

Gadolinium Enhancement in Intracranial Atherosclerotic Plaque and Ischemic Stroke: A Systematic Review and Meta-Analysis

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Background—Gadolinium enhancement on high-resolution magnetic resonance imaging (MRI) has been proposed as a marker of inflammation and instability in intracranial atherosclerotic plaque. We performed a systematic review and meta-analysis to summarize the association between intracranial atherosclerotic plaque enhancement and acute ischemic stroke.

Methods and Results—We searched the medical literature to identify studies of patients undergoing intracranial vessel wall MRI for evaluation of intracranial atherosclerotic plaque. We recorded study data and assessed study quality, with disagreements in data extraction resolved by a third reader. A random-effects odds ratio was used to assess whether, in any given patient, cerebral infarction was more likely in the vascular territory supplied by an artery with MRI-detected plaque enhancement as compared to territory supplied by an artery without enhancement. We calculated between-study heterogeneity using the Cochrane Q test and publication bias using the Begg-Mazumdar test. Eight articles published between 2011 and 2015 met inclusion criteria. These studies provided information about plaque enhancement characteristics from 295 arteries in 330 patients. We found a significant positive relationship between MRI enhancement and cerebral infarction in the same vascular territory, with a random effects odds ratio of 10.8 (95% CI 4.1–28.1, $P < 0.001$). No significant heterogeneity ($Q = 11.08$, $P = 0.14$) or publication bias ($P = 0.80$) was present.

Conclusions—Intracranial plaque enhancement on high-resolution vessel wall MRI is strongly associated with ischemic stroke. Evaluation for plaque enhancement on MRI may be a useful test to improve diagnostic yield in patients with ischemic strokes of undetermined etiology. (*J Am Heart Assoc.* 2016;5:e003816 doi: 10.1161/JAHA.116.003816)

Key Words: cerebral infarction • culprit artery • enhancement gadolinium • ischemic stroke • magnetic resonance imaging • plaque • vessel wall imaging

Intracranial atherosclerosis is one of the most common causes of ischemic stroke worldwide^{1,2} and is associated with a high rate of recurrence.³ The most commonly used

imaging techniques to assess intracranial atherosclerosis, such as computed tomographic angiography or magnetic resonance angiography, provide information on the degree of narrowing of the vascular lumen. Most classification schemes for ischemic stroke etiology require plaque to cause $\geq 50\%$ stenosis for a given stroke to be attributable to large-artery atherosclerosis.⁴ However, magnetic resonance imaging (MRI) studies of the extracranial carotid arteries suggest that many atherosclerotic plaques have high-risk features despite the absence of significant luminal narrowing.^{5,6} It is unknown to what extent similar nonstenosing *intracranial* atherosclerotic plaque might be responsible for a proportion of the approximately 1 in 3 ischemic strokes for which no clear etiology can be determined.⁷

Recent investigations have begun to address this problem by leveraging high-resolution, multiplanar MRI to detect high-risk abnormalities of the intracranial vessel walls. Previous studies in both the coronary and extracranial carotid arteries have shown that abnormal plaque enhancement after the administration of gadolinium contrast agent is a marker of

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An accompanying Data S1 is available at <http://jaha.ahajournals.org/content/5/8/e003816/DC1/embed/inline-supplementary-material-1.pdf>

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Received April 28, 2016; accepted July 19, 2016.

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inflammation, neovascularity, and plaque instability.^{8,9} For this reason, plaque enhancement has been recently studied as a potential high-risk plaque feature in the intracranial circulation (Figure 1). Plaque enhancement is a particularly attractive MRI biomarker because it can be rapidly detected, qualitatively assessed, and does not require significant image postprocessing to analyze. While research to date on the detection of nonstenosing intracranial atherosclerotic lesions based on enhancement characteristics has been promising, individual studies have been small, making it difficult to draw firm conclusions about this emerging technique. We therefore performed a systematic review and meta-analysis to evaluate the association between abnormal plaque enhancement on high-resolution MRI and acute ischemic stroke.

Methods

We performed this study following the Meta-Analysis of Observational Studies in Epidemiology (MOOSE) group guidelines¹⁰ and the Preferred Reporting Items for Systematic Reviews and Meta-Analyses statement.¹¹ Because this was an

analysis of previously published data, this study did not undergo or require Institutional Review Board approval.

Data Sources and Searches

Two research librarians performed comprehensive searches of Ovid MEDLINE, Ovid Embase, and the Cochrane Library from database inception to December 21, 2015. We first conducted our search in Ovid MEDLINE. We then adapted headings and keywords for other databases and identified additional records by employing the “Cited by” and “View references” features in Scopus (see Data S1 for search methodology details).

Study Selection

We included studies evaluating the association of abnormal plaque enhancement on vessel wall MRI with recent ischemic stroke. Specific inclusion criteria were as follows: (1) studies of patients with acute cerebral infarction; (2) studies of patients who underwent MRI of the intracranial vessels with assessment of plaque signal abnormalities within 30 days of

