast-enhanced MR angiography. The correlation between MR angiography and DSA for determination of minimal, moderate, and severe stenoses and occlusion was statistically significant ($r = 0.91$, $P < .001$).

CONCLUSIONS: This investigation with a large number of patients confirms that contrast-enhanced MR angiography could become a diagnostic alternative to DSA in the treatment of patients with carotid artery disease.

Several randomized controlled clinical trials, such as the North American Symptomatic Carotid Endarterectomy Trial (NASCET) and the European Carotid Surgery Trial proved the benefit of endarterectomy in patients with symptomatic, severe ($\geq 70\%$) carotid stenosis (1–4). The indication for carotid endarterectomy has been extended also to patients with asymptomatic carotid artery stenosis ($\geq 60\%$), as presented in the Asymptomatic Carotid Atherosclerosis Study (ACAS) (5). Final NASCET results demonstrated a surgical benefit for selected patients with stenosis as low as 50% (6). These studies gave rise to discussion

and kindled the interest in and necessity for accurate measurements of carotid stenosis. In these studies, digital subtraction angiography (DSA) was the accepted criterion standard to determine the degree of carotid stenosis with respect to surgical intervention.

Since 1996, several preliminary studies have shown the usefulness of contrast material-enhanced MR angiography for imaging supraaortic vessels (7–38). Developments in hardware and software have overcome initial limitations and resulted in high-quality images of the carotid arteries comparable to those with DSA. Different strategies for sampling k space can be used for acquisition of 3D MR angiographic sequences according to the clinical application. The most conventional protocols include a large imaging volume, a spatial resolution adequate for carotid assessment, and an acquisition time of approximately 30 seconds. Other sequences improve the spatial resolution of the images by using a 512×512 matrix with submillimetric voxels. Another strategy is to improve the temporal resolution by using an acquisition time of 5–10 seconds, to obviate bolus timing and provide some hemodynamic data.

Received July 24, 2001; accepted after revision November 12.

From the Departments of Neuroradiology (L.R., K.-O.L., G.S.), Neurosurgery (P.S., A.B.), and Neurology (M.A.), University of Bern, Switzerland.

Presented at the 39th annual meeting of the American Society of Neuroradiology, Boston, April 2001.

Address reprint requests to Luca Remonda, MD, University of Bern, Inselspital, Department of Neuroradiology, Freiburgstrasse 10, CH-3010 Bern, Switzerland.

The aim of this study was to compare an ultrafast time-resolved contrast-enhanced MR angiographic technique with DSA for use in detecting stenosis at the carotid bifurcation.

Methods

Patients

We examined 120 consecutive patients (30 female and 90 male patients; mean age, 65 years \pm 12 (SD); age range, 9–88 years) who underwent contrast-enhanced MR angiography and DSA over a period of 30 months. The patients were examined at our institution for suspected extracranial carotid artery disease based on clinical and color-coded duplex sonographic findings. The MR imaging protocol included intracranial conventional spin-echo T1-weighted and fast spin-echo T2-weighted sequences, as well as diffusion-weighted imaging. DSA and contrast-enhanced MR angiography were performed within a maximum of 2 weeks of each other. A total of 240 carotid bifurcations were evaluated in this study.

MR Angiographic Technique

All MR angiographic investigations were performed with a 1.5-T imaging system (Magnetom Vision; Siemens Medical Systems, Erlangen, Germany) equipped with a gradient overdrive. The maximum achievable gradient amplitude was 25 mT/m, and the slew rate was 180 T/m/s.

Spoiled 3D fast low-angle-shot contrast-enhanced MR angiography was performed by using a 4×2 circularly polarized phased-array neck coil. The sequence was performed with 32–36 coronal partitions each 1.67–1.94 mm thick, 2.84–3.15/1.03–1.11/1 (TR/TE/NEX), 35–40° flip angle, 70–60 \times 140 \times 280-mm³ field of view by a 36 \times 92 \times 256 image matrix, and an imaging time of 9–9.5 seconds. Reconstruction was performed without zero filling. The pixel size was 1.5 \times 1.0. The spatial resolution or voxel size was 1.8 \times 1.4 \times 1.0. The k space was filled in sequential order. Four consecutive 3D images were obtained, starting at approximately 3 seconds after the start of the bolus injection of 0.1 mmol/kg gadodiamide (Omniscan; Nycomed, Oslo, Norway). In our experience, this contrast dose is enough to provide adequate diagnostic image quality with a time-resolved contrast-enhanced MR angiographic technique. Considering the variability in patient size and weight, as well as cardiac output and hydration, we believe that only occasionally could a diagnostically significant higher image quality be attained with a higher dose of contrast material. Because of the rapid acquisition time, we believe that use of other timing strategies such as a test dose or a triggered acquisition would not noticeably improve image quality. In non-time-resolved contrast-enhanced MR angiography, timing strategies have a more important role. No breath holding was used in the current study. All injections were performed by hand (a power injector was not available) by means of a 150-cm-long catheter into an antecubital vein. Each bolus was followed immediately by a 15-mL saline flush, by using a three-way stop-cock to facilitate a smooth transition between contrast material and saline injections. Maximum duration of the total injection was 10 seconds.

