ABSTRACT

BACKGROUND AND PURPOSE: There is need to improve image acquisition speed for MR imaging in evaluation of patients with acute ischemic stroke. The purpose of this study was to evaluate the feasibility of a 3T MR stroke protocol that combines low-dose contrast-enhanced MRA and dynamic susceptibility contrast perfusion, without additional contrast.

METHODS: Thirty patients with acute stroke who underwent 3T MR imaging followed by DSA were retrospectively enrolled. TOF-MRA of the neck and brain and 3D contrast-enhanced MRA of the craniocervical arteries were obtained. A total of 0.1 mmol/kg of gadolinium was used for both contrast-enhanced MRA (0.05 mmol/kg) and dynamic susceptibility contrast perfusion (0.05 mmol/kg) (referred to as half-dose). An age-matched control stroke population underwent TOF-MRA and full-dose (0.1 mmol/kg) dynamic susceptibility contrast perfusion. The cervicocranial arteries were divided into 25 segments. Degree of arterial stenosis on contrast-enhanced MRA and TOF-MRA was compared with DSA. Time-to-maximum maps (>6 seconds) were evaluated for image quality and hypoperfusion. Quantitative analysis of arterial input function curves, SNR, and maximum T2* effects were compared between half- and full-dose groups.

RESULTS: The intermodality agreements (κ) for arterial stenosis were 0.89 for DSA/contrast-enhanced MRA and 0.63 for DSA/TOF-MRA. Detection specificity of >50% arterial stenosis was lower for TOF-MRA (89%) versus contrast-enhanced MRA (97%) as the result of overestimation of 10% (39/410) of segments by TOF-MRA. The DWI-perfusion mismatch was identified in both groups with high interobserver agreement (r = 1). There was no significant difference between full width at half maximum of the arterial input function curves (P = .14) or the SNR values (0.6) between the half-dose and full-dose groups.

CONCLUSIONS: In patients with acute stroke, combined low-dose contrast-enhanced MRA and dynamic susceptibility contrast perfusion at 3T is feasible and results in significant scan time and contrast dose reductions.

ABBREVIATIONS: AIF = arterial input function; CE-MRA = contrast-enhanced MRA; FWHM = full width at half maximum; GRAPPA = generalized autocalibrating partially parallel acquisition; Tmax = time-to-maximum

Published February 6, 2014 as 10.3174/ajnr.A3848

Copyright 2014 by American Society of Neuroradiology.
contrast dosages as low as 0.05 mmol/kg have been used with promising results.\textsuperscript{10,11}

The purpose of this study was to prospectively evaluate the feasibility of a combined low-dose CE-MRA and DSC perfusion acute stroke protocol at 3T that requires no additional contrast during diagnostic assessment and to compare the results with a routine stroke protocol that employs TOF-MRA. Two hypotheses were tested: 1) low-dose CE-MRA provides comparable or superior diagnostic performance in the detection of both extracranial and intracranial arterio-occlusive disease in comparison with TOF-MRA, with potential for significant reduction in acquisition time; 2) DSC perfusion by use of half of the conventional dose is feasible with comparable image quality.

**MATERIALS AND METHODS**

This retrospective study was conducted between August 2011 and April 2013. All examinations were performed in accordance with institutional review board guidelines with an approved study protocol. Inclusion criteria included: 1) patients with clinical suspicion of acute ischemic stroke who presented within the first 9 hours from symptom onset; 2) acquisition of 3T MR imaging as the initial imaging test; and 3) DSA within 24 hours after symptom onset. DSA was performed either because of failed response to conventional treatment, including IV tPA (n = 4), or detection of a proximal clot and major arterial occlusion requiring further endovascular treatment (n = 26). Exclusion criteria included patients with severe contrast allergy, MR-incompatible implants, and severe claustrophobia. The baseline NIHSS scores and median time from stroke onset to MR imaging were documented for each patient.