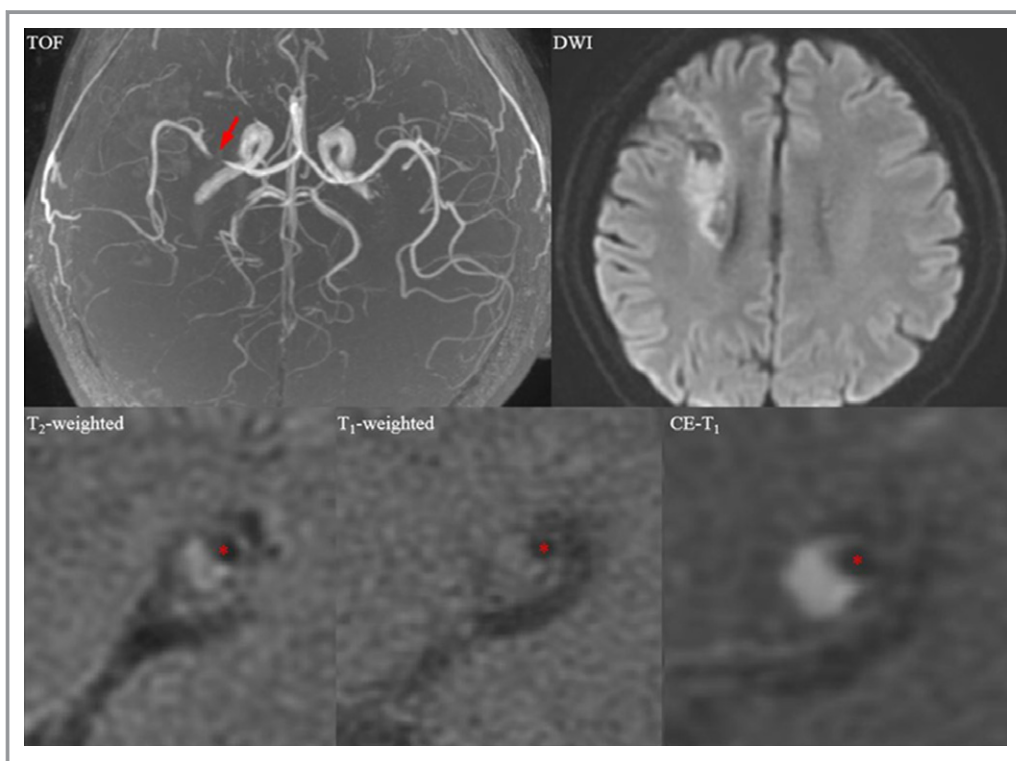


Figure 1. Magnetic resonance images showing an enhancing atherosclerotic plaque in the right middle cerebral artery in a symptomatic patient who had suffered a recent right cerebral hemispheric acute ischemic stroke. A mixture of acute and chronic infarction involving the right periventricular and frontal subcortical regions is seen in the diffusion-weighted image (DWI); the plaque is shown by an arrow in the time-of-flight (TOF) image, and a cross section at the most stenotic site is shown in T₂, T₁, and contrast-enhanced (CE) T₁ images (red asterisks: lumen).

the ischemic stroke; (3) specific assessment of the presence or absence of plaque postgadolinium enhancement; (4) reporting of the prevalence of plaque enhancement in the arteries directly supplying blood to the territory of the infarction compared to the prevalence of enhancement in an unrelated vascular territory (such as contralateral hemisphere for an anterior circulation infarction); and (5) studies with ≥ 10 subjects to avoid the inclusion of case reports or small case series. We only included peer-reviewed journal articles rather than conference proceedings or abstracts. We did this to ensure that the studies included in our analysis provided sufficient information to allow for the collection of patient characteristics, MRI study protocols, and MRI test results in a fashion that would allow us to perform a detailed systematic review and meta-analysis. In otherwise eligible studies, we excluded patients with infarctions that were likely secondary to nonatherosclerotic etiologies such as vasculitis or reversible cerebral vasoconstriction syndrome. We also did not include patient data for chronic ischemic strokes or transient ischemic attacks. If based on review of study inclusion dates, authors appeared to have published data from a single cohort or medical center more than once, the single article with the largest sample size was included to minimize analysis of duplicate or overlapping samples. When necessary we attempted to contact the corresponding author for additional details to clarify our data extraction.

Data Extraction and Quality Assessment

A single investigator read the title and abstract of all references produced by our database search. After preliminary articles were shortlisted as potentially eligible, 2 readers read the articles in their entirety to determine eligibility, with disagreements resolved by consensus. We extracted data in duplicate using a prespecified data collection template. A third tie-breaking reader resolved disagreements in data extraction. We extracted the following study characteristics: first author; study design (prospective or not); major study inclusion criteria; country of the study; total number of subjects; basic study demographics and the prevalence of stroke risk factors in the studied populations, including age, sex, hypertension, diabetes, atrial fibrillation, coronary artery disease, hyperlipidemia, and smoking history; definitions of ischemic stroke and delineation of vascular territories; and specific MRI techniques employed, including definitions of abnormal plaque enhancement. We classified strokes in each of 2 groups: (1) all strokes in the vascular territory of an artery with plaque enhancement and (2) all strokes in the vascular territory of arteries free of plaque enhancement. This allowed for the calculation of a pooled odds ratio (OR) comparing the prevalence of infarction in tissue supplied by an artery with enhancing plaque versus tissue supplied by unaffected

arteries, with each patient serving as his or her own control; an OR >1 would indicate an association between plaque enhancement and infarction.

We adapted risk of bias assessments in previously published meta-analyses of MRI biomarkers of stroke risk^{12,13} and generated 8 specific questions to evaluate for potential selection, detection, reporting, and confounding bias (see Data S1). Two readers assessed for risks of bias using this questionnaire, with disagreements in assessment resolved by a third tie-breaking evaluator.

Data Synthesis and Analysis

We performed a meta-analysis of the individual study ORs (ie, odds of the strength of association between enhancement and ipsilateral infarction) using R package “meta” (version 4.3-2). We also performed a sensitivity analysis limited to prospective studies. We pooled ORs using a random-effects (DerSimonian-Laird) model and generated a forest plot to display the individual study ORs and the pooled ORs. We used a random-effects model because we conservatively assumed that individual studies did not have the same effect size given the high possibility of between-study heterogeneity in terms of sample size, subject characteristics, and imaging methods. The random-effects analysis allows for more variability in the individual study OR estimates when generating the pooled OR. A continuity correction of 0.5 was applied to studies with zero cell frequencies. To assess the combinability of the ORs, we calculated the *P*-value from the Cochrane Q statistical heterogeneity test. For each meta-analysis, the presence of publication bias was evaluated through a funnel plot. The Begg-Mazumdar rank-correlation test was used to statistically assess the presence of publication bias. All *P* < 0.05 were considered statistically significant.

