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Original research

Histological evaluation of acute ischemic stroke thrombi may indicate the occurrence of vessel wall injury during mechanical thrombectomy

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ABSTRACT

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To cite: Mereuta OM, Abbasi M, Fitzgerald S, et al. J NeuroIntervent Surg Epub ahead of print: [please include Day Month Year]. doi:10.1136/ neurintsurg-2021-017310 **Background** Several animal studies have demonstrated that mechanical thrombectomy (MT) for acute ischemic stroke (AIS) may cause vessel wall injury (VWI). However, the histological changes in human cerebral arteries following MT are difficult to determine.

Objective To investigate the occurrence of VWI during MT by histological and immunohistochemical evaluation of AIS clots.

Methods As part of the multicenter STRIP registry, 277 clots from 237 patients were analyzed using Martius Scarlett Blue stain and immunohistochemistry for CD34 (endothelial cells) and smooth muscle actin (smooth muscle cells).

Results MT devices used were aspiration catheters (100 cases), stentriever (101 cases), and both (36 cases). VWI was found in 33/277 clots (12%). There was no significant correlation between VWI and MT device. The degree of damage varied from grade I (mild intimal damage, 24 clots), to grade II (relevant intimal and subintimal damage, 3 clots), and III (severe injury, 6 clots). VWI clots contained significantly more erythrocytes (p=0.006*) and less platelets/ other (p=0.005*) than non-VWI clots suggesting soft thrombus material.

Thrombolysis correlated with a lower rate of VWI ($p=0.04^*$). VWI cases showed a significantly higher number of passes (2 [1–4] vs 1 [1–3], $p=0.028^*$) and poorer recanalization outcome ($p=0.01^*$) than cases without VWI.

Conclusions Histological markers of VWI were present in 12% of AIS thrombi, suggesting that VWI might be related to MT. VWI was associated with soft thrombus consistency, higher number of passes and poorer revascularization outcome. There was no significant correlation between VWI and MT device.

INTRODUCTION

Mechanical thrombectomy (MT) for acute ischemic stroke (AIS) has a risk of arterial wall damage that may lead to long-term side effects, such as intima hyperplasia, de novo stenosis, and occlusions and arterial dissection.^{1–3} The injury may occur in relation to the forces exerted by the device against the vessel wall,¹ the occurrence of vasospasm,² vascular anatomy,³ and the presence of underlying stenotic conditions, such as atherosclerotic plaque^{4 5} or calcification of the vessel wall.⁶

Several animal studies have evaluated the histological and angiographic changes in the vessel wall due to potential damage during MT but the results have been inconsistent.^{1 5 7-11} Initial studies found that Trevo and Solitaire devices can cause severe disruption of the intima^{8 9} and vascular damage extending to the medial layer.¹⁰ Another study that compared an aspiration device (Penumbra) with stentrievers (Merci, Catch, and Solitaire) demonstrated that all devices caused endothelial denudation and disruption of the internal elastic lamina, but the aspiration device was associated with more intimal and medial layer edema.¹¹ However, a recent histological and ultrastructural study comparing direct aspiration first pass technique with stentrievers (Solitaire) showed that stentrievers can be more harmful to all layers of the vessel wall-in particular, the endothelium.¹

Moreover, based on angiographic follow-up, Kurre *et al*² observed a decrease in arterial diameter at the site at which the thrombectomy was performed. A further study reported the appearance of a concentric arterial wall thickening and enhancement after MT using high-resolution contrast-enhanced vessel wall MRI.¹²



Ischemic stroke

However, little is known about the extent and mechanism of vessel wall injury (VWI) caused by MT in human cerebral arteries. Therefore, the aim of our study was to investigate the presence of histological markers that may be indicative of VWI during MT by histological and immunohistochemical evaluation of endothelial cells, smooth muscle cells, and subendothelial connective tissue in AIS clots.

MATERIALS AND METHODS Clot collection

This study was performed as part of the multi-institutional Stroke Thromboembolism Registry of Imaging and Pathology (STRIP). The study was approved by the institutional review board and was compliant with the Health Insurance Portability and Accountability Act. A waiver of consent was granted. Patients were included in the study if they were >18 years, had undergone mechanical thrombectomy treatment for AIS, and had embolic material available for analysis.