The 3D image set acquired during the arterial phase of the bolus passage was identified by means of visual inspection, and that data set was subsequently subtracted from the precontrast (usually the first) data set, after the disturbing background signal intensity of fat was eliminated. Postprocessing subvolumes were generated interactively by using the MR postprocessing console to isolate each carotid artery and create 19 maximum intensity projection (MIP) images at 10° increments each. Voxels of interest on magnified images were used to facilitate determination of the extent of disease.

The total time for the 3D contrast-enhanced MR angiographic examination was 5–10 minutes, the largest part of

which was taken up by handling and positioning the patient. Over the 2½-year period of this study, we did not substantially change the technical parameters of the study protocol.

DSA Technique

Selective cerebral DSA was performed via the femoral artery, starting routinely with imaging of the aortic arch followed by selective injections of contrast material (iopamidol [Iopamiro 300]; Bracco, Milan, Italy) in both common carotid and subclavian arteries and at least one vertebral artery. The injected volume of contrast medium was 5–8 mL. Biplane DSA was performed routinely in the anteroposterior and lateral projections with a 33-cm field of view and a 1024 \times 1024 matrix. The spatial resolution was 0.32 \times 0.32 mm. If a bifurcation was not adequately visualized, additional DSA was performed in oblique projections.

Stenosis Measurement and Image Analysis

Carotid stenosis was measured on both DSA images and MR angiograms as a percentage of the diameter by using the NASCET criteria (1, 2, 39–45). The image that demonstrated the most severe stenosis was used. The diameter of the most severe stenosis was divided by the diameter of the distal cervical internal carotid artery beyond the stenosis. Carotid stenoses were measured at the same level on the DSA images and contrast-enhanced MR angiograms. The value was subtracted from 1 and then multiplied by 100 to yield the percentage diameter stenosis. The NASCET categories were arbitrarily modified to define a 98% stenosis as the presence of a signal void on MIP images at the level of the stenosis when flow was visible distal to the stenosis (9). Negative values were defined as 0% stenosis. The obstructions were classified as mild stenosis (0–29%), moderate stenosis (30–69%), severe stenosis (70–99%), or occlusion (100%). For each examination, the measurement of the exact degree of stenosis was made at the level of maximum stenosis by using high magnification.

A neuroradiologist (L.R.) reviewed image quality of the MR angiograms and DSA images. Images were evaluated for overall quality, including vascular signal intensity, venous suppression, and presence of artifacts. Evaluation criteria for overall quality were 1, excellent; 2, more than adequate for diagnosis; 3, adequate for diagnosis; and 4, nondiagnostic.

The DSA images and MR angiograms were rated in a blinded manner. The patient's name was removed from the images. The DSA images of the right carotid bifurcation were randomized, and the percentage of stenosis was determined. The left carotid images were then randomized and reviewed in a similar manner. The DSA images were evaluated by an experienced neuroradiologist (G.S.).

The MIP MR angiograms obtained in the right and left carotid bifurcations were randomized and reviewed identically for stenosis. The MIP images were reviewed by a neuroradiologist (L.R.) and a neurosurgeon in training at our neurology department (P.S.). No analysis of the source images was performed.

Statistical Analysis

A Spearman rank test was used to find any correlation between findings at DSA and those at MR angiography. Finally, the values of sensitivity, specificity, and positive and negative predictive values were established for the presence or absence of disease that would necessitate surgery (to distinguish stenoses \geq 70% from those $<$ 70%). For each MR angiogram, the degree of agreement in the interpretation of the degree of stenosis was determined by using the κ coefficient. Agreement was classified as mild ($\kappa > 0.40$ –0.69), good ($\kappa > 0.70$ –0.89), or excellent ($\kappa > 0.90$ –1.00).



FIG 1. Agreement between findings at DSA and those at MR angiography.

A, DSA image of the right carotid bifurcation shows a focal severe stenosis (arrow) of the internal carotid artery.

B, The 3D contrast-enhanced MR angiographic MIP image (magnification factor, 2) shows the focal severe stenosis (arrow), correlating well with the DSA image.

Agreement of Contrast-Enhanced MR Angiography with DSA in Evaluation of Carotid Artery Stenosis

Stenosis Grade	DSA	MRA	Agreement (%)	Estimation	
				Over	Under
Mild (0–29%)	101	91	90	10	0
Moderate (30–69%)	38	26	68	6	6
Severe (70–99%)	73	68	93	2	3
Occlusion (100%)	28	28	100	0	0

Note.—MRA indicates MR angiography. Except for agreement, data are the number of arteries.