**Image Acquisition**

All MR examinations were performed on a 3T (Trio) clinical MR scanner (Siemens, Erlangen, Germany). A combination 12-channel head (n = 8) and neck coil (n = 4) was used for radiofrequency signal reception. All patients underwent routine MR stroke protocol, which included DWI, FLAIR, gradient refocused echo, brain 3D TOF-MRA and 2D TOF-MRA of the neck, and DSC perfusion imaging.

MRA sequence parameters were brain TOF-MRA: 3D multi-slab gradient-echo sequence (TR/TE: 25/3.8 ms, flip angle: 18°, 5 slabs each with 30 partitions, FOV: 200 × 200 mm, matrix: 256 mm, section thickness: 0.7 mm, generalized autocalibrating partially parallel acquisition (GRAPPA) × 2, acquisition time: 5:50 minutes); neck TOF-MRA: 2D gradient-echo sequence (TR/TE: 21/5.9 ms, flip angle: 45°, 110 partitions, section thickness: 2 mm, FOV: 250 × 180 mm, matrix: 256 mm, GRAPPA × 2, acquisition time: 3:30 minutes).

A high spatial resolution 3D CE-MRA with centric k-space ordering was performed just before the perfusion imaging by use of a fast spoiled–gradient refocused echo sequence in the coronal plane: (TR/TE: 3/1.2 ms, flip angle: 25°, FOV: 340 × 280 mm, matrix: 448 mm). One hundred twenty sections with a thickness of 0.8 mm were acquired in the coronal plane, providing arterial coverage from the aortic arch to the cranial vertex. With the use of a GRAPPA algorithm\textsuperscript{12} with an acceleration factor of 4, a near isotropic sub-millimeter voxel size 3D dataset (0.8 × 0.9 × 0.8 mm\textsuperscript{3}) was achieved during a 22-second acquisition. A modified 2-phase contrast injection scheme\textsuperscript{13} was used to perform CE-MRA and DSC perfusion imaging, without need for additional contrast. To accomplish this, a total of 0.1 mmol/kg of gadolinium (MultiHance, Bracco Diagnostics, Princeton, New Jersey) that is normally used routinely for MR perfusion was diluted with normal saline to a total volume of 50 mL. Three mL of contrast solution was injected and flushed with 20 mL of saline to determine the transit time from the arm vein to the cervical carotid arteries. A total of 22 mL of contrast solution was then injected and flushed with 20 mL of saline for the CE-MRA acquisition. The contrast injection rate for timing run and CE-MRA was 1.5 mL/s. Subsequently, the remaining 25 mL of contrast solution was injected and flushed with 20 mL of saline at 5 mL/s for the MR perfusion scan, which was performed last. The DSC perfusion was performed by use of a single-shot, gradient echo-planar sequence (TR/TE: 1920/32 ms, flip angle: 90°, 25 partitions, 5-mm section thickness, and an in-plane voxel size of 1.7 × 1.7 mm\textsuperscript{2}). With the use of GRAPPA × 2, 45 dynamic frames were obtained over a 90-second acquisition time.

**Image Analysis**

**MRA.** The source data and reconstructed MRA images by use of MIP were used for image analysis. The cervicocranial arteries were divided into 25 segments, including 1,2: bilateral common carotid arteries; 3,4: bilateral cervical ICAs; 5,6: bilateral petro cavernous ICAs; 7,8: bilateral supraclinoid ICAs; 9–12: bilateral anterior cerebral arteries (A1 and A2 segments); 13–16: bilateral MCAs (M1 and M2 segments); 17,18: bilateral cervical vertebral arteries; 19,20: bilateral intradural segments of vertebral arteries; 21: basilar artery; and 22–25: bilateral posterior cerebral arteries (P1 and P2 segments).

