Role of CT Angiographic Plaque Morphologic Characteristics in Addition to Stenosis in Predicting the Symptomatic Side in Carotid Artery Disease

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AJNR Am J Neuroradiol 2010, 31 (7) 1254-1260
doi: https://doi.org/10.3174/ajnr.A2078
http://www.ajnr.org/content/31/7/1254
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BACKGROUND AND PURPOSE: Atherosclerotic disease of the carotid artery is an important cause of ischemic stroke. We evaluated carotid plaque morphologic features by using CTA in addition to stenosis in the setting of symptomatic hemispheric TIA/stroke to identify factors that may predict plaque activity.

MATERIALS AND METHODS: Six hundred seventy-three patients (408 men; ages, 18–91 years; mean, 65.8 ± 15.2 years) presenting with hemispheric ischemic symptoms and having a CTA that included imaging of both carotid arteries within 24 hours were studied. Scans were interpreted for morphologic features, such as plaque length and width, attenuation, shape, surface, presence and degree of calcification, and ILT in addition to stenosis.

RESULTS: Univariable analysis showed that carotid occlusions ($P = .01$, OR = 5.27), high-grade stenosis (70%–99%) ($P = .06$, OR = 1.8), and the presence of ILT ($P = .01$, OR = 4.33) were highly predictive of the symptomatic side. Smooth plaque ($P = .01$, OR = 0.73) and extensive calcification ($P = .03$, OR = 0.72) were more commonly associated with the asymptomatic side. There was no correlation between plaque hypoattenuation ($P = .7$, OR = 1.06) or ulcerated plaque ($P = .74$, OR = 0.955) in predicting the symptomatic side. In a multivariable logistic regression model, the presence of ILT was still found to be significantly associated with the symptomatic side ($P = .048$, OR = 3.1) and the presence of extensive calcification, with the asymptomatic carotid artery ($P = .047$, OR = 0.69).

CONCLUSIONS: In addition to higher stenosis grades, the presence of ILT is highly predictive of the symptomatic side in carotid disease. Smooth plaque and extensive calcification seem to afford a protective effect. This information may be useful in radiologic risk stratification in carotid disease in addition to the current evidence available based on stenosis criteria alone.

ABBREVIATIONS: AHA = American Heart Association; CTA = CT angiography; ILT = intraluminal thrombus; MI = myocardial infarction; NASCET = North America Symptomatic Carotid Endarterectomy Trial; NIHSS = National Institutes of Health Stroke Scale; OR = odds ratio; TIA = transient ischemic attack

Received June 15, 2009; accepted after revision December 31.

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DOI: 10.3174/ajnr.A2078

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carotid artery atherosclerotic disease is an important cause of ischemic stroke, and thromboembolism is the predominant relevant stroke mechanism. Current clinical guidelines for revascularization strategies in carotid disease are based on stenosis criteria alone. These guidelines are derived from strong clinical trial data that examined primarily the long-term risk of stroke.1–3 Although degree of stenosis is a good surrogate marker for atherosclerotic vascular disease, based on the plausible assumption that vessel narrowing is caused by plaque accumulating in the lumen of the artery, there may be other factors that could potentially predict clinical behavior of atherosclerotic plaques. Plaque characteristics such as lipid-rich core, fibrous cap thickness, and intraplaque hemorrhage have been demonstrated on noninvasive imaging modalities such as sonography4–6 and MR imaging7–14 and it is suggested that these ancillary features, taken in conjunction with the degree of stenosis, may be important predictors of immediate and long-term stroke risk. More recently, especially with advances in multidetector technology, CT and CTA have also been found useful in the evaluation of patients with symptomatic carotid disease.15–17

In clinical practice, patients are seen who have clinical events that are not explained by the stenosis severity and who have negative findings on a work-up for other potential proximal embolic sources. With the increasing availability of noninvasive vascular imaging modalities such as Doppler sonography, MR imaging, and CTA, patients are also encountered with no clear clinical event but who have relatively high stenosis grades. It is also common to see severe stenotic disease on the side contralateral to the presenting hemispheric event.