Data regarding suspected stroke etiology, occlusion location, and clot length on non-contrast computed tomography prior to MT, approach used for MT, number of procedural passes required to remove the clot, and final modified Thrombolysis in Cerebral Infarction (mTICI) score were self-reported at each center and captured on the STRIP registry data abstraction form. Stroke etiology was classified using the Trial of Org 10172 in Acute Stroke Treatment (TOAST) system: large artery atherosclerosis, cardioembolic stroke of other determined etiology, and cryptogenic.¹³

Histological analysis of thrombi and extracted clot area measurement

On retrieval, each clot was immediately fixed in 10% phosphate-buffered formalin. Clots were shipped to the histology core facility. On arrival at the core facility, gross photos were taken of each clot. All clots were then processed using a standard tissue processing protocol, embedded in paraffin and cut into 3 μ m sections. Two representative sections for each clot were stained with Martius Scarlett Blue (MSB) to identify the main components of clots (red blood cells, white blood cells, fibrin, platelets/other) as shown in figure 1.^{14 15}

Stained slides underwent whole slide scanning (Motic Easyscan Pro, Motic Digital Pathology) at $\times 20$ magnification. MSB quantification of clot components was performed using Orbit Image Analysis software (www.orbit.bio).¹⁶ ¹⁷ ImageJ software (https://imagej.nih.gov/ij/) was utilized to measure total extracted clot area (ECA) of each clot using thegross image of the clot. We used the equations proposed by Duffy et al to calculate the overall contribution of each component to whole thrombus material.¹⁸



Figure 1 Thrombus histology and immunohistochemistry in representative cases selected by the degree of vessel wall injury and mechanical thrombectomy approach. Grade I: aspiration catheter (A), stentriever (B), and combination of devices (C); grade II: stentriever (D); grade III: aspiration catheter (E), stentriever (F), and combination (G). Martius Scarlett Blue (MSB) staining identifies the standard components of clots: red blood cells (yellow), fibrin (red). and platelets/other (light pink). Collagen can be also identified by MSB (E–G: light blue and arrows). Immunostaining for CD34 and smooth muscle actin (SMA) identifies endothelial cells and smooth muscle cells (purple), respectively. Areas within the squares in the MSB images are magnified in the immunostaining images. Note that clustered endothelial cells are distributed along the edge of clot fragments in grade I injury (A–C) while defined clusters of CD34- and SMA-positive cells are located at the periphery of the clot in grade II (D) and are associated with collagen in grade III (E–G). Scale bar (MSB)=5 mm (A, C), 3 mm (B, D), 3 mm and 300 µm (E–G); scale bar (immunostaining)=700 µm (A), 500 µm (B), 600 µm (C), and 300 µm (D–G).

Ischemic stroke

Immunohistochemical analysis of thrombi

Immunostaining was used to identify endothelial cells (monoclonal mouse anti-human CD34, clone QBEnd-10, Dako M7165), smooth muscle cells (monoclonal mouse anti-human smooth muscle actin, SMA, clone 1A4, Dako M0851), and von Willebrand Factor (monoclonal mouse anti-human vWF, Dako M0616). Positive controls (artery tissue and human tonsil) and negative controls (omitting primary antibodies) were used. Immunohistochemistry was performed on a Leica Bond Max autostainer using a RedMap kit (Bond Polymer Refine Red Detection, Leica Biosystems).

The grading system showing the degree of underlying intimal damage associated with thrombi retrieved by MT was previously established and published by a board-certified neuropathologist.[19] The grading system used in our study was slightly modified by adding SMA immunohistochemistry to identify smooth muscle cells, another component of vessel wall.

