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Symptomatic and Asymptomatic Chronic Carotid Artery Occlusion on High-Resolution MR Vessel Wall Imaging

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ABSTRACT

BACKGROUND AND PURPOSE: Chronic carotid artery occlusion remains a poorly understood risk factor for subsequent stroke, and potential revascularization is dependent on understanding the anatomy and nature of the occlusion. Luminal imaging cannot assess the nature of an occlusion, so the internal structure of the occlusion must be inferred. The present study examines the signal characteristics of symptomatic and asymptomatic carotid occlusion that may point to management differentiation.

MATERIALS AND METHODS: We prospectively recruited patients who were diagnosed with chronic carotid artery occlusion defined as longer than 4 weeks and confirmed by DSA. All patients underwent high-resolution MR vessel wall imaging examinations after enrollment. Baseline characteristics, vessel wall imaging features, and DSA features were collected and evaluated. The vessel wall imaging features included segment involvement, signal intensity, contrast enhancement, and vessel wall thickness. The symptomatic and asymptomatic chronic carotid artery occlusions were compared.

RESULTS: A total of 44 patients with 48 lesions were included in this study from February 2020 to December 2020. Of the 48 lesions, 35 (72.9%) were symptomatic and 13 (27.1%) were asymptomatic. There was no difference in baseline and DSA features. On vessel wall imaging, C1 and C2 were the most commonly involved segments (91.7% and 68.8%, respectively). Compared with symptomatic lesions, asymptomatic lesions were more often isointense (69.2%) in the distal segment (P = .03). Both groups had diffuse wall thickening (80% and 100%).

CONCLUSIONS: Signal characteristics between those with symptomatic and asymptomatic carotid artery occlusions differ in a statistically significant fashion, indicating a different structure of the occlusion.

 $\label{eq:ABBREVIATIONS: CCAO = chronic carotid artery occlusion; VWI = vessel wall imaging$

C hronic carotid artery occlusion (CCAO) accounts for 6.5% of ischemic strokes.¹ CCAO may present as an incidental finding in asymptomatic patients, or it may present clinically as a TIA or a severe disabling stroke. The annual risk of TIA or ischemic stroke of asymptomatic CCAO is estimated to be between 2% and 8%.²⁻⁴ The risk of stroke recurrence in symptomatic CCAO is 8% at 30 days, 10% at 1 year, and 14% at 5 years.⁵

CCAO may cause ischemic stroke due to an embolism from the occluded segment, compromised hemodynamics, and a mixture of both embolic and hemodynamic factors.⁶ The role of hemodynamics in patients with symptomatic and asymptomatic CCAO has been demonstrated in previous studies.^{4,7} The biology and architecture of the occluded segment of a CCAO remain unclear. Conventional imaging modalities including CTA, MRA, ultrasonography, and DSA can evaluate the structural and hemodynamic functional information of nonoccluded vessels but cannot provide sufficient information about the features of vessel segments with no substantial flow. High-resolution MR vessel

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wall imaging (VWI) has been widely used in atherosclerotic diseases of the carotid and intracranial arteries.⁸⁻¹¹ VWI can determine plaque size, intraplaque hemorrhage, large lipid core, and intraluminal thrombus and can evaluate dissection flaps and subintimal plaque.¹² In addition, 3D VWI with variable flip angles can offer imaging with improved longitudinal coverage in a shorter imaging time.^{13,14} VWI has been reported to be practicable and accurate in the diagnosis of CCAO.^{15,16} VWI for CCAO can provide more information and direct visualization of the occluded segment, such as segment involvement, signal intensity, enhancement characteristics, and vessel wall thickness. However, few studies have focused on the characteristics of the occluded segment of CCAO on VWI.¹⁷ Therefore, we conducted this study to assess the differing imaging characteristics between symptomatic and asymptomatic CCAO.