Results

In all 120 patients, the quality of both the MR angiograms and DSA images was graded as adequate for diagnosis (grade 3) or better. All MIP MR angiograms included the aortic arch and the circle of Willis within the field of view. No relevant motion artifacts diminished the quality of the MIP images.

Grades of stenoses on MR angiograms agreed with grades of stenoses on DSA images in 89% of the carotid arteries. The agreement between findings at MR angiography and those at DSA (Fig 1) in the group of severe stenosis (70–99%) was 93% (Table). Underestimation of severe stenosis at MR angiography in comparison to that at DSA occurred in three cases, each with extremely short severe stenoses (Fig 2). Overestimation occurred in two cases, both pseudo-occlusions. In the group of moderate stenosis (30–69%), stenoses of 60–67% in six cases were overestimated (71–84%) at MR angiography.

All 28 carotid occlusions (Fig 3) were accurately detected with MR angiography. In nine cases, a pseudo-occlusion of the internal carotid artery was suspected during DSA following contrast material injection into the proximal segment of the common carotid artery. MR angiography showed pseudo-occlusion (Fig 4) in seven cases and occlusion in two cases.

The sensitivity of all the MIP images was 98%; specificity, 96%; positive predictive value, 95%; and negative predictive value, 98%. The interobserver

agreement for classification of degree of stenosis was judged as good for MR angiography ($\kappa = 89$). By using the Spearman rank test, comparison of the percentage carotid artery stenosis determined at DSA with that at MR angiography showed a significant correlation ($r = 0.91$, $P < .001$).

Discussion

In many preliminary and feasibility studies, various contrast-enhanced MR angiographic techniques have been implemented in the assessment of extracranial disease of the carotid artery. Different strategies are used to optimize image quality, including time-resolved and high-resolution techniques. Most of these reports, although based on small numbers of patients, suggest that contrast-enhanced MR angiography could become a diagnostic alternative as a fast, first screening method in patients suspected of having carotid artery disease, independently from the technique employed in the study.

DSA is still considered to be the criterion standard for imaging of arteriosclerotic supracarotid vessel disease. Harboring the risks of thromboembolic events, this technique can reduce the overall benefit of endarterectomy. Noninvasive techniques for imaging supracarotid vessels prevent this risk related to the diagnostic procedure and can be regarded as a safe alternative to DSA whenever possible.

We chose to compare time-resolved contrast-enhanced MR angiography with high-resolution DSA in 120 patients. The time-resolved technique for MR angiography is comparable to DSA: With one bolus injection of contrast agent, we have the ability to acquire dynamic data during the arterial and venous phases of contrast passage.

In the present study, contrast-enhanced 3D MR angiography was performed with a short TR and a short imaging time, applied in four sequential acquisitions. This sequence has an important advantage: Considering the arteriovenous transit time of 5–15 seconds in the normal cerebrovascular system, and maximal selective intraluminal contrast enhancement

FIG 2. Underestimation of stenoses.

A, DSA image of the left common carotid artery depicts a proximal moderate stenosis (*straight arrow*) and an extremely short distal stenosis (*curved arrow*) of the internal carotid artery.

B, 3D contrast-enhanced MR angiographic MIP image (magnification factor, 2) depicts the moderate proximal stenosis (*straight arrow*) and a short band of decreased enhancement (*curved arrow*) without clear definition of a stenosis.