Arterial segments were examined for stenoses independently by 2 neuroradiologists by use of the CE-MRA and TOF-MRA datasets during separate sessions. Dices were presented in random order to avoid recall bias. Observers were blinded to results of other imaging. The severity of arterial disease was evaluated by use of a 1–4 grading scale: 1, no stenosis; 2, mild stenosis with ≤50% luminal narrowing; 3, significant stenosis with luminal narrowing >50 and <99%; and 4, occlusion. When 2 or more stenoses were detected in the same vessel segment, the most severe stenosis was used for grading and analysis. Finally, if necessary, discordant reads on both the CE-MRA and TOF-MRA datasets evaluated for arterial stenosis or occlusion were resolved by consensus review and used to determine the diagnostic accuracy of each technique by use of preprocedural DSA as the reference standard.

**DSC Perfusion Analysis.** DSC perfusion datasets were processed by use of FDA-approved software (Olea Sphere; Olea Medical, La Ciotat, France). DSC analysis consisted of the following steps: 1) truncation of the first 5 time points in the DSC time-series, 2) calculation of prebolus signal intensity on a voxel wise basis, followed by 3) conversion of truncated DSC time-series to a concentration-time curve on the basis of the T2\textsuperscript{*} relaxivity of the contrast agent. The change in signal intensity with respect to baseline was
converted to the change in tissue T2* as a result of contrast agent injection, taken with respect to precontrast baseline time points. The arterial input function (AIF) was selected automatically by the perfusion software, and the deconvoluted perfusion parameters were calculated by use of a block-circulant singular value decomposition technique.\textsuperscript{14} Quantitative analysis of the AIF curves was performed to determine the peak and full width at half maximum (FWHM). The infarction volume was automatically calculated by means of the software, by use of ADC threshold \(600 \times 10^{-6}\) mm\(^2\)/s. The time-to-maximum maps, with the use of a threshold of \(6\) seconds, were evaluated by observers independently in each group to determine the presence of hypoperfusion and DWI-perfusion mismatch by use of a 3-scale scoring system: 1, mismatch present, confident; 2, mismatch may be present, less confident; 3, no mismatch present.

The volumes of DWI abnormality, hypoperfused region by use of Tmax \(>6\) seconds, and DWI-diffusion mismatch ratios were generated automatically by the software. These values were used as reference to compare the accuracy of subjective mismatch scores (Fig 2).

Quantitative analysis of the SNR values and maximum T2* effect was performed by use of an ROI-based analysis. ROIs approximately 1 cm in diameter were placed on the normal contralateral postcentral gyrus and corresponding subcortical white matter. The signal intensity of these ROIs was calculated over the entire 45 sequential dynamic perfusion frames, and the mean \(\pm\) SD of the signal intensity was noted for each perfusion scan. Maximum T2* effect, which is the maximum signal intensity drop during the intracerebral passage of an intravenous contrast bolus, was also calculated for each scan. To minimize the nonuniform distribution of noise across the field of view caused by parallel imaging, 2 ROIs were placed in the extracranial space (1 at the top and 1 at the bottom of field of view) to determine the background noise. The standard deviation of the mean of these background ROIs in all 45 dynamic frames (90 ROIs) was defined as image noise and was used to calculate the SNR. The baseline SNR was also evaluated by use of the precontrast perfusion images, which were available during the first 4 dynamic frames (initial 8 seconds) before contrast arrival. This was performed to investigate whether the half-dose perfusion scans obtained after CE-MRA were contaminated with recirculating contrast agent and/or had a reduced baseline signal in comparison to the full-dose control perfusion group.

**Statistical Analysis**

Statistical analysis was performed by use of MedCalc (Version 12.2.1, MedCalc Software Mariakerke, Belgium). The degree of interobserver and intermodality agreement for estimation of arterial stenosis was determined by kappa (\(\kappa\)) coefficients and calculation of 95% CI. The sensitivity and specificity of CE-MRA and TOF-MRA for stenosis \(>50\%\) was calculated for each observer, by use of preprocedural DSA images as the standard of reference. Spearman-Rank correlation coefficient was used to evaluate the qualitative scores of perfusion maps and interobserver confidence. The SNR values, maximum T2* effects, and peak and FWHM of the AIF curves were plotted as mean \(\pm\) SD, and the differences between the full- and half-dose groups were evaluated by means of a \(t\) test. The significance level was defined as \(P < .05\) (2-sided).