MATERIALS AND METHODS

Study Population

This was a prospective study, approved by the institutional review board of the hospital. Informed consent was obtained from all patients or their representatives before enrollment. Patients admitted to the Department of Interventional Neuroradiology of Beijing Tiantan Hospital and the Department of Neurology of The First Affiliated Hospital of Zhejiang University who had been diagnosed with carotid artery occlusion by sonography, CTA, or MRA were selected. Carotid artery occlusion by sonography, CTA, or MRA was defined as 100% cross-sectional truncation of the vessel lumen. "Chronic" was defined as the patient having a carotid occlusion diagnosis for >4 weeks, confirmed by DSA, with TICI grade 0.^{18,19} All patients underwent DSA and VWI examinations after enrollment. The symptomatic CCAO group comprised patients who were admitted due to anterior ipsilateral ischemic events in the territory of the occluded vessel. The asymptomatic CCAO group was patients diagnosed with carotid artery occlusion incidentally during the course of routine clinical care. Ischemic events included ischemic stroke and TIA on the ipsilateral hemisphere or retinal artery. The exclusion criteria were the following: 1) contraindications to VWI or DSA examination; 2) near-occlusion or pseudo-occlusion confirmed by DSA; 3) intracranial chronic carotid artery occlusion; and 4) cardioembolic risk factors.

Data Collection

Baseline data including sex, age, body mass index, atherosclerotic risk factors, antithrombotic agents taken before and after diagnosis of CCAO, history of radiation, and time of the ischemic event onset were collected. Risk factors were defined as the following: hypertension defined as a history of hypertension or the patient taking any hypotensive drugs; diabetes mellitus defined as a history of diabetes mellitus, the patient taking any hypoglycemic agents, or a glycosylated hemoglobin level of \geq 7%; hyperlipidemia defined as a history of hyperlipidemia or receiving lipid-lowering treatment; and coronary artery disease defined as a history of myocardial infarction or angina pectoris. Smoking was defined as current smokers or a smoking history. VWI and DSA features were recorded. On VWI, features of segment involvement, signal intensity of the occluded segment, contrast enhancement, and vessel wall thickness were evaluated. On DSA, the imaging features of the vascular stump and the level of the distal reconstitution were also evaluated.

Vessel Wall Imaging

VWI was performed on a 3T Magnetom Trio MR imaging scanner (Siemens) or a 3T Discovery MR750 scanner (GE Healthcare) with a head/neck coil. The 3D T1-weighted turbo spin-echo sequences on the Magnetom Trio MR imaging scanner were obtained with the following parameters: TR/TE = 760/15 ms, section partial Fourier factor = 0.75, turbo factor = 60, echo spacing = 4.52 ms, parallel imaging acceleration = 3. The FOV was 240 \times 220 \times 172 mm (Foot-Head \times Anterior-Posterior \times Right-Left), and the voxel size was 0.54 $\, imes$ 0.54 $\, imes$ 0.54 mm. The 3D T1-weighted turbo spin-echo sequences on the Discovery MR750 were obtained with the following parameters: TR/TE = 675/21 ms, echo-train length = 24, echo spacing = 7.5 ms, bandwidth = 62.50 kHz, phase and HyperSense acceleration (GE Healthcare) = 2, FOV = 180 × 180 mm (Superior-Inferior \times Right-Left). The pixel size was 0.6 \times 0.6 mm, and the section thickness was 0.8 mm. Other MR imaging scans included 3D TOF MRA, T2-weighted imaging, and contrastenhanced T1-weighted imaging. Images were reconstructed using the Reformate tool in the Advantage Workstation 4.5 (GE Healthcare) or the D multiple planer reconstruction tool in the Siemens workstation.

DSA Imaging

All DSA studies were performed by 2 experienced interventional neuroradiologists with >10 years' experience each. During angiography, the bilateral common carotid arteries and bilateral vertebral arteries received contrast injections in the anterior-posterior and lateral projections, respectively.

