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**SUPPLEMENTAL MATERIALS**

2 **Manuscript title:** Vessel wall MRI characteristics associated with intraprocedural

3 stent thrombosis during angioplasty for intracranial atherosclerotic stenosis

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## 1 **2. Supplemental Materials and Methods**

### 2 **2.2. High-resolution magnetic resonance vessel wall image acquisition**

3 All patients were scanned on a 3.0T MRI scanner (Philips Medical Systems, Best, The  
4 Netherlands) with standard 8-channel phased array head coils. The standardized  
5 imaging protocols included diffusion weighted imaging (DWI), three-dimensional  
6 (3D) time-of-flight magnetic resonance angiography (TOF-MRA), and HR-VWI  
7 sequences. TOF-MRA and DWI were acquired in a transverse plane by using the  
8 following parameters: TOF MRA-repetition time /echo time (TR/TE), 27/6.9 ms; flip  
9 angle, 20°; field of view (FOV), 240×160 mm; matrix size = 320 × 256; layer  
10 thickness = 1.6 mm. DWI- TR-TE = 2191/95 ms, layer thickness = 6.5 mm, matrix =  
11 200 mm × 204 mm. The maximum density projection of the TOF-MRA was used as  
12 the localization image of the HR-VWI sequences.

13 The HR-VWI sequences were then performed by using a volumetric isotropic  
14 turbo spin-echo acquisition (VISTA; Philips Healthcare, Best, The Netherlands) in a  
15 coronal plane (40-mm-thick slab) optimized for flow suppression and intracranial  
16 vessel wall delineation. The following parameters were used: TR/T3=800/18 ms;  
17 turbo spin-echo factor, 16 echoes; echo spacing, 6.1 ms; sensitivity encoding factor,  
18 two; number of signals acquired, 1-2; FOV, 200×180×40 mm; matrix, 332×302;  
19 acquired resolution, 0.6\*0.6\*0.6 mm (By interpolation, the acquired voxel is  
20 reconstructed into a size of 0.3\*0.3\*0.3 mm); acquisition time, 378s. A variable flip  
21 angle refocusing scheme was used with a minimum flip angle of 50° and a maximum  
22 flip angle of 120°, enabling high signal-to-noise efficiency and strong black blood  
23 effects. Radial k-space view ordering was used to optimize T1-weighted contrast.

24 Gadopentetate dimeglumine (Magnevist; Schering, Berlin, Germany) was  
25 administered intravenously (0.1 mmol per kilogram of body weight), and  
26 post-contrast 3D T1 imaging was repeated 5 minutes after contrast material  
27 administration. Scan parameters and sites were consistent with pre-contrast 3D T1  
28 scan.

### 29 **2.3. Image processing and analysis**

30 Manual segmentation of the lumen and outer wall boundaries of culprit plaque on MR

1 image was performed using MR Workspace (Philips Healthcare, Best, The  
2 Netherlands). Two experienced neurologists who were blinded to patient identifiers  
3 and clinical data measured image features twice independently.

4 Atherosclerotic plaque on MR images was defined as eccentric wall thickening  
5 with or without luminal stenosis identified on both the reconstructed pre- and  
6 post-contrast 3D T1 images. The two neurologists qualitatively graded plaque contrast  
7 enhancement based on its signal intensity on post-contrast 3D T1 images by using the  
8 corresponding pre-contrast series. The culprit plaque was identified by another two  
9 experienced neurologists based on clinical judgment. The plaque was considered a  
10 culprit plaque when it was 1) the only lesion in the vicinity of the stroke vessel or 2)  
11 the narrowest lesion in the presence of multiple plaques in the same vessel region  
12 when stroke occurred.