**Cohort Controlled Group**

To perform a comparative analysis for the DSC perfusion parameters, an age-matched control population of 30 patients (12 women, 18 men; age range, 43–90 years; average, 69 years) with

---

**FIG 1.** A 55-year-old woman presented with acute onset aphasia; NIHSS score 20. Full-thickness MIP of the entire supra-aortic arteries (A) and focused oblique reformatted (B, C, and D) thin MIP images from CE-MRA, coronal MIP images from TOF-MRA of the brain (E) and neck (F), and focused oblique views from DSA (G) after left common and ICA injections are demonstrated. Both CE-MRA and DSA demonstrate mild stenosis of the left distal common carotid artery (arrow in D and arrowhead in G), high-grade stenosis of the distal left ICA near the petrous segment (arrow on C and G), and occlusion of the ICA bifurcation with extension into the left M1 (arrow in B and black arrow in G). Because of longer acquisition time, both TOF-MRA of the neck (F) and brain (E) are degraded by motion artifact. The mild stenosis at the left common carotid artery is not clearly seen. In addition, because of distal occlusion and sluggish flow, the entire left high cervical, petrocavernous, and supracliniod ICA segments appear to be occluded.
strokes were identified by CE-MRA: mild (≤50%): n = 14; high-grade (>50–99%): n = 18; and occlusion: n = 20. Eighty-two segmental arterial stenoses were identified by TOF-MRA: mild (≤50%): n = 20; high-grade (>50–99%): n = 22; and occlusion: n = 40. The interobserver agreements were k = 0.78; 95% CI, 0.65–0.90 for CE-MRA; k = 0.86; 95% CI, 0.78–0.92 for TOF-MRA.

Preprocedural DSA images were available for 410 arterial segments, and identified 50 arterial stenoses: mild (≤50%): n = 19; high-grade (>50–99%): n = 12; and occlusion: n = 19. Table 1 shows the distribution of arterial stenoses detected by TOF-MRA and CE-MRA in comparison to DSA. Table 2 shows the diagnostic performance and agreement of CE-MRA and TOF-MRA in comparison with DSA as the standard of reference. Compared with DSA, by use of CE-MRA, 1 arterial segment with high-grade stenosis was graded as occlusion and 8 arterial segments with mild stenosis were graded high-grade stenoses. This resulted in a sensitivity and specificity of 100% and 97%, respectively, for the detection of high-grade arterial stenosis (>50%). With the use of TOF-MRA, 21 arterial segments with varying degrees of stenosis were graded as occluded and 18 arterial segments with no to mild stenosis were graded as high-grade stenosis. This resulted in sensitivity and specificity of 100% and 89%, respectively, for the detection of high-grade arterial stenosis (>50%).

**DSC Perfusion**

The mean ± SD of the volume of infarction on DWI and hypoperfusion with Tmax >6 seconds were 16.5 ± 7.6 (mL)/120 ± 48 (mL) for the full-dose group and 18 ± 9 (mL)/137 ± 70 (mL) for the half-dose group (P = .2). The mismatch ratios calculated automatically by the software demonstrated DWI-perfusion mismatch in all patients in both groups. DWI-perfusion mismatch was identified in 100% of the full-dose group and 90% of the half-dose group by both observers, with high confidence (r = 1).

There was no statistically significant difference in the SNR and background noise values between half- and full-dose groups (Table 3). When the baseline SNR of the initial phase of the perfusion study before arrival of contrast was evaluated, there was no significant difference (P = .6) between the mean baseline SNR of the full-dose (342 ± 101) and half-dose groups (330 ± 67), indicating an absence of significant recirculation effects. The maximum T2* signal drop values were higher in the full-dose group (P = .01). There was statistically significant higher AIF peak in the full-dose group (P = .002). There was however, no statistically significant

---

**RESULTS**

A total of 30 patients (18 men; age range, 38–84 years; average, 62 years) met our inclusion criteria. Baseline NIHSS scores ranged from 4–24, with a median of 16. The median time from presentation to MR imaging was 5 hours (range, 1–8 hours).