Imaging Analysis

All MR images were reviewed separately by a neuroradiologist with 6 years' experience and a neurologist with 8 years' experience who were blinded to clinical and medical information. A third reader was used to resolve disagreements in assessing the segment involvement. We divided each ICA into 7 segments (C1–C7) according to the classification of Bouthillier et al.²⁰ To measure signal intensities, we placed ROIs at the occluded segment on the coronal T1-weighted turbo spin-echo sequences, with maximal size and adjacent sternomastoid muscle by the consensus decision of the 2 investigators. Signal intensity was categorized as hypointensity (less than the signal intensity of the adjacent sternocleidomastoid muscle), isointensity (equal to that of the adjacent sternocleidomastoid muscle), hyperintensity (more than that of the adjacent sternocleidomastoid muscle), mastoid muscle), and heterogeneous intensity (a mix of the above).

The occluded segment was divided into the origin segment (origin of the ICA) and the distal segment (beyond the origin of the ICA) to assess the signal intensity separately. Contrast enhancement was determined by comparing pre- and postcontrast T1-weighted images focused on the intraluminal thrombus. Vessel wall thickness was evaluated by T1-weighted turbo spinecho sequences in the petrous ICA. The petrous ICA was divided into proximal, middle, and distal segments for measuring vessel wall thickness. Diffuse wall thickneing was defined as a mean wall thickness of >1.5 mm.²¹ Time intervals between the last

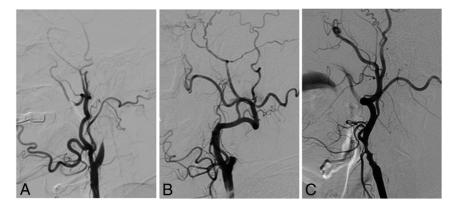


FIG 1. Stump condition: tapered stump (A), blunt stump (B), no stump (C)

Table 1: Baseline data of	patients with	asymptomatic and	symptomatic CCAO
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	Asymptomatic	Symptomatic	Р
	(n = 9)	(n = 35)	Value
Male (No.) (%)	8 (88.9)	34 (97.1)	.37
Age (mean) (yr)	59 (SD, 13.35)	60.51 (SD, 11.46)	.73
Body mass index (mean)	25.30 (SD, 3.49)	24.40 (SD, 3.12)	.46
Hypertension (No.) (%)	7 (77.8)	26 (74.3)	>.99
Diabetes mellitus (No.) (%)	3 (33.3)	12 (34.3)	>.99
Hyperlipidemia (No.) (%)	7 (77.8)	28 (80.0)	>.99
Coronary artery disease (No.) (%)	1 (11.1)	3 (8.6)	>.99
Smoker (No.) (%)	4 (44.4)	15 (42.9)	>.99
Antithrombotic treatment before diagnosis of	1 (11.1)	8 (22.9)	.75
CCAO (No.) (%)			
Antithrombotic treatment after diagnosis of	3 (33.3)	18 (51.4)	.55
CCAO (No.) (%)			
History of radiation (No.) (%)	0 (0.0)	1 (2.9)	>.99
Time intervals between DSA and VWI (median)	1 (1, 2.5)	2 (1, 4)	.74
(IQR) (days)			

Note:—IQR indicates interquartile range.

ischemic event onset to VWI were divided into 4 stages (<7 days, 7–30 days, 30–90 days, and \geq 90 days). The difference in signal intensity across the 4 time points was evaluated.

DSA images were reviewed by the 2 operators. According to the contrast filling within the ICA from the common carotid artery and the shape, the morphology of the carotid stump was categorized as tapered, blunt, and no stump (Fig 1). The level of the distal collateral reconstitution was defined as the most proximal site visible at the distal end of the occlusion, divided into proximal or distal cavernous segments.

Statistical Analysis

We performed statistical analysis using SPSS software, Version 20.0 (IBM). Normal distribution measurement data were expressed as mean (SD), while skew distribution data were expressed as median and interquartile range. Counting data were expressed in frequencies and percentages. We compared baseline data of symptomatic and asymptomatic patients. DSA features and VWI features were compared between those with symptomatic and asymptomatic CCAO. The Student *t* test (normal distribution) or Mann-Whitney *U* test (skewed distribution) and the χ^2 test or Fisher exact test were used according to the situation. *P* < .05 was considered a statistically significant difference.