The slides were examined by an experienced pathologist blinded to clinical data, and a grading system to classify the vessel wall damage was established (figure 1): grade I – clustered endothelial cells (CD34-positive cells) at the periphery of the clot: mild intimal damage; grade II – clustered endothelial cells associated with smooth muscle cells (SMApositive cells) at the periphery of the clot: relevant intimal and subintimal damage; grade III - clustered endothelial cells and/or smooth muscle cells associated with subendothelial connective tissue (collagen) at the margin of the clot: severe injury of the vessel wall. The presence of single endothelial cells or small clusters of endothelial cells was probably related to the physiological turnover of the endothelium.¹⁹

Statistical analysis

Data were analyzed using IBM SPSS 25 software. Graphpad Prism 8 was used to generate graphs. A Kolmogorov-Smirnov test indicated that quantitative variables did not follow a standard normal distribution and therefore, the non-parametric Kruskal-Wallis test was applied to make comparisons. Correlations between categorical variables were assessed using χ^2 test. A level of statistical significance for all analyses was set at p<0.05 (two-sided). Results were reported as mean±SD, median (IQ1– IQ3), or number (% of cases), as appropriate.

RESULTS

Clinical data

Two-hundred and thirty seven patients who underwent MT between October 2016 and November 2019 for large vessel occlusion in the anterior or posterior cerebral circulation were included in the study. Two hundred and seventy-seven clots were retrieved. Aspiration was used in 100 cases, stentriever in 101 cases, and a combined approach of stentriever and aspiration device was used in 36 cases. Suspected stroke etiologies were large artery atherosclerosis (43 cases, 18.1%), cardioembolic (100 cases, 42.2%), other determined etiology (19 cases, 8%), and cryptogenic (58 cases, 24.5%). The etiology was not reported in 17 cases (7.2%). Eighty patients (33.8%) were treated with intravenous recombinant tissue plasminogen activator (IV rtPA). Two hundred and seventeen patients (91.6%) had an occlusion in the anterior circulation, 16 cases (6.8%) in the posterior circulation, and one case (0.4%) had a dual occlusion involving both territories. The clots were extracted at first attempt in 130 cases (54.9%). Multiple procedural passes were required in 107 cases

(45.1%). A final mTICI score 2b/3 was achieved in 214 patients (90.3%).

Histopathologic findings

The overall contribution of each component based on the quantification of MSB staining was $52.16 \pm 16.88\%$ red blood cells (RBCs), $29.15 \pm 12.83\%$ fibrin, $15.17 \pm 14.6\%$ platelets/other and $3.48 \pm 2.57\%$ white blood cells for the VWI clots. Moreover, VWI clots contained significantly more RBCs ($52.16 \pm 16.88\%$ vs $41.78 \pm 21.32\%$, p=0.006*) and less platelets/other ($15.17 \pm 14.6\%$ vs $23.12 \pm 17.88\%$, p=0.005*) than non-VWI clots suggesting soft thrombus material. Total ECA was not significantly different between VWI and non-VWI group ($43.17 \pm 36.28 \text{ mm}^2$ vs $62.75 \pm 82.52 \text{ mm}^2$, p=0.233). Data are summarized in Supplementary Table 1.

Degree of vessel wall damage

We found evidence of VWI in 33 clots (12%). The degree of damage was categorized as follows: grade I – 24 clots (72.7%), grade II – three clots (9.1%), and grade III – six clots (18.2%).

Grade I damage was the predominant type of injury. We observed that grade I injury was characterized by the presence of clustered endothelial cells (CD34-positive cells, purple) distributed along the edge of the clot (figure 1A–C), whereas VWI grade II (figure 1D) and grade III (figure 1E–G) were represented by focal lesions with clusters of endothelial cells, smooth muscle cells (SMA-positive cells, purple), and collagen (light blue on MSB staining, arrows) at the periphery of the clot.

In addition, small foci of CD34- and SMA-positive cells were present inside fragments in two other cases, suggesting early organization of the clots (online supplemental figure 1A–C). A pavement-like surface endothelium representing surface endothelialization was also observed in another case. Dissection of the vertebral artery occurred in one patient.

CD34 is not a specific marker for endothelial cells and can be also expressed by hematopoietic stem cells. To deal with this problem, we performed immunohistochemistry for vWF, showing that CD34-positive endothelial cells are also vWFpositive (online supplemental figure 1D,E).