All studies were completed successfully. In 2 patients (7%), evaluation of the aortic arch and proximal aspects of supra-aortic arteries on CE-MRA was degraded by motion artifact. In 15 studies (50%), the image quality of TOF-MRA of the neck was degraded by motion artifact (Fig 1). A total of 52 segmental arterial

---

**FIG 2.** A, 62-year-old man with acute stroke; NIHSS score, 14. MR imaging was performed at 3T, 5 hours after the onset (from cohort group: 0.1 mmol/kg of gadolinium used for dynamic susceptibility contrast perfusion). B, 58-year-old man with acute stroke; NIHSS score, 16. MR imaging was performed at 3T, 3 hours after the onset (from study group: 0.05 mmol/kg gadolinium was used for dynamic susceptibility contrast perfusion). Serial DWI, T_max, and superimposed DWI-Tmax demonstrate diagnostic image quality in both patients. A large perfusion-diffusion mismatch was identified with high confidence in both patients. The corresponding arterial input function curves are shown. There is approximately 24% lower arterial peak in half dose (6.7) compared with full dose (8.9). There is only small difference between the full width at half maximum of full dose (13) versus half dose (15).
Intermodality agreement for TOF-MRA, Intermodality agreement for CE-MRA: Max indicates maximum.

Note: between full dose (0.1 mmol/kg) and half dose (0.05 mmol/kg)

Table 3: Comparative quantitative analysis of DSC perfusion

| Arterial Segment                  | TOF-MRA Stenosis Severity | CE-MRA Stenosis Severity | DSA Stenosis Severity |
|-----------------------------------|---------------------------|--------------------------|-----------------------|
|                                   | ≤50% | >50%, <99% | Occlusion | ≤50% | >50%, <99% | Occlusion | ≤50% | >50%, <99% | Occlusion |
| Common carotid artery             | 3    | 2         | 3         | 5    | 1         | 0         | 3    | 0         | 0         |
| Cervical ICA                      | 2    | 1         | 6         | 4    | 4         | 4         | 7    | 5         | 4         |
| Petrocerebral ICA                 | 2    | 3         | 4         | 1    | 1         | 1         | 0    | 2         | 1         |
| Supraclinoid ICA                  | 3    | 4         | 6         | 1    | 5         | 3         | 1    | 1         | 3         |
| MCA                               | 3    | 6         | 14        | 1    | 2         | 11        | 2    | 1         | 10        |
| Anterior cerebral artery          | 2    | 2         | 2         | 0    | 1         | 0         | 2    | 1         | 0         |
| Cervical vertebral artery         | 0    | 0         | 1         | 1    | 0         | 1         | 2    | 0         | 1         |
| Intradural vertebral artery       | 3    | 0         | 3         | 1    | 0         | 0         | 0    | 0         | 0         |
| Basilar artery                    | 2    | 1         | 0         | 0    | 0         | 0         | 0    | 0         | 0         |
| Posterior cerebral artery         | 0    | 3         | 1         | 0    | 2         | 0         | 2    | 1         | 0         |
| Total                             | 20   | 22        | 40        | 14   | 18        | 20        | 19   | 12        | 19        |

DISCUSSION

Our results suggest that the described protocol combining low-dose CE-MRA and DSC perfusion imaging is feasible in the evaluation of patients with acute stroke, without the need for additional contrast. With the use of the described low-dose protocol, near isotropic sub-millimeter voxel sizes of neck and brain arteries were obtained with diagnostic accuracy equal or superior to TOF-MRA when compared with DSA. DSC perfusion imaging with reduced contrast dose was feasible at 3T with comparable qualitative results to a full-dose control group. Importantly, the presence of contrast in the circulating blood of the CE-MRA half-dose group did not negatively affect the image quality nor the quantitative analysis of perfusion data when compared with the control full-dose group.

MRA is used not only to detect the presence of intracranial arterio-occlusive disease but also to detect extracranial tandem stenosis to characterize the stroke mechanism. Despite its broad use in stroke protocols, TOF-MRA requires long acquisition times in the order of 8–11 minutes for the neck and brain arteries.