Results

Study Selection and Characteristics

We screened a total of 4437 titles and abstracts from which we identified 8 articles^{14–21} that met all inclusion criteria for the systematic review. Study selection steps are summarized in Figure 2. In total, the 8 studies included a combined 330 individual subjects in whom data from 295 atherosclerotic plaques provided data eligible for meta-analysis. Of the 8 articles meeting inclusion criteria (Table 1), 5^{14,18–21} were prospective cross-sectional studies and 3^{15–17} were retrospective cross-sectional studies. Three studies were conducted in China,^{19–21} 2 in the United States,^{17,18} and 1 each in Canada,¹⁶ the Netherlands,¹⁴ and South Korea.¹⁵ A preponderance of men were studied in 7 of the 8 included studies, with the range of percent of men in each study

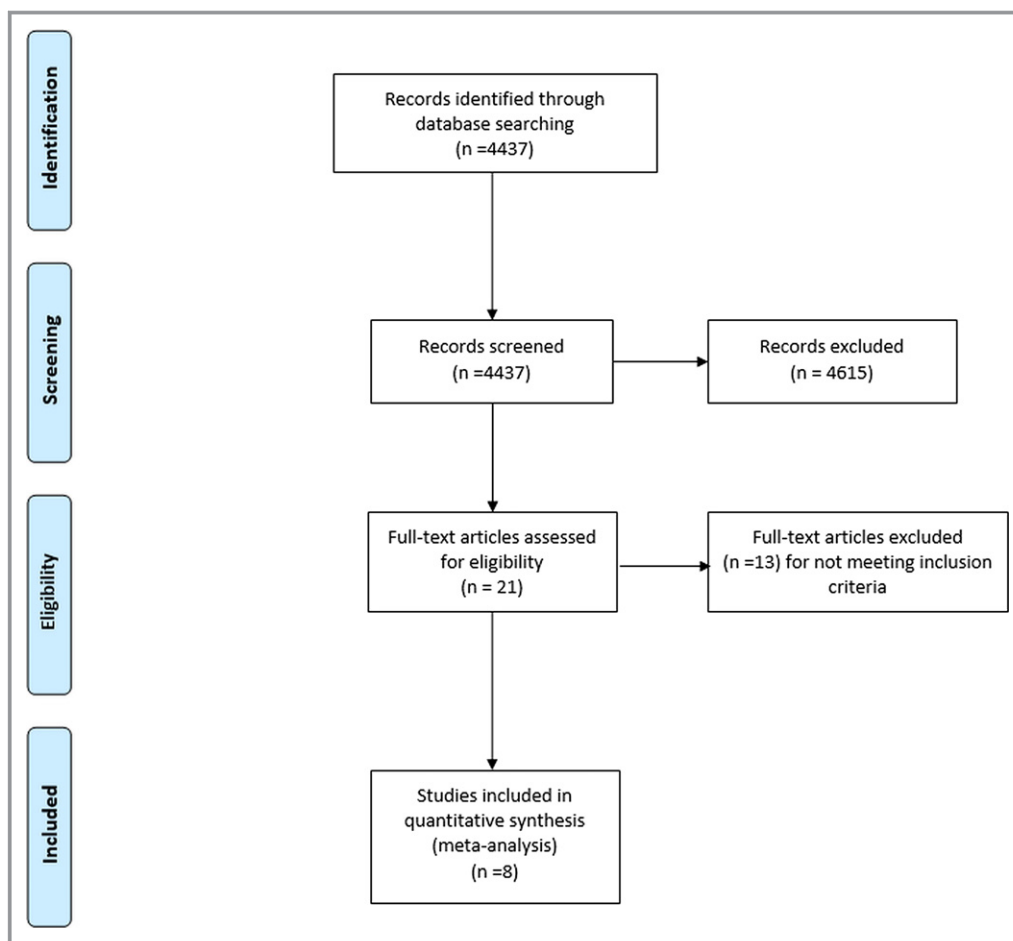


Figure 2. Study selection flow diagram.

ranging from 46.9% to 88.9%. All studies had a mean age above 50 years (range 54.6–68.7 years). There were differences in the degree of intracranial luminal stenosis measurements required for patients to be included in the individual studies, with some studies^{17,18,20} requiring $\geq 50\%$ or $\geq 70\%$ stenosis while others focused on patients without significant stenosis (ie, $< 50\%$).^{19,21} Though some studies included patients with transient ischemic attacks or nonacute ischemic strokes, all included studies provided adequate information to collect MRI enhancement data on the subset of relevant patients with acute ischemic stroke occurring within 30 days of imaging.

Definitions of Abnormal Plaque Enhancement and Delineation of Vascular Territories

Evaluation for abnormal plaque enhancement on all studies occurred on postcontrast T1-weighted sequences, with most studies using a high-resolution technique capable of submillimeter resolution (Table 2). Six studies^{15,16,18–21} were performed on 3.0-T MRI scanners, 1¹⁴ on a 7.0-T scanner, and

1¹⁷ on a 1.5-T scanner. All but 1 study¹⁶ involved more than 1 reader evaluating vessel wall MRI for abnormal vessel wall enhancement. Though there were some between-study differences in the definition of abnormal plaque enhancement, abnormal plaque was most commonly defined as enhancement judged to be the same or greater than the degree of physiologic enhancement present in the adjacent pituitary gland. Similarly, though specific definitions varied, all studies explicitly differentiated acute neuroimaging-confirmed infarctions occurring in the vascular territory of an artery with enhancing plaque, versus infarctions occurring in the vascular territory of an artery without enhancing plaque. All studies evaluated for enhancement in either the intracranial internal carotid artery or the middle cerebral artery.