We acknowledge that VWI markers could be also present in a chronic thrombus material and may arise due to a vessel pathology, such as a ruptured atherosclerotic plaque. Therefore, we excluded the presence of foam cells, calcium deposits, fibrous caps or recanalization vessels in VWI clots. In addition, the conventional histopathologic evaluation confirmed that all 33 VWI thrombi were fresh thrombi mainly composed of RBCs and fibrin. These findings indicate that the presence of the endothelial cells, smooth muscle cells and collagen within fresh thrombi was caused by the vessel wall tearing associated with the MT procedure.

Table 1	Distribution of vessel wall injury (VWI) degree of damage			
based on mechanical thrombectomy device used.				

	No of patients (%)			
VWI grade	Aspiration	Stentriever	Combination	
Grade I	13 (39.4)	8 (24.2)	3 (9.1)	
Grade II	1 (3)	2 (6.1)	-	
Grade III	2 (6.1)	2 (6.1)	2 (6.1)	



Figure 2 Number of procedural passes and revascularization outcome in cases associated with vessel wall injury (VWI). (A) Violin plots show that patients with VWI had a significantly higher number of passes than non-VWI cases. (B) Graphical representation of final modified Thrombolysis in Cerebral Infarction (mTICI) score for the two groups of patients. VWI was associated with a significantly lower mTICI score in patients with no injury than in patients without VWI.

Degree of vessel wall damage and MT devices

As shown in table 1, an aspiration catheter was used in 16 cases (48.5%), a stentriever device in 12 cases (36.4%), and a combined approach in five cases (15.2%). Within the stentriever only group, the stentriever device was changed during the procedure in one case. The same aspiration catheter was used during the entire procedure in each case within the aspiration only group. Within the combination group, the technique was changed using the same stentriever/aspiration catheter during the procedure in each case. There was no significant difference between MT technique with regard to presence of VWI (p=0.7) or the degree of injury (p=0.5).

Thrombolytic treatment

The IV rtPA administration was correlated with a lower rate of VWI. The rate of VWI was 8% (6/80 cases) for the rtPA group and 17% (27/157 cases) for non-rtPA group (χ^2 =3.93, p=0.04*).

Number of passes and revascularization outcome

As represented in figure 2A, the cases associated with VWI (n=33) showed a significantly higher number of procedural passes than non-VWI cases (n=204; two passes (1-4) vs one pass(1-3) H1=4.83, p=0.028*). Moreover, the degree of injury significantly increased with the number of passes (H2=6.72, $p=0.035^*$).

As a limitation of our study, the clots were not collected per pass in all cases and therefore, we could not appreciate in some cases the exact pass when VWI occurred. Nevertheless, more than 3 passes were required to extract the clots in 11 cases with VWI. In particular, in four cases VWI occurred at pass 4, 5, 7 and 11.

Figure 2B highlights that VWI-positive cases (n=33) had a significantly poorer recanalization outcome than non-VWI patients (n=202; H1=6.62, p=0.01*). In particular, there was a significant difference between bad/successful (mTICI \leq 2c) and complete revascularization (mTICI 3) outcome for the incidence of VWI (21%, 23/112 cases vs 9%, 10/123 cases, χ^2 =6.48, p=0.01*).

Clinical and procedural information for the two groups of patients are summarized in table 2.

DISCUSSION

In this study, we performed a comprehensive histological and immunohistochemical analysis of clots retrieved by MT aimed to identify the presence of vessel wall components, such as endothelial cells, smooth muscle cells, and subendothelial connective tissue which may indicate a VWI caused by MT devices. We found evidence of VWI in 12% of the clots (14% of the cases investigated). The degree of damage varied from grade I (mild intimal damage with CD34-positive cells extended along the edge of the clot), representing the predominant type of injury, to grade II (relevant intimal and subintimal damage), and grade III (severe wall damage) characterized by focal clusters at the margin of the clot containing endothelial cells, smooth muscle cells, and collagen, respectively. There was no significant correlation between VWI and MT technique used, although aspiration was predominantly associated with VWI grade I. IV rtPA administration correlated with a lower rate of VWI. We also found that VWI clots contained significantly more RBCs and less platelets/ other compared to non-VWI clots. VWI cases were associated with a significantly higher number of passes and lower mTICI score than the cases without VWI.