In addition, spin saturation and phase dispersion caused by slow, in-plane, turbulent, or complex flow can result in overestimation of arterial stenosis and occlusion and increase false-positive rates. It is not uncommon to notice the complete absence of flow signal in intracranial branches distal to a sub-occlusive thrombus or clot (Fig 1).

CE-MRA has not been widely incorporated into routine stroke protocols because of its relatively lower spatial resolution for evaluation of intracranial stenosis and need for extra contrast dose, which, until now, has been required in addition to the contrast used for DSC perfusion imaging. Introduction of multi-coil technology with inherent improved SNR and fast imaging techniques such as GRAPPA have significantly improved the speed and spatial resolution of CE-MRA while also permitting use of very low contrast doses.

With the use of the proposed CE-MRA protocol, sub-millimeter voxel sizes of cervicocranial arteries were obtained by use of 0.05 mmol/kg of gadolinium contrast. The specificity of CE-MRA was higher than TOF-MRA because of more frequent overestimation of arterial stenosis on TOF-MRA, predominantly caused by slow flow and phase dispersion in arterial segments at or distal to a high-grade stenosis or occlusion. Acquisition time required for MR stroke protocol including DWI, FLAIR, gradient refocused echo, and DSC perfusion is approximately 10 minutes with the use of CE-MRA as compared with approximately 20 minutes if TOF-MRA is used. Therefore, the use of CE-MRA instead of TOF-MRA can cut total scan time in half and result in a time savings of approximately 10 minutes.

Although the role of DSC perfusion in the acute stroke setting is still being defined, it is being used with increasing frequency in multiple prospective clinical trials and has become an important component of acute MR stroke protocols in large stroke cen-

Table 1: Distribution of arterial stenoses detected by TOF-MRA and CE-MRA in comparison with DSA (total arterial segments available for comparison = 410)

| Arterial Segment                  | TOF-MRA Stenosis Severity | CE-MRA Stenosis Severity | DSA Stenosis Severity |
|-----------------------------------|---------------------------|--------------------------|-----------------------|
|                                   | ≤50% | >50%, <99% | Occlusion | ≤50% | >50%, <99% | Occlusion | ≤50% | >50%, <99% | Occlusion |
| Common carotid artery             | 3    | 2         | 3         | 5    | 1         | 0         | 3    | 0         | 0         |
| Cervical ICA                      | 2    | 1         | 6         | 4    | 4         | 4         | 7    | 5         | 4         |
| Petrocerebral ICA                 | 2    | 3         | 4         | 1    | 1         | 1         | 0    | 2         | 1         |
| Supraclinoid ICA                  | 3    | 4         | 6         | 1    | 5         | 3         | 1    | 1         | 3         |
| MCA                               | 3    | 6         | 14        | 1    | 2         | 11        | 2    | 1         | 10        |
| Anterior cerebral artery          | 2    | 2         | 2         | 0    | 1         | 0         | 2    | 1         | 0         |
| Cervical vertebral artery         | 0    | 0         | 1         | 1    | 0         | 1         | 2    | 0         | 1         |
| Intradural vertebral artery       | 3    | 0         | 3         | 1    | 0         | 0         | 0    | 0         | 0         |
| Basilar artery                    | 2    | 1         | 0         | 0    | 0         | 0         | 0    | 0         | 0         |
| Posterior cerebral artery         | 0    | 3         | 1         | 0    | 2         | 0         | 2    | 1         | 0         |
| Total                             | 20   | 22        | 40        | 14   | 18        | 20        | 19   | 12        | 19        |