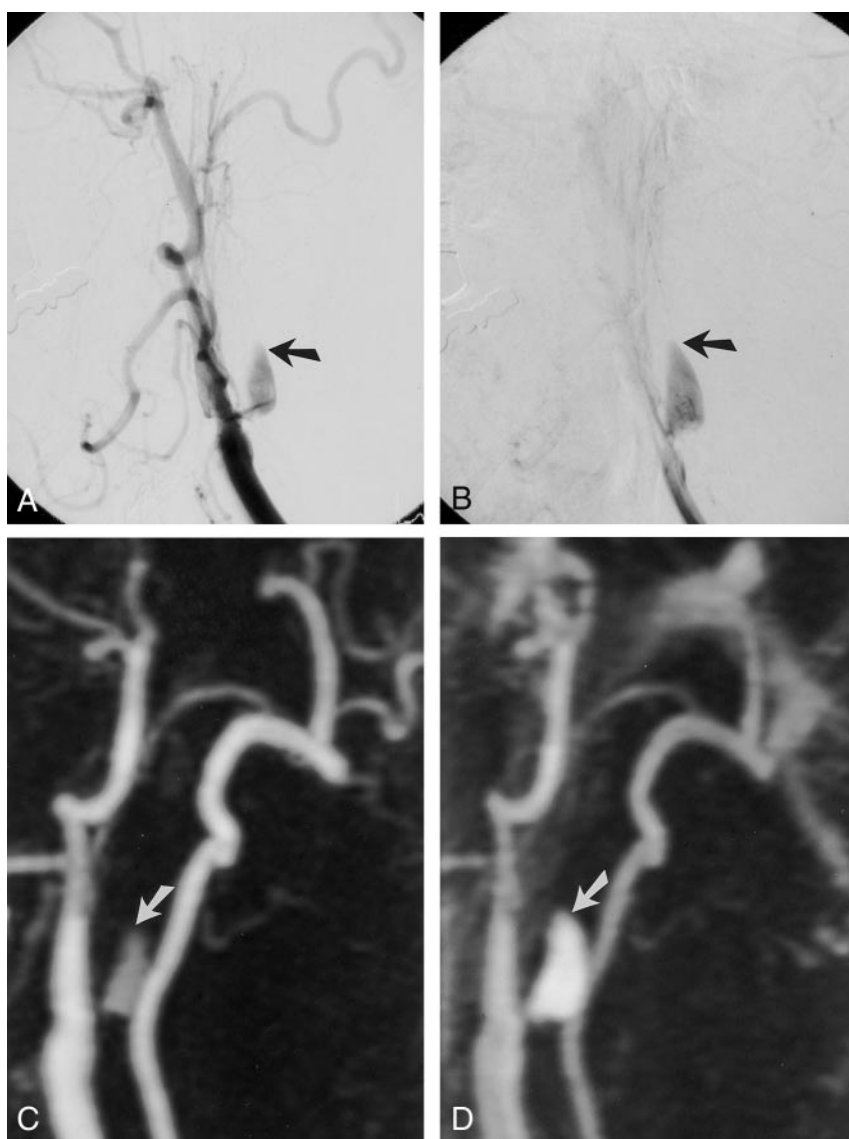
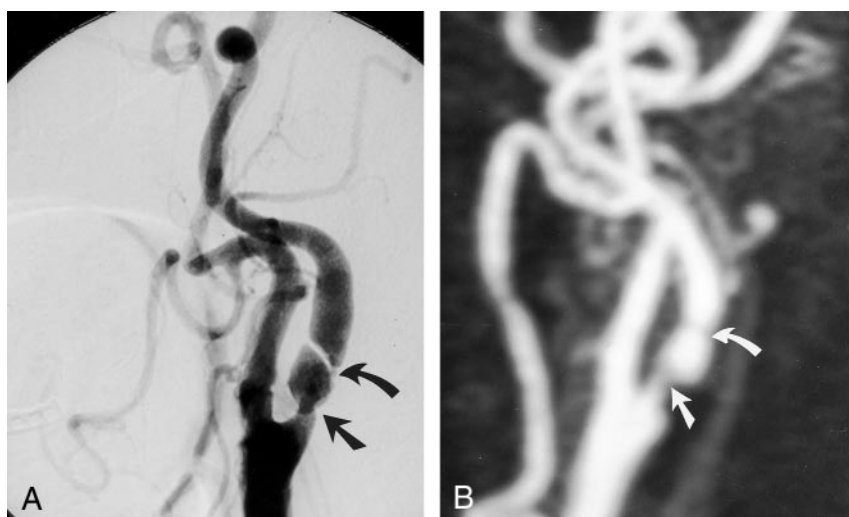


FIG 3. Carotid occlusions.

A and B, DSA images and, C and D, 3D contrast-enhanced MR angiographic MIP images (magnification factor, 1.5) of the left common carotid artery in the early (A and C) and delayed (B and D) phases demonstrate a stump (*arrow*) at the origin of the internal carotid artery and occlusion of the distal internal carotid artery.

of the carotid arteries of 10–25 seconds after intravenous administration of a bolus of contrast material, selective arterial enhancement can be obtained from

the aortic arch and the supraaortic arteries up to the circle of Willis. A delay in the arteriovenous transit time due to the presence of an arterial stenosis may