RESULTS Patient Characteristics

Overall, a total of 44 patients with 48 lesions were included in this study from February 2020 to December 2020: 22 (50.0%) in Beijing Tiantan Hospital and 22 (50.0%) in The First Affiliated Hospital of Zhejiang University. Of the 44 patients, 35 (79.5%) were symptomatic and 9 (20.5%) were asymptomatic. In symptomatic patients, 32 patients had ischemic stroke (29 ipsilateral hemispheres and 3 retinal arteries) and 3 patients had experienced TIA (1 ipsilateral hemisphere and 2 retinal arteries). One (2.0%) patient had a history of radiation. Baseline characteristics are shown in Table 1; there was no difference in baseline characteristics between symptomatic and asymptomatic patients. The time intervals between DSA and VWI were 1 day (interquartile range, 1-2.5 days) in the asymptomatic group and 2 days (interquartile range, 1-4 days) in the symptomatic group (P = .74).

Comparison between Symptomatic and Asymptomatic CCAO

DSA showed no difference in stump morphology and the level of distal collateral reconstitution between the 2 groups (Online Supplemental Data).

On VWI, C1 and C2 were the most commonly involved segments in both

symptomatic CCAO (97.1% and 68.6%) and asymptomatic CCAO (76.9% and 69.2%). Two patterns of signal intensity were detected in the origin segment (isointensity and heterogeneous intensity). There were 7 (20.0%) instances of isointensity and 27 (77.1%) of heterogeneous intensity in symptomatic CCAO (Fig 2). There were 5 (38.5%) instances of isointensity and 5 (38.5%) of heterogeneous intensity in those with asymptomatic CCAO (P = .15). Three patterns of signal intensity were detected in the distal segment (hyperintensity, isointensity, and heterogeneous intensity). There was a higher prevalence of isointensity (69.2%) in the asymptomatic CCAO compared with the symptomatic CCAO (P = .03) group. There were 14 patients (15 lesions) with contrast-enhanced scans. Nine scans in 13 patients with symptomatic CCAO were demonstrated to be contrast-enhanced, as well as 2 scans in 2 patients with asymptomatic CCAO (P = .95). Both patients with symptomatic, and asymptomatic CCAO had a high prevalence of diffuse thickening of the vessel wall (80% and 100% respectively).

Difference in Signal Intensity at Different Stages of Symptomatic CCAO

In symptomatic CCAO, both the origin segment and distal segment showed no difference at the following 4 stages: <7 days, 7–30 days, 30–90 days, and \geq 90 days (Tables 2 and 3).

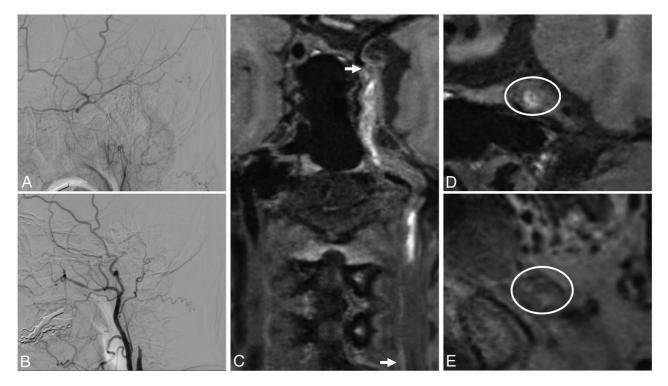


FIG 2. An adult man who presented with blurred vision for 1 month and right-limb weakness for 10 days. On DSA, the occluded segment is from C1 (*B*) to C6 (*A*). On VWI, 3D TI-weighted turbo spin-echo sequences show that the occluded segment is C1 (*arrow*) to C6 (*arrow*) (*C*). The signal intensity of the distal segment in a elliptical region-of-interest (ROI) (*D*), and the signal intensity of the origin segment is heterogeneous intensity in a elliptical ROI (*E*).