Table 2: Diagnostic performance and agreement of CE-MRA and TOF-MRA for arterial stenosis and occlusion in comparison with DSA

| Arterial Segment                  | No Disease | Mild Stenosis (≤50%) | High Grade (>50%, <99%) | Occlusion |
|-----------------------------------|------------|----------------------|-------------------------|-----------|
|                                   | DSA        | TOF-MRA              | CE-MRA                  | TOF-MRA   | CE-MRA | TOF-MRA |
| No disease                        | 353        | 324                  | 7                       | 17        | 0      | 14      | 0      | 5        |
| Mild stenosis (≤50%)              | 5          | 4                    | 7                       | 3         | 1      | 7       | 4      | 0        |
| High grade (>50%, <99%)           | 0          | 0                    | 0                       | 0         | 11     | 4       | 1      | 8        |
| Occlusion                         | 0          | 0                    | 0                       | 0         | 0      | 0       | 19     | 19       |

Table 3: Comparative quantitative analysis of DSC perfusion between full dose (0.1 mmol/kg) and half dose (0.05 mmol/kg)

| Image Parameter Evaluated | Full Dose | Half Dose | t Test |
|---------------------------|-----------|-----------|--------|
| SNR                       | 342 ± 101 | 330.3 ± 67 | 0.6    |
| Max T2* signal drop       | 120 ± 54  | 81.6 ± 48  | 0.01   |
| Background noise           | 2.5 ± 0.6 | 2.7 ± 0.9  | 0.4    |
| Peak of AIF               | 9.1 ± 2.4 | 7.2 ± 2.0  | 0.002  |
| FWHM of AIF               | 16.8 ± 2.6| 18.2 ± 2.8 | 0.14   |

Note: Data presented as mean ± standard deviation. Max indicates maximum.
CBV and CBF probably varied between the full-dose and half-noise contamination parameters calculated from deconvolution through additional results and performance of CE-MRA should be interpreted in this context. Second, the 50% cutoff used for hemodynamic significant arterial stenosis is somewhat arbitrary because there are no firm guidelines to determine the threshold of hemodynamic significance for the intracranial circulation. Third, the full-dose control group for DSC perfusion analysis was different from our patient population, with differences in underlying hemodynamic status, which can potentially complicate direct comparisons of image quality and quantitation. However, the limitation imposed by total contrast dose precludes the use of the same patients as their own control group. By matching the age, sex, and sequence parameters, we attempted to control for these potential confounding factors. Nevertheless, the comparison of image quality scores between 2 different groups should be interpreted with some caution. The choice of a longer TR (1.9 seconds) used in the current study is less ideal for DSC perfusion. This value was chosen to compensate for a possible SNR penalty that may result when reducing contrast dose. It is possible that this choice of TR could have resulted in undersampling of the AIF and therefore overestimation of perfusion parameters. Finally, the combination of parallel acquisition (GRAPPA) and contrast dose reduction may accentuate the effect of noise on image quality and DSC quantitative values, though higher SNR at 3T can help to mitigate this effect. Larger-scale clinical studies with concurrent use of different contrast dosage in the same individuals would help to validate our results.

CONCLUSIONS

Combining low-dose CE-MRA and DSC perfusion at 3T is feasible, with diagnostic image quality and without the need for additional contrast. The proposed protocol is significantly faster and more accurate than is the routine stroke protocol by use of TOF-MRA and may be adopted into routine clinical practice to streamline the treatment of patients with acute stroke. Our study demonstrates the clinical feasibility of performing DSC perfusion by use of a lower dose of contrast with qualitative measures comparable to a conventional dose group.

Disclosures: J. Paul Finn—UNRELATED: Grants/Grants Pending: Siemens Medical Solutions. Comments: Research grant only indirectly related to the work in this project; Payment for Lectures (including service on speakers bureau): Honorarium for lecture from Siemens Medical Solutions. David Liebeskind—UNRELATED: Consultancy: Covidien,* Stryker*; Grants/Grants Pending: NIH,* Comments: Research grants (*money paid to institution).