Association Between Plaque Enhancement and Cerebral Infarction

We were able to obtain sufficient raw data to calculate a pooled OR expressing the strength of association between MRI enhancement and cerebral infarction in all of our included

Table 1. Study Characteristics

Study Number	Study First Author and Year	Study Design	Major Inclusion Criteria	Country	Total Number of Subjects	Mean Age	Men, no. (% male)	Hypertension (%)	Diabetes Mellitus (%)	Atrial Fibrillation (%)	Coronary Artery Disease (%)	Hyperlipidemia (%)	Smoking History (%)
1	van der Kolk 2011 ¹⁴	Prospective	Patients admitted with ischemic infarct or TIA involving the anterior cerebral circulation; Healthy volunteers were also included	Netherlands	32	59 years (range: 26–83 years)	18 (51%)	N/A	N/A	N/A	N/A	N/A	N/A
2	Kim 2012 ¹⁵	Retrospective	Patients admitted with infarction involving the MCA territory or TIA due to MCA disease	South Korea	34	64.9 years (SD: ±11.22)	21 (62%)	70.60%	38.00%	N/A	23.50%	44.10%	N/A
3	Skarpathiotakis 2013 ¹⁶	Retrospective	Patients with documented prior ischemic stroke based on DWI and ≥1 intracranial atherosclerotic plaque on MRI	Canada	29	61.2 years (range: 35–84)	18 (62%)	N/A	N/A	N/A	N/A	N/A	N/A
4	Vakil 2013 ¹⁷	Retrospective	Patients with severe intracranial atherosclerosis causing ≥70% stenosis with detectable atherosclerotic plaque on MRI	United States (Illinois)	19	68.7 years (SD: ±9.6)	13 (68%)	89.50%	47.40%	0%	N/A	84.20%	10.5% (active)
5	Qiao 2014 ¹⁸	Prospective	Patients with: (1) Intracranial stenosis ≥50% in a large intracranial artery; (2) TIA or stroke in the distribution of the narrowed vessel	United States (Baltimore, MD)	27	56.8 years (SD: ±12.4)	19 (70%)	81%	33%	N/A	N/A	59%	15% (active)

Continued

Table 1. Continued

Study Number	Study First Author and Year	Study Design	Major Inclusion Criteria	Country	Total Number of Subjects	Mean Age	Men, no. (% male)	Hypertension (%)	Diabetes Mellitus (%)	Atrial Fibrillation (%)	Coronary Artery Disease (%)	Hyperlipidemia (%)	Smoking History (%)
6	Teng 2016 ¹⁹	Prospective	(1) Absence of significant carotid arterial stenosis (<30%); (2) absence of atrial fibrillation; (3) absence of ascending aortic arch atheroma; and (4) ≥ 1 atherosclerotic risk-factors	Shanghai, China	139	57.1 years	90 (64.7%)	71.20%	34.50%	N/A	9.40%	Pre-admission statin: 32.3%	29%
7	Xu 2015 ²⁰	Prospective	Unilateral MCA stenosis confirmed by MRI; age >50 years; $\geq 50\%$ MCA stenosis; absence of nonatherosclerotic cerebrovascular disease	Xuzhou, China	32	65.8 years (SD: ± 13.3)	15 (46.9%)	78.10%	28.10%	N/A	37.50%	N/A	34.40%
8	Zou 2015 ²¹	Prospective	(1) Recent single infarction confirmed by DWI; (2) no ipsilateral MCA stenosis based on MRI; (3) one or more risk factors for atherosclerosis	Beijing, China	18	54.6 years (range: 40–70)	16 (88.9%)	66.70%	22.20%	N/A	N/A	55.60%	72.20%

DWI indicates diffusion-weighted imaging; MCA, middle cerebral artery; N/A, data not available; MRI, magnetic resonance imaging; TIA, transient ischemic attack.

Table 2. MRI Protocols and Study Definitions

Study Number	Study First Author and Year	MRI Field Strength	MRI Scanner Platform	Coil Type Used	T1-Weighted Imaging Sequence Parameters	Field of View	Matrix	Acquired Resolution	Contrast Agent Administered	Number of MRI Readers	Definitions of Abnormal Vessel Wall Enhancement	Vessel Site Evaluated for Enhancement	Culprit and Non-Culprit Plaque Definition
1	van der Kolk 2011 ¹⁴	7.0-T	Philips Healthcare	16-channel receive coil	Magnetization preparation inversion recovery turbo spin-echo sequence; TR/TE 6050/23, inversion time 1770 ms	220×180 × 13 mm ³	N/A	0.8×0.8 ×0.8 mm ³	0.1 mL/kg of gadobutrol	2 with consensus	After registering pre- and postcontrast scans, the images were subtracted and directly compared to determine enhancing areas and concomitant vessel wall anatomy. The infundibulum was used to assess whether normal contrast enhancement had taken place	Intracranial internal carotid artery	Culprit (defined during data extraction for meta-analysis); Vessel immediately upstream of a stroke of undetermined etiology or stroke attributable to ipsilateral large vessel atherosclerosis; Non-culprit plaque: vessels studied in the intracranial circulation but not meeting definition of culprit plaque
2	Kim 2012 ¹⁵	3.0-T	Conventional MRI: Sigma, GE Medical Systems; High Resolution-MRI: Veiro, Siemens Healthcare	8-channel head coil	T1-weighted: TR/TE 600/12	120×120 mm	384×269	N/A	Gadolinium	2 with a 3rd reader available to resolve disagreements	Atheromatous plaque: Eccentric or focal signal intensity within the vessel lumen on MRI. Abnormal enhancement: enhancement on T1-enhanced images. Vulnerable symptomatic plaque: eccentric or focal signal intensity within vessel lumen on MRI accompanied by intraplaque heterogeneous signal intensity on T1 and T2 images and plaque enhancement on T1-enhanced images. Stable symptomatic plaque: eccentric or focal signal intensity within vessel lumen on MRI in the absence of the features described earlier in vulnerable symptomatic plaque. Culprit plaque defined as plaques upstream of ipsilateral ischemic infarctions	Middle cerebral artery	Vulnerable symptomatic plaque: eccentric or focal signal intensity within vessel lumen on MRI accompanied by intraplaque heterogeneous signal intensity on T1 and T2 images and plaque enhancement on T1-enhanced images. Stable symptomatic plaque: eccentric or focal signal intensity within vessel lumen on MRI in the absence of the features described earlier in vulnerable symptomatic plaque. Culprit plaque defined as plaques upstream of ipsilateral ischemic infarctions
3	Skampathidakis 2013 ¹⁶	3.0-T	HDX platform, GE healthcare	8-channel head coil	T1 FLAIR TR/TE 2108/12, inversion time 860 ms	16×22 cm	N/A	384×384	7.5-mL gadobutrol	1	Enhancement similar to pituitary parenchyma—strong; less than pituitary parenchyma—mild; no change from precontrast images—absent	Middle cerebral artery	Number of atherosclerotic plaques were tabulated and divided into plaques in stroke territory (culprit) and nonstroke territory (non-culprit) vasculature