We evaluated CTA features of carotid plaque that would potentially predict symptomatic carotid disease in addition to the degree of stenosis. At our dedicated stroke center, we perform CTA in patients with acute ischemic events as part of our clinical protocol for vascular and parenchymal imaging. We hypothesized that certain morphologic characteristics of atherosclerotic plaque could serve as an adjunct to stenosis grade
in potentially helping to decide revascularization options among patients who may not have been offered these same options based on stenosis criteria alone, especially if recurrent strokes occur with maximal medical therapy.

**Materials and Methods**

We retrospectively evaluated CTA data, obtained as part of standard clinical care for patients with stroke presenting to our institution, with institutional review board approval. At our center, patients with suspected acute ischemic stroke undergo evaluation with a plain CT followed by CTA of the cervical and intracranial vasculature as per institutional protocol. We identified consecutive patients from June 2004 to August 2007 who presented to the stroke neurology service, usually through the emergency department, with hemispheric ischemic symptoms and/or TIA, including amaurosis fugax, and who also had a CTA that included imaging of both carotid arteries within 24 hours of admission. All patients were reviewed clinically by a stroke neurologist and had a baseline NIHSS score documented on admission. Patients with technically inadequate CTA studies were excluded. The affected hemispheric side was determined, and the ipsilateral carotid artery was designated as the “case” carotid. The contralateral carotid artery served as the “control.” The use of contralateral carotid arteries has advantages because it automatically accounts for age, sex, and cardiovascular risk factors.

**CTA Technique**

CTA was performed by using multidetector scanners. Both a 4-section CT scanner (LightSpeed Plus; GE Healthcare, Milwaukee, Wisconsin) and a 64-section scanner (SOMATOM Sensation 64; Siemens, Erlangen, Germany) were used. The distribution of patients between both types of scanners was roughly equal. Both scanners used an automated trigger technique for injection of contrast at 5 mL/s for a total of 90–120 mL. Scanning was performed from the aortic arch to the vertex. The 4-section scanner used a collimation of 4 × 1.25 mm, a pitch of 0.9, and a rotation time of 0.5 seconds to acquire the raw data, which was reconstructed at 2.5-mm-thick overlapping sections for axial images. Thinner sections (1.25 mm at 50% overlap) were used to reconstruct axial images of the circle of Willis. Images of the cervical carotid arteries were reconstructed in the sagittal plane at a 4-mm thickness. With the 64-section scanner, a collimation of $64 \times 0.6$ mm and pitch of 0.9 for a rotation time of 0.5 seconds was used to acquire the raw data, which were reconstructed at 1-mm overlapping sections for the axial images. Reconstructions for axial, coronal, and sagittal images of the circle of Willis as well as the carotid arteries on each side were performed at a 3-mm thickness at 1-mm intervals. 

**Evaluation of Findings**

The CTA images were evaluated by 4 readers, who analyzed the images in groups of 2, 1 neurologist and 1 neuroradiologist who were experienced in the evaluation of CT and CTA images in patients with acute stroke. The readers were blinded to the symptom side as well as other clinical information. Stenosis was calculated from sagittal images reconstructed along the long axis of the internal carotid artery, by using the NASCET approach. The morphologic features including plaque attenuation (Fig 1), plaque shape (Figs 2 and 3), plaque surface (Fig 4), and degree of calcification (Fig 5) were determined from a review of all images. The various plaque shapes were classified as linear, crescent, sessile, pedunculated, or circumferential. The plaque surface was classified as smooth, irregular without ulceration, and ulcerated. The length and width of the plaque were also documented as was the presence of an intraluminal thrombus (Fig 6). An “intraluminal thrombus” was defined as an eccentric pedunculated filling defect projecting into the lumen of the vessel. Discrepant findings were resolved by consensus.