REFERENCES

1. Lloyd-Jones D, Adams RJ, Brown TM, et al. Heart disease and stroke statistics–2010 update: a report from the American Heart Association. Circulation 2010;121:e66–e215
2. Saver JL. Time is brain–quantified. Stroke 2006;37:263–66
3. Sohn CH, Sevick RJ, Frayne R. Contrast-enhanced MR angiography of the intracranial circulation. Magn Reson Imaging Clin N Am 2003;11:599–614
4. Bernstein MA, Huston J 3rd, Lin C, et al. High-resolution intracranial and cervical MRA at 3.0T: technical considerations and initial experience. Magn Reson Medicine 2001;46:955–62
5. Takano K, Utunomiya H, Ono H, et al. Dynamic contrast-enhanced subtraction MR angiography in intracranial vascular abnormalities. Eur Radiol 1999;9:1909–12
6. Tomassian A, Salamon N, Lohan DG, et al. Supraaortic arteries: contrast material dose reduction at 3.0-T high–spatial-resolution MR angiography–feasibility study. Radiology 2008;249:980–90
7. Nael K, Moriarty JM, Finn JP. Low dose CE-MRA. Eur J Radiol 2011;80:2–8
8. Heiland S, Reith W, Forsting M, et al. How do concentration and dosage of the contrast agent affect the signal change in perfusion-weighted magnetic resonance imaging? A computer simulation. Magn Reson Imaging 2001;19:813–20
9. Alger JR, Schaewe TJ, Lai TC, et al. Contrast agent dose effects in cerebral dynamic susceptibility contrast magnetic resonance perfusion imaging. J Magn Reson Imaging 2009;29:52–64
10. Manka C, Traber F, Gieseke J, et al. Three-dimensional dynamic susceptibility-weighted perfusion MR imaging at 3.0 T: feasibility and contrast agent dose. Radiology 2005;234:869–77
11. Alger JR, Schaewe TJ, Liebeskind DS, et al. On the feasibility of reduced dose dynamic susceptibility contrast perfusion MRI for stroke. Paper presented at International Society of Magnetic Resonance Medicine, Montréal, Québec, Canada, May 7–13, 2011
12. Griswold MA, Jakob PM, Heidemann RM, et al. Generalized auto-calibrating partially parallel acquisitions (GRAPPA). Magn Reson Med 2002;47:1202–10
13. Habibi R, Krishnam MS, Lohan DG, et al. High-spatial-resolution lower extremity MR angiography at 3.0 T: contrast agent dose comparison study. Radiology 2008;248:680–92
14. Wu O, Ostergaard L, Weisskoff RM, et al. Tracer arrival timing-insensitive technique for estimating flow in MR perfusion-weighted imaging using singular value decomposition with a block-circulant deconvolution matrix. Magn Reson Med 2003;50:164–74
15. Lin W, Tkach JA, Haacke EM, et al. Intracranial MR angiography: application of magnetization transfer contrast and fat saturation to short gradient-echo, velocity-compensated sequences. Radiology 1993;186:753–61
16. Isoda H, Takehara Y, Isogai S, et al. MRA of intracranial aneurysm models: a comparison of contrast-enhanced three-dimensional MRA with time-of-flight MRA. J Comput Assist Tomogr 2000;24:308–15
17. Nael K, Villablanca JP, Pope WB, et al. Supraaortic arteries: contrast-enhanced MR angiography at 3.0 T–highly accelerated parallel acquisition for improved spatial resolution over an extended field of view. Radiology 2007;242:600–09
18. Phan T, Huston J 3rd, Bernstein MA, et al. Contrast-enhanced magnetic resonance angiography of the cervical vessels: experience with 422 patients. Stroke 2001;32:2282–86
19. Butcher KS, Parsons M, MacGregor L, et al. Refining the perfusion-diffusion mismatch hypothesis. Stroke 2005;36:1153–59
20. Sobesky J, Zaro Weber O, Lehnhardt FG, et al. Does the mismatch match the penumbra? Magnetic resonance imaging and positron emission tomography in early ischemic stroke. Stroke 2005;36:980–85
21. Carpenter TK, Armitage PA, Bastin ME, et al. DSC perfusion MRI: quantification and reduction of systematic errors arising in areas of reduced cerebral blood flow. Magn Reson Med 2006;55:1342–49