Continued

Table 2. Continued

Study Number	Study First Author and Year	MRI Field Strength	MRI Scanner Platform	Coil Type Used	T1-Weighted Imaging Sequence Parameters	Field of View	Matrix	Acquired Resolution	Contrast Agent Administered	Number of MRI Readers	Definitions of Abnormal Vessel Wall Enhancement	Vessel Site Evaluated for Enhancement	Culprit and Non-Culprit Plaque Definition
4	Vakil 2013 ¹⁷	1.5-T	Avanto/Espree, Siemens	Receive-only head coil	T1-weighted spin-echo; TR/TE 663/15	220 to 240 mm	256×256	N/A	0.1 mmol/kg of gadopentetate dimeglumine	2 with a 3rd reader available to resolve disagreements	Qualitative assessment: 1—definite non-enhancement, 2—suspected non-enhancement, 3—uncertain, 4—suspected enhancement, 5—definite enhancement. A plaque with an average >3-enhancing, while plaque with an average <3—nonenhancing. Plaque enhancement was quantified as the relative increase in lesion T1-signal postcontrast: mean T1 signal intensity in the outer wall area on postcontrast divided by its precontrast signal	Supracarotid internal carotid artery	A symptomatic intracranial atherosclerotic plaque (culprit) was defined as presenting with acute neurologic symptoms and/or restricted diffusion weighted abnormalities corresponding to the vascular distribution of the intracranial stenosis. Non-culprit plaques were asymptomatic plaques
5	Qiao 2014 ¹⁸	3.0-T	Achieva and Vista, Philips Healthcare	Body coil for transmission and 8-channel head coil for reception	3D black blood T1: TR/TE 2000/38	180×180×40 mm	450×450×100	0.4×0.4×0.4 mm	0.1 mmol/kg of gadopentetate dimeglumine	1 reader for plaque identification and 2 for assessment of enhancement	Qualitative 3-point scale: grade 0 enhancement similar to or less than normal vessels in the same individual; grade 1: enhancement was greater than that of grade 0 but less than that of the pituitary infundibulum; grade 2: enhancement was similar to or greater than that of the infundibulum	Cavernous and supracarotid internal carotid artery	1. Culprit: (a) the only lesion within the vascular territory of the stroke; (b) the most stenotic lesion when multiple plaques were present within the same vascular territory of the stroke. 2. Probably culprit: when the plaque was not the most stenotic lesion within the same vascular territory of the stroke. 3. Nonculprit: when the plaque was not within the vascular territory of the stroke

Continued

Table 2. Continued

Study Number	Study First Author and Year	MRI Field Strength	MRI Scanner Platform	Coil Type Used	T1-Weighted Imaging Sequence Parameters	Field of View	Matrix	Acquired Resolution	Contrast Agent Administered	Number of MRI Readers	Definitions of Abnormal Vessel Wall Enhancement	Vessel Site Evaluated for Enhancement	Culprit and Non-Culprit Plaque Definition
6	Teng 2015 ¹⁹	3.0-T	HDX, GE Healthcare	8-channel phased array brain coil	T1-weighted: TR/TE 567/16	10×10 cm ²	320×256	N/A	0.2 mmol/kg gadopentetic acid	3 (2 residents, 1 neuroradiologist) with consensus	Average signal intensity enclosed between the lumen and outer wall normalized to adjacent gray matter. If normalized value >1.0=plaque enhancement	MCA	Culprit plaque: a lesion arising on the ipsilateral side to an ischemic stroke on neuro-imaging with accompanying clinical symptoms. Non-culprit plaque: either a plaque occurring in a contralateral artery of a symptomatic patient or one in asymptomatic controls
7	Xu 2015 ²⁰	3.0-T	Discovery MP750, GE Healthcare	16-channel phase array coil	T1-weighted: TR/TE 600/12 ms	12×12 cm	256×256	N/A	0.1 mmol/kg of gadopentetate dimeglumine	2 with consensus	Enhancement between the outer wall boundary and vessel lumen on postcontrast images	Middle cerebral artery	Culprit: Atherosclerotic plaques upstream and ipsilateral to diffusion-weighted imaging hyperintensity in the same vascular territory. All other plaques studied considered non-culprit
8	Zou 2015 ²¹	3.0-T	Magnetom Verio, Siemens	32-channel head coil	T1-weighted SPACE: TR/TE 938/24 ms	N/A	N/A	0.5 to 0.7 mm ³	0.1 mmol/kg of gadopentetate dimeglumine	2 with consensus	Mean signal intensities of the MCA vessel wall on registered pre- and post-contrast T1w-SPACE were measured and normalized to gray matter. Signal intensity change ≥20% was defined as plaque enhancement	Middle cerebral artery	Culprit plaque: a lesion arising on the ipsilateral side to an ischemic stroke confirmed on brain MRI; Non-culprit plaque: plaque occurring in a contralateral artery of a symptomatic patient

MCA indicates middle cerebral artery; MRI, magnetic resonance imaging; N/A, data not available; SPACE, sampling perfection with application optimized contrasts using different flip-angle evolution; T, Tesla; TE, time to echo; TR, time to repetition.

Table 3. MRI Test Results

Study Number	Study First Author and Year	Total Number of Suspected Culprit Plaques Causing Downstream Ischemic Stroke Only Imaged in Acute Phase	Number of Enhancing Suspected Culprit Plaques Causing Downstream Ischemic Stroke Imaged in Acute Phase	Total Number of Non-Culprit Plaques	Number of Enhancing Non-Culprit Plaques
1	van der Kolk 2011 ¹⁴	5	3	12	1
2	Kim 2012 ¹⁵	21	14*	8 [†]	0
3	Skarpathiotakis 2013 ^{‡16}	13	13	2	0
4	Vakil 2013 ¹⁷	7	4	3	0
5	Qiao 2014 ¹⁸	21	19	45	10
6	Teng 2015 ^{§19}	82	41	27	7
7	Xu 2015 ²⁰	15	12	17	4
8	Zou 2015 ²¹	12	11	5	4

MRI indicates magnetic resonance imaging.