Table 2: Difference in signal intensity at the origin segment of symptomatic CCAO at different stages

	Hyperintensity (n = 0)	lsointensity (n = 7)	Hypointensity (n = 0)	Heterogeneous Intensity (n = 27)	<i>P</i> Value
<7 days (No.) (%)	0 (0.0)	0 (0.0)	0 (0.0)	5 (14.7)	.69
7–30 days (No.) (%)	0 (0.0)	4 (57.1)	0 (0.0)	16 (47.1)	
30–90 days (No.) (%)	0 (0.0)	2 (28.6)	0 (0.0)	10 (29.4)	
≥90 days (No.) (%)	0 (0.0)	1 (14.3)	0 (0.0)	3 (8.8)	

Table 3: Difference in signal intensity at the distal segment of symptomatic CCAO at different stages

	Hyperintensity (n = 9)	lsointensity (n = 14)	Hypointensity (n = 0)	Heterogeneous Intensity (n = 12)	<i>P</i> Value
<7 days (No.) (%)	2 (22.2)	1 (7.1)	0 (0.0)	2 (16.7)	.29
7–30 days (No.) (%)	2 (22.2)	6 (42.9)	0 (0.0)	8 (67.7)	
30–90 days (No.) (%)	4 (44.4)	4 (28.6)	0 (0.0)	2 (16.7)	
≥90 days (No.) (%)	1 (11.1)	3 (21.4)	0 (0.0)	0 (0.0)	

on VWI than DSA, and a tiny lumen could be seen beyond the occluded segment on VWI but was considered occluded on DSA (Fig 3).

DISCUSSION

The present study demonstrates that asymptomatic CCAO lesions have a higher prevalence of isointensity compared with symptomatic CCAO lesions. The most common occluded segments in CCAO lesions were C1 and C2. Most of the CCAO lesions had diffuse thickening of the vessel wall on VWI, so VWI may provide detailed characterization of the occlusion for patients with CCAO.

VWI for the diagnosis of CCAO has been reported to have high accuracy and sensitivity using DSA as a reference.^{15,22,23} A few studies have focused on the signal intensity of the occluded segment of the CCAO. Previous studies have reported a higher prevalence of heterogeneous intensity at the begin-

Comparison of DSA with VWI

The accuracy of VWI for the diagnosis of total CCAO was 100%. The level of occlusion differed between DSA and VWI, with a consistency rate of 27.1%. Most of the lesions were shown to be shorter

ning of the occluded segment and a 47.5% presence of high signal intensity in the occluded segment.^{15,17} Atherosclerosis is the major cause (about 70%) of CCAO, and most lesions are located at the origin of the ICA.²⁴ The deterioration of the atherosclerotic plaque

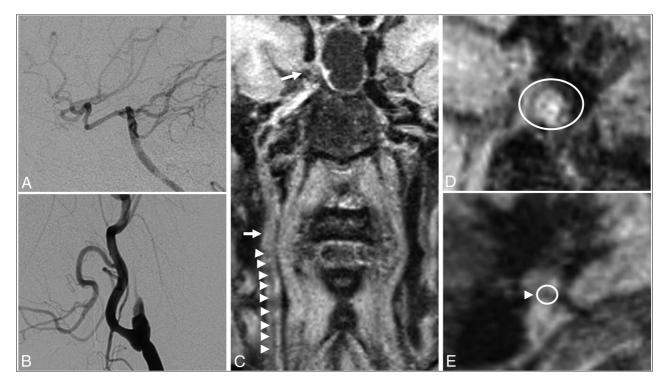


FIG 3. An adult woman who presented with dizziness for 2 months. On DSA, the occluded segment is from C1 (*B*) to C7 (*A*). On VWI, 3D TI-weighted turbo spin-echo sequences show that the occluded segment was C2 (*arrow*) to C4 (*arrow*), and C1 did not have total occlusion (*arrowheads, C*). The signal intensity of distal segment is isointensity in a elliptical region-of-interest (ROI) (*D*). The origin segment showed a tiny lumen (*arrowhead*), in a circular ROI which was considered an occluded segment on DSA (*E*).

leads to occlusion of the carotid artery and formation of the intraluminal thrombus within the distal segment. The signal intensities of the origin and distal segments are usually different. It is difficult to describe the whole occluded segment with uniform signal intensity. In this study, we divided the occluded segment into 2 parts to evaluate the signal intensity: the origin segment (the origin of the ICA) and the distal segment (distal C1 and above). Heterogeneous signal intensity was frequently encountered in the origin segment, but there was no difference between the 2 groups. The most common signal intensity in the distal segment was isointensity in asymptomatic lesions, which was not the case in symptomatic lesions (P = .03).