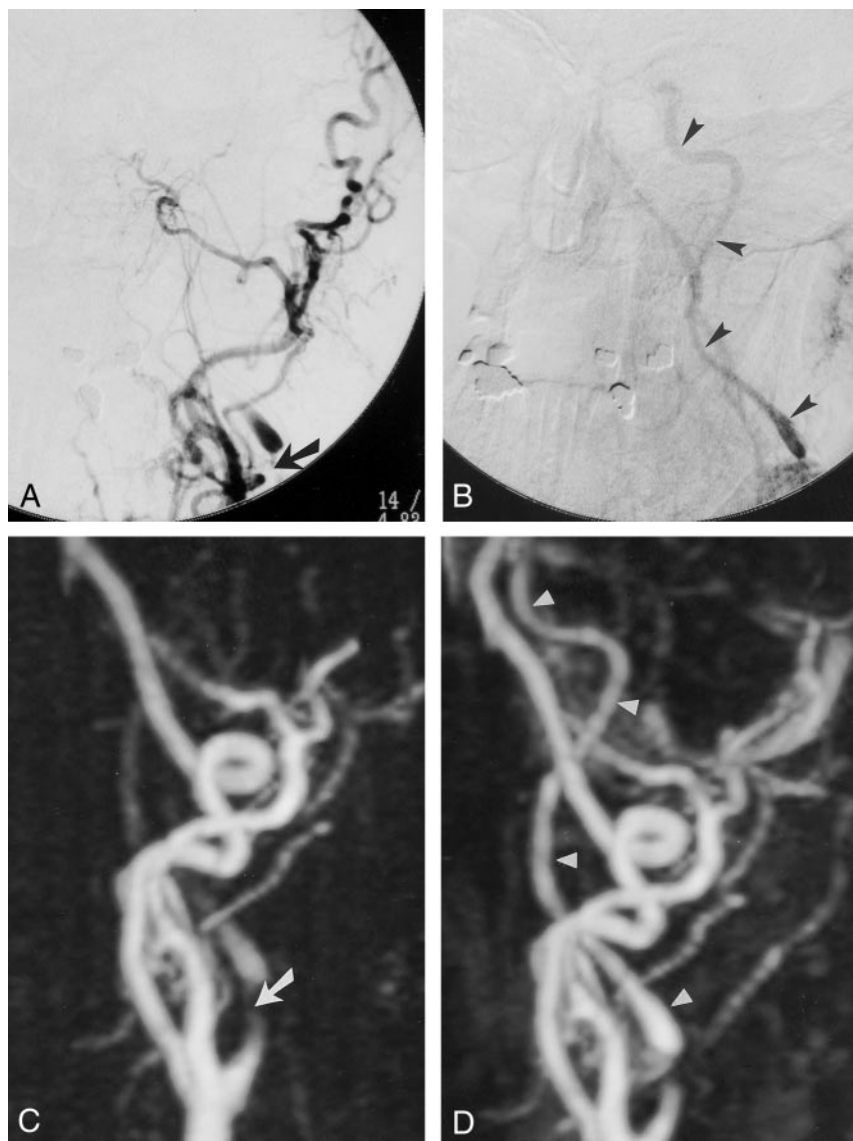


FIG 4. A and B, DSA images of the left common carotid artery demonstrate an extremely severe stenosis (arrow) at the origin of the internal carotid artery in the early phase (A), with collapse of the distal lumen ("string" sign [arrowheads]) in the delayed phase (B).

C and D, The 3D contrast-enhanced MR angiographic MIP images (1.5 magnification factor) demonstrate the extremely severe stenosis (arrow in C) at the origin of the internal carotid artery and opacification of the collapsed distal lumen in only the delayed phase (arrowheads in D).

actually improve the increase in intraluminal signal intensity during the first pass of the contrast-enhanced blood in the carotid and vertebral arteries. Using a short TE, we were able to reduce flow voids in the intra- and poststenotic regions of the carotid artery, because intravoxel spin dephasing due to fast or turbulent flow can be minimized with this technique. This results in a reduction in the subsequent overestimation of the stenosis, if the remaining vessel lumina are 1 mm or less. In addition, the high signal-to-noise ratio allows visualization of vessels in subvoxel volumes during the first passage of gadolinium-based contrast material. However, the effective resolution exceeds the numerical resolution. In our experience, the minimum accessible vessel diameter is estimated to be between 0.3 and 0.5 mm, thus approaching the resolution limit of DSA. At the level of a stenosis, the minimum accessible vessel diameter increases to approximately 0.8–1 mm.

Because of the rapid acquisition time with four consecutive measurements used in our study, other

timing strategies such as a test dose or triggered acquisition would not noticeably improve image quality. The major advantages of this MR angiographic sequence are the rapid acquisition time with subsequent reduction in flow- and patient-related movement artifacts, the high spatial resolution, and the large acquisition volume. However, results of other studies using high-spatial-resolution sequences are also very promising, although larger groups of patient are needed to confirm these results (9, 13, 14, 22, 25, 29).

The NASCET study showed that patients with an internal carotid artery stenosis of at least 70% diameter benefit from endarterectomy (1, 2). More recently, the indication for carotid endarterectomy has been established also for patients with asymptomatic carotid artery stenosis ($\geq 60\%$) as presented in the ACAS (5).

In our series, the agreement between MR angiography and DSA in the group of severe stenosis (70–99%) was 93%. In three cases, a short severe stenosis of the extracranial internal carotid artery was under-

estimated with MR angiography compared with DSA. In six cases, a 60–67% internal carotid artery obstruction was overestimated (71–84%) with MR angiography.