*Data obtained via direct author correspondence.

[†]The nonculprit group had no evidence of any significant middle cerebral artery (MCA) plaque (and no abnormal wall or plaque enhancement).

[‡]Three patients had multiple plaques in the same territory for which individual plaque-level data were not available.

[§]The nonculprit group included plaques in separately recruited asymptomatic subjects.

^{||}Authors provided enhancement data specifically for superiorly situated MCA plaques ipsilateral and contralateral to infarctions.

8 studies (Table 3). In this analysis, information about plaque enhancement characteristics was available from 295 atherosclerotic lesions, 143 of which supplied a vascular territory containing an acute infarction and 152 of which supplied an infarct-free vascular territory. We found a significant association between MRI enhancement of an artery and stroke within the vascular territory of that same artery, with a random effects OR of 10.8 (95% CI 4.1–28.1, $P < 0.001$, Figure 3). These results were robust to a sensitivity analysis excluding the 2 retrospective studies (OR 8.4, 95% CI 3.1–22.8, $P < 0.001$). There was neither statistically significant

heterogeneity ($Q = 11.08$, $P = 0.14$) nor significant publication bias (Begg-Mazumdar test for publication bias $P = 0.80$, Figure 4) present in this analysis.

Assessment of the Quality of the Included Studies

The results from the quality assessment questionnaire are shown in Table 4. All studies provided detailed description of study inclusion/exclusion criteria and 6 of 8 studies^{14,17–21} were prospectively conducted. Seven^{15–21} of 8 studies had as

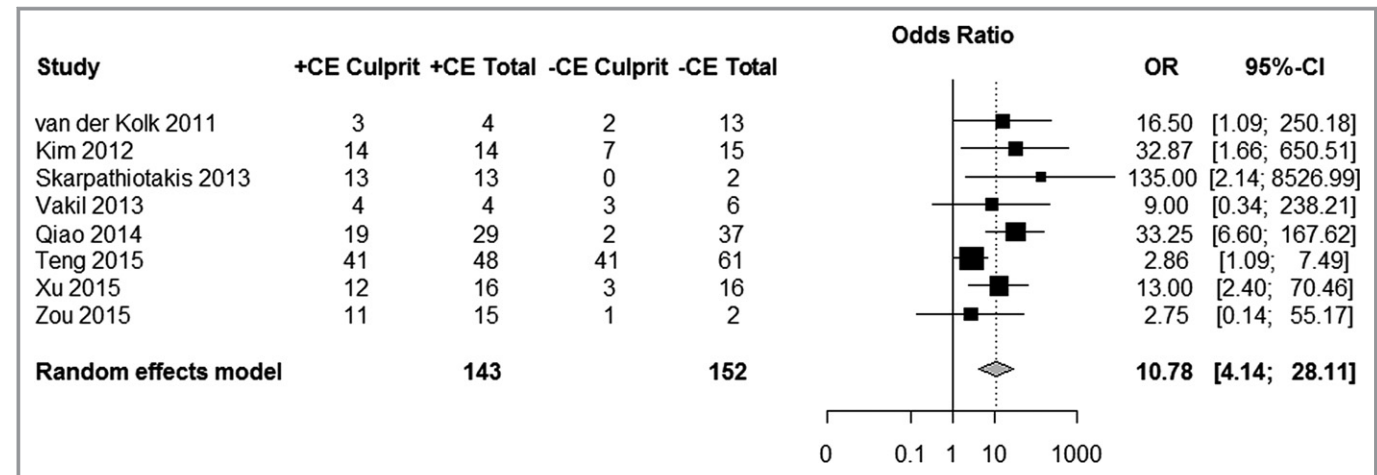


Figure 3. Forest plot of the association between magnetic resonance imaging–determined plaque contrast enhancement (CE) and acute ischemic stroke. The meta-analysis was calculated using a random-effects model, with the pooled odds ratio (OR) shown in the forest plot. Each square represents the point estimate of any given study’s effect size. The size of the squares is proportional to the inverse of the variance of the estimate, while the horizontal lines represent each study’s 95% CIs. The diamond represents the pooled estimate with the width of the diamond representing the pooled 95% CI.

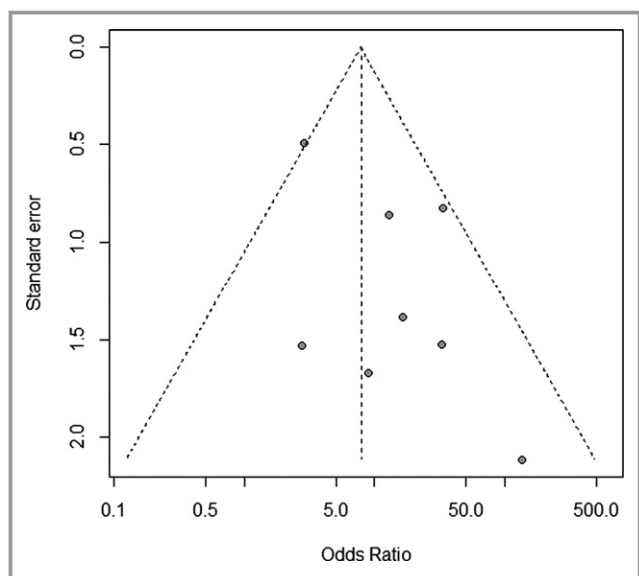


Figure 4. Funnel plot to evaluate for publication bias. Individual study effect sizes expressed as odds ratios are shown on the x -axis and each study's standard error is shown on the y -axis. Larger and more precise studies are plotted at the top, near the combined (pooled) odds ratio, whereas smaller and less precise studies will show a wider distribution below. If there is no publication bias, the studies would be expected to be symmetrically distributed on both sides of the pooled odds ratio line. In the case of publication bias, the funnel plot may be asymmetrical, since the absence of studies would distort the distribution on the scatter plot.

a primary objective to determine whether enhancement was associated with acute ischemic change in the brain. Similarly, 7^{15–21} of 8 studies had investigators blinded to the location of infarction during vessel wall enhancement detection and utilized more than 1 reader to evaluate for plaque enhancement, though only 2 studies^{17,18} reported interreader reproducibility measures. The classification of strokes as occurring in the territory of enhancing arterial plaque versus in territory without enhancing plaque occurred by an adjudicated panel of more than 1 investigator in only 1 study.¹⁸ Finally, no studies consistently matched enhancing plaques leading to downstream infarction versus plaque in infarct-free territory in regard to stenosis severity.