**Statistical Methods**

Statistical analysis was performed with the STATA statistical software, Version 10 (StataCorp, College Station, Texas). For evaluation, the degree of stenosis was categorized as follows: $<30\%$, $30\%–49\%$, $50\%–69\%$, $70\%–99\%$, and occlusion. Data were reported by using standard descriptive summary statistics. Univariable analysis was used to assess the relationship between each individual plaque feature in addition to the category of stenosis and symptomatic side status. A multivariable logistic regression model was developed by using backward stepwise elimination of imaging factors to provide an overall assessment of the role of these factors.
in predicting the symptomatic side. A parsimonious model was sought so that those factors not significant at $P < .05$ were dropped. Age and sex were forced variables in this model. We accounted for the dependence of variables within patients by clustering the analysis within patients.

**Results**

There were 988 patients who had evaluations of the carotid arteries between June 2004 and August 2007. Of these patients, 972 patients had scans that were adequate for interpretation. Of these patients, 673 (408 male patients and 265 female patients; ages, 18–91 years; mean, 65.8 ± 15.2 years) were identified who had a clear hemispheric event, and these patients constituted the study population. Three hundred forty-nine patients had right-hemispheric symptoms, and 324 patients had left-hemispheric symptoms. The ipsilateral carotid arteries served as cases ($n = 673$) and the contralateral side, as controls ($n = 673$). The demographic information for the study population is given in Table 1.

Univariable analysis (Table 2) showed that carotid occlusions ($P = .01$, OR = 5.27), high-grade stenosis (70%–99%) ($P = .06$; OR = 1.9), and the presence of intraluminal thrombus ($P = .01$, OR = 4.33) were highly predictive of the symptomatic side. In addition, some features were more commonly associated with the asymptomatic side, such as smooth plaque surface ($P = .01$, OR = 0.73) and extensive calcification ($P = .03$, OR = 0.72). There was no correlation between plaque hypoattenuation ($P = .7$; OR = 1.06) or ulcerated plaque ($P = .74$, OR = 0.955) in predicting the symptomatic side.

In a multivariable logistic regression model accounting for age, sex, and degree of stenosis, the presence of intraluminal thrombus was still found to be significantly associated with the symptomatic side ($P = .048$, OR = 3.1) and the presence of extensive calcification, with the asymptomatic carotid ($P = .047$; OR = 0.69). This also demonstrated a graded increase in the magnitude of effect as the degree of stenosis increased toward occlusion (Table 2). The fre-
quency of significant findings in cases versus controls is shown in Table 3.

**Discussion**

Existing measures for the effectiveness of endarterectomy among patients with symptomatic carotid disease have been based on the degree of stenosis from strong clinical trial data.\(^1\)\(^-\)\(^3\),\(^18\) The degree of stenosis is also the basis for surgical intervention in asymptomatic patients\(^19\) and in endovascular treatment for carotid disease.\(^20\) The premise that various pathologic stages in the evolution of carotid atheroma are predictive of clinical events remains to be convincingly proved in patients with stroke. Our understanding of this process in carotid atheromas has been extrapolated from the histologic classification of coronary plaques developed by the AHA.\(^21\),\(^22\) Noninvasive modalities can image atherosclerotic carotid vascular disease and depict these various pathologic stages. Plaque characteristics such as lipid-rich core, fibrous cap thickness, and intraplaque hemorrhage have been demonstrated on sonography\(^4\)\(^-\)\(^6\) and MR imaging,\(^7\)\(^-\)\(^14\),\(^23\)\(^-\)\(^25\) and it is suggested that these ancillary features, taken in conjunction with the degree of stenosis, may be important predictors of future stroke risk. CT angiography is being increasingly used to noninvasively image carotid stenosis\(^26\),\(^27\) and has also been shown to be useful in the evaluation of carotid plaque morphology.\(^15\),\(^17\),\(^28\)\(^-\)\(^31\)