#### 13 **2.4. Perioperative management and stenting procedures**

14 We performed DSA for the responsible artery using a guiding catheter and determined  
15 the degree of stenosis according to the Warfarin-Aspirin Symptomatic Intracranial  
16 Disease (WASID) criteria. All patients were administered dual antiplatelet therapy  
17 (100 mg aspirin and 75 mg clopidogrel or cilostazol 200mg daily for at least 5 days  
18 before stenting) and received the interventional procedure under general anesthesia. In  
19 this study, all patients were used the same surgical approach and medical consumables.  
20 Pre-dilation was achieved using the Gateway balloon (Gateway-Wingspan system,  
21 Stryker, USA) with a balloon size of 80% of the normal segment diameter of the  
22 narrow distal. The Wingspan stent (Gateway-Wingspan system, Stryker, USA) was  
23 selected according to the diameter and length of the stenosis (stent extending at least 3  
24 mm on either side of the lesion). In this study, intraoperative angiography was  
25 performed at intervals of approximately 5 minutes and at least 30 minutes after  
26 stenting for detection of IPST. IPST was recorded when intraoperative angiography  
27 showed thrombus within or adjacent to the stent. Intra-arterial injection followed by  
28 intravenous tirofiban was used as salvage treatment for patients diagnosed with IPST.  
29 After endovascular treatment, all patients underwent computed tomography (CT)  
30 scans within 24 hours to detect bleeding or early cerebral infarction. Generally, dual

1 antiplatelet therapy (aspirin and clopidogrel or cilostazol orally) and statins  
2 (atorvastatin or rosuvastatin orally) were continued for 6 months post-procedure.  
3 After 6 months, one antiplatelet agent was discontinued.

#### 4 **2.5. Statistical analysis**

5 Clinical and imaging characteristics between patients in the IPST and non-IPST  
6 groups were compared. We presented mean  $\pm$  standard deviation (SD) or median  $\pm$   
7 interquartile range (IQR) for continuous variables and frequency (%) for categorical  
8 variables. The plaque characteristics between the IPST and non-IPST groups were  
9 compared using t-test for continuous variables with normal distribution, the  
10 Mann-Whitney U test for continuous variable with non-normal distribution, and the  
11  $\chi^2$  test or Fisher's exact test for categorical variables. Patients with IPST (all male)  
12 and those without IPST were matched using a 1:2 matching algorithm with age and  
13 sex as covariates. Univariate logistic regression analysis was performed for screening  
14 clinical and imaging factors associated with IPST at  $P < 0.05$ , and then all variables  
15 with  $P < 0.05$  in the univariate analysis were considered candidates for stepwise  
16 logistic regression analysis, where the entry-level probability was set at 0.05, and the  
17 removal level was set at 0.10. Two-tailed  $P$  value  $< 0.05$  was considered statistically  
18 significant. Multivariable binary logistic regression analysis was used to detect  
19 clinical and imaging factors associated with IPST.

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1 **Table S1.** Characteristics of study population

Characteristics	IPST group (n = 8)	Non-IPST group (n = 16)	P value
Age (years), mean (SD)	58.7 (2.47)	58.7 (1.69)	0.999
Male, n (%)	8 (100)	16 (100)	–
Hypertension, n (%)	6 (75.00)	10 (62.50)	0.549
DM, n (%)	2 (25.00)	3 (18.75)	0.728
Dyslipidemia, n (%)	3 (37.50)	7 (43.75)	0.770
LDL, mean (SD)	2.19 (0.60)	1.89 (0.55)	0.235
HDL, mean (SD)	1.08 (0.19)	1.00 (0.15)	0.303
TG, mean (SD)	3.67 (0.78)	3.33 (0.71)	0.299
TC, mean (SD)	1.60 (0.60)	1.57 (0.54)	0.910
Ischemic stroke history, n (%)	4 (50.00)	6 (37.50)	0.558
NIHSS, mean (SD)	0.75 (1.39)	2.00 (2.22)	0.162
Medication, n (%)			
Aspirin	8 (100.00)	16 (100.00)	–
Clopidogrel	8 (100.00)	16 (100.00)	–
Aspirin resistance and/or Clopidogrel resistance, n (%)	0 (0.00)	0 (0.00)	–
Smoker, n (%)	4 (50.00)	12 (75.00)	0.231
Alcohol abuse, n (%)	7 (62.50)	13 (81.25)	0.328
Stent location-anterior, n (%)	3 (37.50)	7 (43.75)	0.770
Diameter stenosis degree	77.7(4.3)	75.5(5.5)	0.341
Time interval between HR-VWI and intracranial stenting, mean (SD)	2.9 (1.5)	2.8 (1.6)	0.542

2 IPST, intraprocedural stent thrombosis; DM, diabetes mellitus; LDL, low-density  
3 lipoprotein; HDL, high-density lipoprotein; TG, triglyceride; TC, triglyceride; NIHSS,  
4 National Institutes of Health Stroke Scale.

5 Continuous variables were represented as median (interquartile range) and mean  
6 (standard deviation). Categorical variables were represented as frequency and  
7 percentage.

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