An exact differentiation between occlusion and pseudo-occlusion is important for the therapeutic approach and may be a problem when using duplex sonography. All occlusions were correctly detected with MR angiography. In seven cases, pseudo-occlusions of the internal carotid artery were clearly detected with MR angiography. In two cases, pseudo-occlusions assessed with DSA were overestimated (occlusion) with MR angiography. However, retrospectively, a careful analysis of the later-phase MR angiogram showed that the pseudo-occlusion similar to that on the DSA image, according to a very slow blood flow. Thus, in cases of suspected occlusion or pseudo-occlusion, careful analysis of the later-phase image of time-resolved MR angiography is mandatory. A spatial resolution of 1 mm or less is needed.

As an alternative to MIP images, multiplanar reconstruction images can be used for stenosis measurement. We did not use this approach because it is time-consuming and previous reports indicate that measurements with MIP and those with multiplanar reconstruction images do not differ significantly (43, 45). In selected cases, the study of source images could be critical to evaluate subtle lesions (43, 45).

Although no transient or permanent neurologic complication occurred after DSA in our study, carotid angiography remains associated with a substantial number of risks and potential complications. Whereas mortality is low, the frequency of neurologic events varies from 0.45% to 2.6%. In contrast, MR imaging and MR angiography have no known adverse effects and contrast-enhanced MR angiography has a low complication rate. The unlimited number of views from a single acquisition of a carotid bifurcation MR angiographic study can represent an important advantage compared with DSA, in which a minimum of two injections of contrast material are necessary to obtain four views with standard biplane imaging. Another advantage of the contrast-enhanced MR angiographic approach is the ability to display the carotid bifurcation anatomy in a format similar to that of DSA, also in the time resolution. The major advantages of contrast-enhanced MR angiography is the reduction of flow and patient-related movement artifacts, the good spatial resolution, and the large acquisition volume from the aortic arch to the circle of Willis. This allows a good estimation of the degree of carotid stenosis, a differentiation between occlusion and pseudo-occlusion, and detection of tandem stenoses, which is important for clinical and therapeutic management.

In our hospital, contrast-enhanced MR angiography in association with Doppler sonography has now mostly replaced DSA for diagnostic imaging of the cervical arteries. We use contrast-enhanced MR angiography not only as a screening procedure but also for preoperative (endarterectomy) or preinterventional (percutaneous angioplasty and stent placement) evaluation. We still continue to perform DSA

in unclear cases. For postoperative follow-up, Doppler sonography plus contrast-enhanced MR angiography is the method of choice to display morphologic results (46).

We believe the further developments in the hardware and software of the new generation of MR imagers will permit a clinically even more sufficient and stable quality of contrast-enhanced MR angiography (47).

Conclusion

Since several clinical trials have proved the benefit of carotid thromboendarterectomy in patients with moderate or severe stenosis, imaging of the carotid artery has been the focus of considerable attention. To reduce risks related to DSA, many noninvasive imaging approaches have been advocated, such as contrast-enhanced MR angiography. Many preliminary and feasibility studies suggest that contrast-enhanced MR angiography could become a diagnostic alternative as a fast screening method in patients suspected of having carotid artery disease, independently from the technique used in the study. The time-resolved technique for contrast-enhanced MR angiography is comparable to DSA: With one bolus injection of contrast agent, we have the ability to acquire dynamic data during the arterial and venous phases of contrast passage.

Our results indicated a statistically significant correlation between contrast-enhanced MR angiography and DSA for minimal, moderate, and severe stenoses and occlusions. This investigation with a large number of patients confirms that contrast-enhanced MR angiography could become a diagnostic alternative to DSA in the treatment of patients with carotid artery disease.

References

1. North American Symptomatic Carotid Endarterectomy Trial Collaborators. **Beneficial effect of carotid endarterectomy in symptomatic patients with high-grade carotid stenosis.** *N Engl J Med* 1991;325:445–453
2. The (NASCET) North American Symptomatic Carotid Endarterectomy Trial Steering Committee. **North American Symptomatic Carotid Endarterectomy Trial: methods, patients characteristics and progress.** *Stroke* 1991;22:711–720
3. European Carotid Surgery Trials Collaborative Group. **MRC European carotid surgery trial: interim results for symptomatic patients with severe (70–99%) or mild (0–29%) carotid stenosis.** *Lancet* 1991;337:1235–1243
4. Hobson RW II, Weiss DG, Fields WS, et al. **Efficacy of carotid endarterectomy for asymptomatic carotid stenosis. The Veterans Affairs Cooperative Study Group.** *N Engl J Med* 1993;328:221–227
5. **Endarterectomy for asymptomatic carotid artery stenosis. Executive Committee for the Asymptomatic Carotid Atherosclerosis Study.** *JAMA* 1995;273:1421–1428
6. Barnett HJ, Taylor DW, Eliasziw M, et al. **Benefit of carotid endarterectomy in patients with symptomatic moderate or severe stenosis: North American Symptomatic Carotid Endarterectomy Trial Collaborators.** *N Engl J Med* 1998;339:1415–1425
7. Levy RA, Prince MR. **Arterial-phase three-dimensional contrast-enhanced MR angiography of the carotid artery.** *AJR Am J Roentgenol* 1996;167:211–215
8. Krinsky G, Rofsky N, Flyer M, et al. **Gadolinium-enhanced three-dimensional MR angiography of acquired arch vessel disease.** *AJR Am J Roentgenol* 1996;167:981–987