The findings from our study indicate that apart from increasing degree of stenosis, the presence of an intraluminal thrombus is highly predictive of the symptomatic side in patients with carotid disease. In addition, we observed that ex-
tensive calcification and smooth plaques were more likely to be associated with the asymptomatic side. Whether this in fact correlates with protective mechanisms requires large-scale prospective trials looking at natural history data. Other studies have shown that patients with extensive calcification of plaques on endarterectomy specimens are less likely to have symptomatic disease.32

The presence of an intraluminal thrombus has implications in the management of patients with complicated carotid plaques33,34, and was seen in ≈1.8% of patients in the NASCET trial.1 In our series, this was defined as an eccentric protruding filling defect within the lumen, which, on cross-section, appears to lie centrally within the lumen, surrounded by contrast (Fig 6). CT angiography may be more sensitive to the detection of intraluminal thrombus because it is less prone to the volume averaging effects from surrounding contrast as in a conventional catheter angiogram. Some of the patients with intraluminal thrombus have shown resolution of the thrombotic filling defect on follow-up imaging after aggressive antithrombotic medical treatment. These observations suggest that the presence of this finding might indicate plaque activity based on an interaction of factors such as platelet activation and aggregation and continued exposure to thrombogenic plaque contents.

We also observed that the presence of a smooth-appearing plaque and extensive calcification seems to afford a protective effect. As a corollary to the discussion on intraluminal thrombus, it is plausible that unless it is causing a hemodynamic effect, a smooth appearance of a plaque may indicate relative plaque stability and a lower grade plaque without complication (AHA I-V).21,22 Calcification may be protective by preventing adhesion, and activation and subsequent aggregation forming platelet-rich thrombi and corresponding to AHA type VII.22

Wintermark et al.,17 in a smaller cross-sectional retrospective study, by using a custom automated computer algorithm to evaluate various wall descriptors, showed that wall volume, fibrous cap thickness, the number and location of lipid clusters, and the number of calcium clusters were significantly associated with acute carotid stroke. Of 136 patients, they identified 40 patients with “acute carotid stroke” and 50 patients with “noncarotid strokes.” They analyzed features between the 2 groups and also between the symptomatic and asymptomatic sides for the 40 patients with acute carotid stroke.

Table 1: Patient demographics

<table>
<thead>
<tr>
<th>Demographic</th>
<th>Total (N = 673), Instances (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male</td>
<td>408</td>
</tr>
<tr>
<td>Female</td>
<td>265</td>
</tr>
<tr>
<td>Median age (range)</td>
<td>69 (18–91)</td>
</tr>
<tr>
<td>Median NIHSS score at baseline (range)</td>
<td>5 (0–25)</td>
</tr>
<tr>
<td>Vascular risk factors</td>
<td></td>
</tr>
<tr>
<td>Hypertension</td>
<td>401 (59.6)</td>
</tr>
<tr>
<td>Diabetes</td>
<td>114 (16.9)</td>
</tr>
<tr>
<td>Atrial fibrillation</td>
<td>95 (14.1)</td>
</tr>
<tr>
<td>Valvular heart disease</td>
<td>25 (3.7)</td>
</tr>
<tr>
<td>History of TIA/stroke</td>
<td>129 (19.2)</td>
</tr>
<tr>
<td>Recent MI</td>
<td>10 (1.5)</td>
</tr>
<tr>
<td>Family history of stroke</td>
<td>65 (9.7)</td>
</tr>
<tr>
<td>Current smoker</td>
<td>165 (24.5)</td>
</tr>
<tr>
<td>Former smoker</td>
<td>74 (11.0)</td>
</tr>
<tr>
<td>Hypercholesterolemia</td>
<td>195 (29.0)</td>
</tr>
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</table>