VWI has been reported in animal models and may differ based on the MT device used.¹⁵⁷⁻¹¹ The presence of VWI histological markers in AIS emboli suggests that VWI may be caused by the MT device during the intervention procedure. However, it is also possible that they may have arisen due to a vessel pathology, such as a ruptured atherosclerotic plaque. Histological evaluation of thrombi may provide new insights into the complex interaction between the vessel wall, thrombus, and MT devices.

Two previous studies have assessed the occurrence of VWI using histological analysis of retrieved thrombi.^{19 20} In one study, CD34-positive endothelial cells were identified in 11 out of 48 thrombi (23%) removed with a stentriever, but no subendothelial connective tissue was found. Retrieved clots were classified as fibrin dominant (13 clots), RBCs-dominant (15 clots and mixed (20 clots).¹⁹ In a recent study, in a total of 150 thrombi from 101 patients treated with Penumbra aspiration catheters (42 patients), stentrievers (21 patients), or both techniques (38 patients), collagen fibers and internal elastic lamina were shown in 16% and 8% of specimens, respectively. No difference between aspiration and stentrievers was observed. A low proportion of red blood cells, a high frequency of M2 segment of middle cerebral artery/P2

non-VWI cases). We identified the presence of collagen fibers
in six cases (VWI grade III).
Interestingly, in our study, VWI clots contained more RBCs
and less platelets/other than non-VWI clots suggesting soft consis-
tency, a finding that argues with the previous study showing
that a stiffer clot is associated with higher number of passes and
consequently, with VWI. The number of VWI cases in our study
is small which may be a factor. On the other hand, Funatsu et
al reported that vessel wall components included fibrous caps of
plaque. We excluded in our cases the presence of fibrous caps or
other plaque components associated with a chronic thrombus
material which would indicate a LAA origin rather than a VWI
during MT. In our cohort, 7 cases with VWI had LAA etiology. It
has been recently reported that LAA clots are richer in RBCs in
passes 1-3 than all other suspected etiologies whereas fibrin and
platelet/other content increases in clots removed in later passes.
²¹ We can only assume that VWI in LAA cases may have occurred
during early procedural passes.
We also found that IV rtPA administration correlated with a
lower rate of VWI. A recent study demonstrated that throm-
bolysis significantly reduced the clot size, ²² which may help to
retrieve the clot without damaging the occluded vessel. Further-
more, a recent three-dimensional scanning electron microscopy
analysis of AIS emboli showed that rt-PA treatment resulted in
thinning of fibrin fibers and disruption of fibrin network which
may explain the reduced clot size. ²³
A recent study in an animal model reported more intima and
medial layer damage after stentriever devices than with aspira-
tion. Several factors may account for this difference, such as the
mechanism of clot removal. Aspiration exerts a force to the prox-
imal base of the clot, whereas stentrievers apply a continuous
radial force against the vessel wall which may lead to a deeper
damage. ¹ Moreover, the radial force varies greatly between sten-
trievers, ²⁴ and the extent of vascular injury may increase with
the increasing radial force of the stentriever. ¹⁰ Using an in vitro
live cell artificial vessel system, Teng et al demonstrated that the
degree of endothelial injury was related to the vessel diameter in
a device-specific manner, with different patterns of endothelial
damage caused by different MT devices. ²⁵ In the present study,
VWI grade I was observed in aspiration-retrieved clots more
than with a stentriever and was characterized by the presence of
endothelial cells extended along the edge of the clot suggesting a
large circumferential injury. VWI grades II and III were described
more in clots retrieved with stentrievers/combination of devices,

at each site, which might have resulted in some site-to-site vari-

ability. Owing to the lack of angiographic data, we were not able to evaluate the occurrence of vasospasm or the presence of an underlying stenosis. Also, a high-resolution MRI scan may

provide a better visualization of vessel wall changes. We did

not assess in our study other potential factors that could mini-

mize VWI including adjunctive devices such as balloon guiding

catheters, anesthesia type, etc. Finally, we included in this study

various devices characterized by different radial forces and

designs. Further studies could define specific patterns of VWI

based on the specific MT device used.