9. Remonda L, Heid O, Schroth G. Carotid artery stenosis, occlusion, and pseudo-occlusion: first-pass, gadolinium-enhanced, three-dimensional MR angiography—preliminary study. *Radiology* 1998; 209:95–102
10. Leclerc X, Gauvrit JY, Nicol L, Pruvo JP. Contrast-enhanced MR angiography of the craniocervical vessels: a review. *Neuroradiology* 1999;41:867–874
11. Sardanelli F, Zandrino F, Parodi RC, De Caro G. MR angiography of internal carotid arteries: breath-hold Gd-enhanced 3D fast imaging with steady-state precession versus unenhanced 2D and 3D time-of-flight techniques. *J Comput Assist Tomogr* 1999;23:208–215
12. Cloft HJ, Murphy KJ, Prince MR, Brunberg JA. 3D gadolinium-enhanced MR angiography of the carotid arteries. *Magn Reson Imaging* 1996;14:593–600
13. Serfaty JM, Chirouss P, Chevallier JM, Ecochard R, Froment JC, Douek PC. Accuracy of three-dimensional gadolinium-enhanced MR angiography in the assessment of extracranial carotid artery disease. *AJNR Am J Roentgenol* 2000;175:455–463
14. Cloft HJ, Kallmes DF. Gadolinium-enhanced MR angiography for carotid artery disease. *AJR Am J Roentgenol* 1999;172:833–834
15. Scarabino T, Carriero A, Giannatempo GM, et al. Contrast-enhanced MR angiography (CE MRA) in the study of the carotid stenosis: comparison with digital subtraction angiography (DSA). *Neuroradiology* 1999;26:87–91
16. Slosman F, Stolpen AH, Lexa FJ, et al. Extracranial atherosclerotic carotid artery disease: evaluation of non-breath-hold three-dimensional gadolinium-enhanced MR angiography. *AJR Am J Roentgenol* 1998;170:489–495
17. Willig DS, Turski PA, Frayne R, et al. Contrast-enhanced 3D MR DSA of the carotid artery bifurcation: preliminary study of comparison with unenhanced 2D and 3D time-of-flight MR angiography. *Radiology* 1998;208:447–451
18. Martinat P, Leclerc X, Gauvrit JY, Giboreau F, Pruvo JP. Contribution of fast-sequence three-dimensional MRI angiography with gadolinium injection in the evaluation of supra-aortic vessels. *J Radiol* 1998;79:673–680
19. Jager HR, Moore EA, Bynevelt M, et al. Contrast-enhanced MR angiography in patients with carotid artery stenosis: comparison of two different techniques with an unenhanced 2D time-of-flight sequence. *Neuroradiology* 2000;42:240–248
20. Huston J 3rd, Fain SB, Wald JT, et al. Carotid artery: elliptic centric contrast-enhanced MR angiography compared with conventional angiography. *Radiology* 2001;218:138–143
21. Sardanelli F, Zandrino F, Parodi RC, De Caro G. MR angiography of internal carotid arteries: breath-hold Gd-enhanced 3D fast imaging with steady-state precession versus unenhanced 2D and 3D time-of-flight techniques. *J Comput Assist Tomogr* 1999;23:208–215
22. Leclerc X, Martinat P, Godefroy O, et al. Contrast-enhanced three-dimensional fast imaging with steady-state precession (FISP) MR angiography of supraaortic vessels: preliminary results. *AJNR Am J Neuroradiol* 1998;19:1405–1413
23. Kim JK, Farb RI, Wright GA. Test bolus examination in the carotid artery at dynamic gadolinium-enhanced MR angiography. *Radiology* 1998;206:283–289
24. Leclerc X, Gauvrit JY, Nicol L, Martinat P, Pruvo JP. Gadolinium-enhanced fast three-dimensional angiography of the neck: technical aspect. *Invest Radiol* 1999;34:204–210
25. Kollias SS, Binkert CA, Ruesch S, Valavanis A. Contrast-enhanced MR angiography of the supra-aortic vessels in 24 seconds: a feasibility study. *Neuroradiology* 1999;41:391–400
26. Aoki S, Nakajima H, Kumagai H, Araki T. Dynamic contrast-enhanced MR angiography and MR imaging of the carotid artery: high-resolution sequences in different acquisition planes. *AJNR Am J Neuroradiol* 2000;21:381–385
27. Fellner FA, Fellner C, Wutke R, et al. Fluoroscopically triggered contrast-enhanced 3D MR DSA and 3D time-of-flight turbo MRA of the carotid arteries: first clinical experiences in correlation with ultrasound, x-ray angiography, and endarterectomy findings. *Magn Reson Imaging* 2000;18:575–585