Table 2: Results of univariable and multivariable logistic regression

<table>
<thead>
<tr>
<th>Variable</th>
<th>Odds Ratio</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Univariable logistic regression</td>
<td></td>
<td></td>
</tr>
<tr>
<td>30%–49% Stenosis</td>
<td>0.90</td>
<td>.64</td>
</tr>
<tr>
<td>50%–69% Stenosis</td>
<td>1.27</td>
<td>.28</td>
</tr>
<tr>
<td>70%–99% Stenosis</td>
<td>1.90</td>
<td>.06</td>
</tr>
<tr>
<td>Occlusions</td>
<td>5.27</td>
<td>.01</td>
</tr>
<tr>
<td>Plaque length</td>
<td>0.99</td>
<td>.51</td>
</tr>
<tr>
<td>Plaque width</td>
<td>0.98</td>
<td>.43</td>
</tr>
<tr>
<td>Smooth surface</td>
<td>0.73</td>
<td>.01</td>
</tr>
<tr>
<td>Irregular surface</td>
<td>0.86</td>
<td>.20</td>
</tr>
<tr>
<td>Ulcerated surface</td>
<td>0.96</td>
<td>.74</td>
</tr>
<tr>
<td>No calcification</td>
<td>0.76</td>
<td>.08</td>
</tr>
<tr>
<td>Mild calcification</td>
<td>0.83</td>
<td>.16</td>
</tr>
<tr>
<td>Moderate calcification</td>
<td>0.87</td>
<td>.28</td>
</tr>
<tr>
<td>Extensive calcification</td>
<td>0.72</td>
<td>.03</td>
</tr>
<tr>
<td>Hypodense plaque</td>
<td>1.06</td>
<td>.70</td>
</tr>
<tr>
<td>Isodense plaque</td>
<td>0.83</td>
<td>.29</td>
</tr>
<tr>
<td>Heterogeneous plaque</td>
<td>0.81</td>
<td>.05</td>
</tr>
<tr>
<td>Intraluminal thrombus</td>
<td>4.33</td>
<td>.01</td>
</tr>
</tbody>
</table>

Multivariable logistic regression of degree of stenosis accounting for age and sex

<table>
<thead>
<tr>
<th>Variable</th>
<th>Odds Ratio</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>30%–49%</td>
<td>1.00</td>
<td>.99</td>
</tr>
<tr>
<td>50%–69%</td>
<td>1.38</td>
<td>.26</td>
</tr>
<tr>
<td>70%–99%</td>
<td>1.90</td>
<td>.07</td>
</tr>
<tr>
<td>Occlusions</td>
<td>4.99</td>
<td>.00</td>
</tr>
</tbody>
</table>

Multivariable logistic regression of ILT and calcification accounting for age, sex, and degree of stenosis

<table>
<thead>
<tr>
<th>Variable</th>
<th>Odds Ratio</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Intraluminal thrombus</td>
<td>3.10</td>
<td>.05</td>
</tr>
<tr>
<td>Extensive calcification</td>
<td>0.69</td>
<td>.05</td>
</tr>
</tbody>
</table>

Table 3: Percentage of significant findings in cases versus controls

<table>
<thead>
<tr>
<th></th>
<th>Cases (%)</th>
<th>Controls (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Occlusions</td>
<td>44 (6.54)</td>
<td>9 (1.34)</td>
</tr>
<tr>
<td>ILT</td>
<td>17 (2.53)</td>
<td>4 (0.59)</td>
</tr>
<tr>
<td>Hypodense plaque</td>
<td>64 (9.51)</td>
<td>56 (8.32)</td>
</tr>
<tr>
<td>Plaque ulceration</td>
<td>85 (12.63)</td>
<td>82 (12.18)</td>
</tr>
<tr>
<td>Smooth plaque</td>
<td>103 (15.30)</td>
<td>130 (19.32)</td>
</tr>
<tr>
<td>Extensive calcification</td>
<td>53 (7.88)</td>
<td>68 (10.10)</td>
</tr>
</tbody>
</table>

* OR of a finding being associated with the symptomatic side and P value.
stroke. Their study was robust in that their use of a comput-
erized algorithm to quantitatively assess a battery of carotid
features yielded objective parameters for studying carotid ath-
erosclerotic disease. Also using a noncarotid stroke population
as a control allows a truer determination of factors contribut-
ing to atherosclerotic carotid disease. In our study, we in-
cluded many of those wall descriptors but in a manner that
allows their identification by visual inspection from scans ob-
tained within existing clinical protocols, and we used only
contralateral carotid arteries as controls.