Discussion

Increasing the diagnostic confidence that a specific atherosclerotic lesion is the cause of a patient's ischemic stroke is important because it can guide optimal secondary stroke prevention measures. By characterizing plaque characteristics beyond luminal stenosis, MRI of the vessel wall is a promising technique to detect specific plaque features associated with acute ischemic infarction. In this systematic

review and meta-analysis we studied nearly 300 plaques imaged with vessel wall MRI pooled together from 8 individual studies, and focused on 1 such high-risk atherosclerotic plaque feature, enhancement after gadolinium administration. We found that plaque enhancement is strongly associated with downstream acute infarction. Infarction was 10 times more likely in tissue supplied by an enhancing artery than in tissue supplied by nonenhancing arteries.

The mechanisms of plaque enhancement in vulnerable intracranial atherosclerotic plaques are likely complex and multifactorial. Much of our understanding of the pathophysiology of gadolinium enhancement in the cerebral arterial vessel wall arises from literature on the extracranial carotid arteries where endarterectomy specimens are more readily accessible for pathological correlation. In these studies, carotid plaque gadolinium enhancement has been shown to spatially correlate with histological markers of vessel wall neovascularization and inflammation, both well-known markers of unstable atherosclerotic plaque.⁹ Enhancement detectable on MRI may be related to endothelial dysfunction present in the diseased intraplaque microvasculature of atherosclerotic vessels.²² Such compromised microvascular endothelium likely results in the vascular leakage needed for gadolinium to accumulate in the perivascular spaces and become detectable on T1-weighted MRI sequences. However, it is important to note that unlike the extracranial carotid arteries, most intracranial vessels do not have a vasa vasorum, though there is evidence that arteriosclerosis can promote the development of vasa vasorum in the proximal intracranial vasculature.^{23,24} As a result, though it is likely that a leaky endothelial barrier due to inflammation is responsible for the gadolinium enhancement seen in the studies included in this meta-analysis, the specific contribution of vasa vasorum to this enhancement requires further histopathological evaluation.

Understanding the time course of intracranial plaque enhancement after an acute ischemic event would help optimize the timing and diagnostic yield of intracranial vessel wall MRI in patients presenting with ischemic stroke. One study¹⁶ in this meta-analysis also focused on the longitudinal pattern of plaque enhancement after ischemic stroke and found that the strength of enhancement was its highest in the acute stroke phase (within 4 weeks), and that enhancement steadily decreased as the infarct became chronic in age (after 12 weeks). Though there are limited data suggesting the plaque enhancement may also predict future stroke recurrence,²⁵ further work is necessary to more fully elucidate the temporal relationships between MRI plaque enhancement, prior ischemic stroke, and future ischemic stroke risk.

We focused our analysis on gadolinium enhancement rather than other specific proposed high-risk plaque elements such as intraplaque hemorrhage, fibrous cap abnormalities, or lipid-rich necrotic core because of the paucity of studies

Table 4. Risk of Bias Question Results

Question	Answers	van der Kolk 2011 ¹⁴	Kim 2012 ¹⁵	Skarpathiotakis 2013 ¹⁶	Vakil 2013 ¹⁷	Qiao 2014 ¹⁸	Teng 2015 ¹⁹	Xu 2015 ²⁰	Zou 2015 ²¹
Was the study sample prospectively selected to minimize the risk of selection bias?	Yes (+) or no (-)	+	+	-	-	+	+	+	+
Were the inclusion and exclusion criteria adequately described?	Yes (+) or no (-)	+	+	+	+	+	+	+	+
Was the study's primary objective to assess whether enhancement was associated with ischemic presentations?	Yes (+) or no (-)	-	+	+	+	+	+	+	+
Were the investigators blinded to the location of infarction during vessel wall enhancement detection?	Yes (+) or no (-)	-	+	+	+	+	+	+	+
Did more than 1 investigator assess for the presence of vessel enhancement?	Yes (+) or no (-)	+	+	-	+	+	+	+	+
Was a measure of interreader reproducibility for enhancement detection reported?	Yes (+) or no (-)	-	-	-	+	+	-	-	-
Did more than 1 investigator adjudicate culprit lesion detection?	Yes (+) or no (-)	-	-	-	-	+	-	-	-
Were culprit and nonculprit lesions matched in terms of vessel stenosis severity?	Yes (+) or no (-)	-	-	-	-	-	-	-	-

Note: If data not provided or not specified, recorded as no (-).

evaluating these specific elements in the intracranial circulation. The lack of studies evaluating these features may be because these specific elements are much more difficult to detect in the intracranial vasculature due to its smaller size and deeper location compared to the extracranial carotid arteries where they have been more well characterized, including in studies with histopathological confirmation.¹³ Similarly, though some recent studies have proposed quantitative biomarkers of high-risk plaque such as arterial remodeling ratios or surface area measurements of plaque,¹⁹⁻²¹ such measures require intensive and precise measurements of plaque that may be challenging to implement into clinical practice. Evaluating for the presence or absence of gadolinium enhancement, on the other hand, may offer a relatively simple and rapid method of detecting high-risk plaque that could be used routinely in MRI examinations of patients presenting with acute ischemic stroke.