Cerebral hemorrhage can be chronologically divided into different stages by MR imaging: hyperacute, acute, early subacute, late subacute, and chronic stage.^{25,26} Intraplaque hemorrhage of atherosclerosis on high-resolution MR imaging is also chronologically divided into fresh, recent, and old stages.²⁷ We proposed that the signal intensity of the distal segment may vary with time, similar to the signal intensity of cerebral hemorrhage and intraplaque hemorrhage. We selected different time intervals from the last onset of symptoms to the VWI examination in reference to cerebral hemorrhage and intraplaque hemorrhage. However, there was no difference in signal intensity at different time points. The same phenomenon has been observed in studies about intracranial largevessel arterial occlusion.²⁸ The reason may be that the exact time of the CCAO is uncertain and there is spontaneous recanalization and reformation of the thrombus in the occluded segment.

In this study, all the lesions in the asymptomatic group and 80% in the symptomatic group had a diffuse thickening of the

vessel wall. In a previous study, diffuse wall thickening in the petrous ICA was correlated with proximal ICA steno-occlusive disease.²⁹ With the increasing degree of stenosis, diffuse wall thickening became severe (present in 1.4%, 5.3%, 5.9%, and 80.4% of ipsilateral proximal ICAs, respectively, in patients with stenosis of 1%–49%, 50%–69%, 70%–99%, and total occlusion, respectively).²⁹ The probable cause is edema of the vessel wall due to long-term ischemia. Nutrients of the intima and inner media are supplied by diffusion from the lumen, and the adventitia is supplied by the vasa vasorum.³⁰ The vasa vasorum often arose directly from the lumen of the carotid artery and the arterial branching points.³¹ When the carotid artery was occluded, the vessel wall lost its nutrient supply and edema occurred.

DSA remains the criterion standard for the diagnosis of vascular disease; it can evaluate structural and hemodynamic functional information but cannot provide information on the vessel wall. VWI has been widely used in atherosclerosis of the carotid artery for its sufficient black-blood effects, excellent scan efficiency, high isotropic resolution and signal-to-noise ratio, and large spatial coverage.^{32,33} Compared with DSA, VWI can provide more information about the occluded segment. In this study, 72.9% of occluded segments detected by VWI were shorter than those detected by DSA. A previous study also found a similar result.¹⁶ The tiny lumen can be observed beyond the occluded segment on VWI, but it was considered occluded on DSA. We hypothesize that this was a pseudo-occlusion caused by collapse of the vessel wall due to inadequate compensatory proximal and distal blood flow. Endovascular interventional therapy has become a major treatment for CCAO. The stump condition and distal ICA reconstitution on DSA are important factors affecting successful recanalization.³⁴ It is necessary to identify the true occluded segment, and VWI may provide more information to help improve the success rate of recanalization.¹⁶

There were some limitations of this study. First, the MR imaging scanner was different in the 2 centers. Second, 3D T1weighted turbo spin-echo sequences were used to evaluate occluded segments without additional flow suppression, possibly affecting the evaluation of slow blood flow in the proximal-to-distal occluded segments. Third, 9 patients included in the asymptomatic group were detected occasionally, so selection bias may exist. Due to the small sample size, the findings in the present study need to be interpreted with caution. Fourth, only a portion of patients underwent enhanced scans, possibly not fully demonstrating the characteristics of the occluded segment. Finally, the conclusion of this study needs to be confirmed by a larger amount of data research in the future.

CONCLUSIONS

VWI can provide further insight into the occluded segment of CCAO. The signal intensity was demonstrated to be different between symptomatic and asymptomatic CCAO.

Disclosure forms provided by the authors are available with the full text and PDF of this article at www.ajnr.org.

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