In our series, we did not find evidence to suggest that hy-
poattenuated plaque or the presence of ulceration is signifi-
cantly associated with the symptomatic side. It could be pos-
sible that in the absence of thrombi formation, the mere
presence of a lipid-rich necrotic core may not be a causative
factor for ischemic symptoms in the acute setting or that CT
may not be accurately detecting attenuation changes within
plaque to the level required to achieve statistical significance.
Most interesting, in our patients, the univariable logistic re-
gression did not show that the presence of an ulcer was signifi-
cantly associated with the symptomatic side. In their study of
406 patients by using CTA, de Weert et al\(^30\) showed that com-
plexed plaques, which included both irregular and ulcerated
plaques, were more often present in the symptomatic carotid
artery than in the contralateral asymptomatic carotid artery
(25% versus 18%, \(P < .01\)). A multivariable analysis, however,
suggested that this could be attributed to the significantly
higher stenosis grade present in symptomatic arteries.

MR imaging has been, in general, the more common tech-
nique for detecting complicated plaque. Murphy et al\(^9\) dem-
onstrated the utility of direct thrombus imaging in patients
with carotid disease by using MR imaging to suggest that high
signal intensity on T1-weighted images may represent the in-
traplaque hemorrhage component of complicated plaques.
In a progress review, Wasserman et al\(^11\) suggested that high-res-
olution black-blood MR imaging may predict vulnerable le-
sions even in low-grade carotid stenosis. In a recent study,
Bitar et al\(^13\) showed that T1 hyperintense intraplaque signal
intensity on 3D high spatial resolution on in vivo imaging
represents areas of intraplaque hemorrhage on histologic sam-
ples. The utility of CT to detect intraplaque hemorrhage as a
component of vulnerable plaque is limited by inadequate con-
trast resolution. Wasserman et al\(^12\) also demonstrated that in-
creased contrast enhancement within the fibrous cap and
outer wall of a carotid atheroma after administration of intra-
venous gadolinium might reflect sites of active inflammation.
This may be indicative of the neovascularity seen with plaque
instability. We think that it is unlikely that CT-based tech-
niques would be able to depict such inflammatory change.

Our study should be interpreted in the context of the fol-
lowing limitations: We acknowledge that the study was per-
fomed in a retrospective manner. We also note that there are
multiple factors involved in the pathophysiologic mechanisms
underlying cerebrovascular ischemia, and in our patients, we
believe that symptoms related to hemispheric ischemia origi-
nated in the carotid arteries. Using contralateral carotid arter-
ies as controls has advantages in that various factors such as
age and cardiovascular risk factors are accounted for and an
adequate sample size of controls is obtained. The alternative
would have been to use age-matched controls with noncarotid
disease (such as patients with a proximal embolic source). We
also acknowledge that certain findings may be related to the
inherent nature of CT technology, especially when compared
with MR imaging—based studies. We think that CT may not
be adequate to demonstrate some features of complicated
plate such as intraplaque hemorrhage, neovasularity, and
small surface ulcerations. The inability to detect a significant
difference with hypoattenuated plaques in our study may re-
late to the fact that attenuation values were not objectively
measured. Also, it is unclear whether the differences in scanner
technology used (4-detector scanner versus 64-detector) had
an effect on the data.

In summary, our findings suggest that certain morphologic
characteristics of carotid plaque on CTA are seen more com-
monly with symptomatic carotid disease at the time of presen-
tation and may predict clinical behavior in addition to the
degree of stenosis alone. This may help in the decision-making
process with regard to the need and timing of carotid revascular-
ization in the appropriate clinical setting.

Acknowledgments
We acknowledge the help of Jayanta Roy, Imanuel Dzialowski,
Volker Puetz, Christine O’Reilly, and Sherif Idris.

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