Our study has revealed some limitations about the existing literature evaluating the relationship between plaque MRI enhancement and acute ischemic stroke. First, though there was general agreement on the MRI features of plaque enhancement, including a greater degree of signal intensity than normal vessel wall enhancement, the reference standard by which abnormal enhancement is determined varied by study. The pituitary gland may offer an easily identifiable reference standard by which plaque enhancement can be subjectively assessed since it is located in close proximity to

the circle of Willis vessels, thereby ensuring its presence in the MRI field-of-view. Indeed, good interobserver consistency was shown using the pituitary gland as a reference to judge enhancement in the 1 study¹⁸ reporting interpretive reproducibility. Future studies will be strengthened if authors report interreader reproducibility measures more consistently in their interpretation of intracranial vessel wall MRI studies. Second, we found that studies did not routinely control for or match enhancing plaques leading to downstream infarction versus plaque in infarct-free territory in regard to stenosis severity. Indeed, stenosis severity was not systematically reported in all included studies. This makes it unclear to what extent stenosis severity and plaque volume may have acted as confounding factors in the strong association we found between plaque enhancement and ipsilateral ischemic stroke. Third, we found variability in how rigorously patients with alternative causes of ischemic stroke such as cardioembolism or more proximal extracranial artery-to-artery embolism were excluded from individual studies. Therefore, it is possible that infarcts attributed to an enhancing plaque supplying that vascular territory may have actually been caused by another mechanism. We believe that this risk is relatively low given our within-subjects design, in which infarcts in the territory of enhancement were compared to infarcts in other territories within the same patient. In other words, if an infarct were due to cardioembolism, that infarct should not be more likely to occur in the territory of an

enhancing plaque than in the vascular territory of an artery without enhancing plaque. However, we believe that future studies should be conducted in patients with strictly defined cryptogenic stroke in whom nonstenosing intracranial atherosclerotic plaques are present. Fourth, existing data only provided a crude OR and not covariate-adjusted OR. However, because most studies utilized a *within*-subject design, the risk of confounding systemic vascular risk factors should be relatively low. Fifth, as an emerging diagnostic technique, there are relatively few studies addressing the topic of intracranial plaque enhancement and its association with ischemic stroke. As such, though our study showed no statistical evidence of publication bias, this possibility should be re-evaluated as more studies are published in this field since an increasing study sample size will allow for a more valid and reliable estimate of publication bias. Sixth, there were differences in the MRI protocols used to evaluate for enhancing intracranial plaques, including differences in magnet field strength and specific pulse sequences used by investigators. We believe that for intracranial vessel wall MRI to be a useful tool in both clinical trial and patient care settings, future efforts to further standardize MRI acquisition parameters are warranted.

In summary, our systematic review and meta-analysis suggests that MRI-detected intracranial plaque enhancement is strongly associated with ipsilateral acute ischemic stroke. Plaque enhancement may offer a relatively simple means to detect a high-risk vessel wall feature that may play a complementary role to luminal stenosis measurements in diagnosing stroke etiology. Since conventional stroke etiology classification schemes attribute the cause of a stroke to large vessel intracranial atherosclerosis only when luminal stenosis is $\geq 50\%$, MRI for plaque enhancement may be most useful to evaluate the etiology of strokes currently considered cryptogenic. Future prospective studies in patients with ischemic strokes of undetermined etiology should be undertaken in which patients demonstrating vessel wall enhancement are followed for recurrent ischemic events. Such studies are necessary to test whether proven medical therapies for secondary stroke reduction are beneficial in patients with an enhancing intracranial plaque in the vascular territory of an acute ischemic stroke.

Acknowledgments

The authors wish to thank Dr Jeong-Min Kim, MD, PhD for providing unpublished data used in this meta-analysis.

Sources of Funding

This study was funded by grant K23NS082367 (Kamel) from National Institutes of Health/National Institute of

Neurological Disorders and Stroke (NIH/NINDS) and by grant KL2TR000458 (Gupta) from National Institutes of Health/National Center for Advancing Translational Sciences (NIH/NCATS).

Disclosures

None.

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SUPPLEMENTAL MATERIAL

Supplemental Methods: Data S1

Search Methodology Details

Ovid MEDLINE(R) In-Process & Other Non-Indexed Citations and Ovid MEDLINE(R) 1946 to Present - December 21, 2015

1. exp Stroke/
2. Stroke\$.tw.
3. cerebrovascular.tw.
4. exp Brain Ischemia/
5. ((brain or vascular or lacunar or venous or cerebral or isch?emi\$) adj2 (accident\$ or infarct\$ or event\$ or attack\$)).tw.
6. (cva or cvas).tw.
7. exp Magnetic Resonance Imaging/
8. (magnetic resonance or MR or MRI or MRIs or MRA or MRDTI).tw.
9. Contrast Media/
10. Imaging, Three-Dimensional/
11. exp Image Enhancement/
12. Gadolinium DTPA/
13. (vessel adj3 imag\$).tw.
14. (Gadolinium or Gd?DTPA or Magnograf or Magnevist or gadopentetate).tw.
15. (contrast or enhance\$ or 3?D or Dimension\$).tw.
16. Intracranial Arteriosclerosis/
17. Plaque, Atherosclerotic/
18. (LRNC or ((athero\$ or steno\$ or occlus\$ or ulcer\$ or plaque\$ or intraplaque or h?emorrhag\$ or IPH or narrow\$ or obstruct\$ or constrict\$ or bruit\$ or lipid or fibrous or culprit or lesion\$) adj3 (intracranial or cerebr\$ or brain))).tw.
19. or/1-6
20. or/7-8
21. or/9-15
22. or/16-18
23. and/19-22
24. (animals not (humans and animals)).sh.
25. 23 not 24

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|----|---|
| #1 | MeSH descriptor: [Stroke] explode all trees |
| #2 | stroke |
| #3 | cerebrovascular |

- #4 MeSH descriptor: [Brain Ischemia] explode all trees
- #5 ((brain or vascular or lacunar or venous or cerebral or isch*emi*) near/2 (accident* or infarct* or event* or attack*))
- #6 cva or cvas
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- #10 MeSH descriptor: [Imaging, Three-Dimensional] this term only
- #11 MeSH descriptor: [Image Enhancement] explode all trees
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- #19 #1 or #2 or #3 or #4 or #5 or #6
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10. three dimensional imaging/
11. image enhancement/
12. image quality/
13. digital subtraction angiography/
14. radiography/
15. computer assisted tomography/
16. positron emission tomography/

17. single photon emission computer tomography/
18. gadolinium pentetate/
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J Am Heart Assoc. 2016;5:e003816; originally published August 15, 2016;
doi: 10.1161/JAHA.116.003816

The *Journal of the American Heart Association* is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
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