28. Melhem ER, Serfaty JM, Jones L, et al. Contrast-enhanced MR angiography: the effects of k-space truncation on luminal representation in a carotid artery phantom model. *AJNR Am J Neuroradiol* 2000;21:1028–1031
29. Huston J 3rd, Fain SB, Riederer SJ, Wilman AH, Bernstein MA, Busse RF. Carotid arteries: maximizing arterial to venous contrast in fluoroscopically triggered contrast-enhanced MR angiography with elliptic centric view ordering. *Radiology* 1999;211:265–273
30. Isoda H, Takehara Y, Isogai S, et al. Software-triggered contrast-enhanced three-dimensional MR angiography of the intracranial arteries. *AJR Am J Roentgenol* 2000;174:371–375
31. Korosec FR, Turski PA, Carroll TJ, Mistretta CA, Grist TM. Contrast-enhanced MR angiography of the carotid bifurcation. *J Magn Reson Imaging* 1999;10:317–325
32. Melhem ER, Caruthers SD, Faddoul SG, Tello R, Jara H. Use of three-dimensional MR angiography for tracking a contrast bolus in the carotid artery. *AJNR Am J Neuroradiol* 1999;20:263–266
33. Isoda H, Takehara Y, Isogai S, et al. Technique for arterial-phase contrast-enhanced three-dimensional MR angiography of the carotid and vertebral arteries. *AJNR Am J Neuroradiol* 1998;19:1241–1244
34. Kaandorp DW, Kopinga K, Wijn PF. Venous signal suppression in 3D dynamic Gd-enhanced carotid artery imaging using the eigen-image filter. *Magn Reson Med* 1999;42:307–313
35. Steffens JC, Link J, Heller M. Contrast-enhanced magnetic resonance angiography of the cervical arteries. *Invest Radiol* 1998;33: 573–577
36. Krinsky G, Maya M, Rofsky N, et al. Gadolinium-enhanced 3D MRA of the aortic arch vessels in the detection of atherosclerotic cerebrovascular occlusive disease. *J Comput Assist Tomogr* 1998; 22:167–178
37. Leclerc X, Nicol L, Gauvrit JY, Le Thuc V, Leys D, Pruvo JP. Contrast-enhanced MR angiography of supraaortic vessels: the effect of voxel size on image quality. *AJNR Am J Neuroradiol* 2000;21:1021–1027
38. Leclerc X, Lucas C, Godefroy O, et al. Preliminary experience using contrast-enhanced MR angiography to assess vertebral artery structure for the follow-up of suspected dissection. *AJNR Am J Neuroradiol* 1999;20:1482–1490
39. Vanninen RL, Manninen HI, Koivisto K, et al. Carotid stenosis by digital subtraction angiography reproducibility of the European Carotid Surgery Trial and the North American Symptomatic Carotid Endarterectomy Trial measurement methods and visual interpretation. *AJNR Am J Roentgenol* 1994;15:1635–1641
40. Brown PM, Johnston KW. The difficulty of quantifying the severity of carotid stenosis. *Surgery* 1992;92:468–473
41. Rothwell PM, Gibson RJ, Slattery J, Warlow CP. Prognostic value and reproducibility of measurement of carotid stenosis: a comparison of three methods on 1001 angiograms. *Stroke* 1994;25:2440–2444
42. Fox AJ. How to measure carotid stenosis. *Radiology* 1993;186:316–318
43. Vanninen RL, Manninen HI, Partanen PK, Tulla H, Vainio PA. How should we estimate carotid stenosis using magnetic resonance angiography? *Neuroradiology* 1996;38:299–305
44. Staikov IN, Arnold M, Mattle HP, et al. Comparison of the ECST, CC, and NASCET grading methods and ultrasound for assessing carotid stenosis. European Carotid Surgery Trial. North American Symptomatic Carotid Endarterectomy Trial. *J Neurol* 2000;247: 681–686.
45. Huston J III, Lewis BD, Wiebers DO, Meyer FB, Riederer SJ, Weaver AL. Carotid artery: prospective blinded comparison of two-dimensional time-of-flight MR angiography with conventional angiography and duplex US. *Radiology* 1993;186:339–344
46. Wetzel S, Boos M, Bongartz G, Radu EW. Selection of patients for carotid thromboendarterectomy: the role of magnetic resonance angiography. *J Comput Assist Tomogr* 1999;23(suppl 1):S91–S94
47. Rofsky NM, Adelman MA. Gadolinium-enhanced MR angiography of the carotid arteries: a small step, a giant leap? *Radiology* 1998;209:31–34