 Table 2
 Clinical characteristics of patients grouped by vessel wall
 non-VWI cases). We in six cases (VWI gr

Vessel wall iniury

Characteristic	No injury, 204 (86)	33 (14)	to a set of the second south the manifest standard and
MT device			that a stiffer clot is associated with higher number of passes
Aspiration only	84 (41.2)	16 (48.5)	consequently with VWI. The number of VWI cases in our s
Stentriever only	89 (43.6)	12 (36.4)	is small which may be a factor. On the other hand, Funate
Combination	31 (15.2)	5 (15.2)	al reported that vessel wall components included fibrous ca
Stroke Etiology			plaque. We excluded in our cases the presence of fibrous cap
Large artery atherosclerosis	36 (17.6)	7 (21.2)	other plaque components associated with a chronic thror
Cardioembolic	88 (43.1)	12 (36.4)	material which would indicate a LAA origin rather than a
Other determined etiology	16 (7.8)	3 (9.1)	during M1. In our cohort, 7 cases with VWI had LAA etiolog
Cryptogenic	49 (24)	9 (27.3)	passes 1-3 than all other suspected etiologies whereas fibrin
Not available	15 (7.4)	2 (6.1)	platelet/other content increases in clots removed in later pa
IV rtPA			²¹ We can only assume that VWI in LAA cases may have occu
Yes	74 (36.3)	6 (18.2)	during early procedural passes.
No	130 (63.7)	27 (81.8)	We also found that IV rtPA administration correlated w
Statistical analysis	N=237. H1=3.93. p=0.0	4*	lower rate of VWI. A recent study demonstrated that the
Occlusion location			bolysis significantly reduced the clot size, ²² which may hel
MCA	145 (71 1)	20 (60 6)	retrieve the clot without damaging the occluded vessel. Fur
	26 (12 7)	7 (21 2)	more, a recent three-dimensional scanning electron microse
Vortobrobacilar	12 (5 0)	1 (2)	analysis of AIS emboli snowed that rt-PA treatment resulte thinning of fibrin fibers and discussion of fibrin network w
Tandom occlusion	12 (5.3)	5 (15 2)	may explain the reduced clot size 23
Other dual acclusion*	1 (0.5)	5 (15.2)	A recent study in an animal model reported more intima
	1 (0.5)	-	medial layer damage after stentriever devices than with as
	4 (2)	-	tion. Several factors may account for this difference, such as
Net eveilet le	2 (1)	-	mechanism of clot removal. Aspiration exerts a force to the p
	3 (1.5)	-	imal base of the clot, whereas stentrievers apply a continu
No of passes		12 (26.4)	radial force against the vessel wall which may lead to a de
-	118 (57.8)	12 (36.4)	damage. ⁴ Moreover, the radial force varies greatly between s
2	32 (15.7)	10 (30.3)	the increasing radial force of the stentriever ¹⁰ Using an in y
3	19 (9.3)	1 (3)	live cell artificial vessel system. Teng <i>et al</i> demonstrated that
4	11 (5.4)	2 (6.1)	degree of endothelial injury was related to the vessel diamet
≥5	24 (11.8)	8 (24.2)	a device-specific manner, with different patterns of endoth
Median (IQ1-IQ3)	1 (1-3)	2 (1-4)	damage caused by different MT devices. ²⁵ In the present st
Statistical analysis	N=237, H1=4.83, p=0.028*		VWI grade I was observed in aspiration-retrieved clots r
Final mTICI score	•		than with a stentriever and was characterized by the presence
0	6 (2.9)	4 (12.1)	endothelial cells extended along the edge of the clot suggest
1	5 (2.5)	-	large circumferential injury. VWI grades II and III were descr
2a	4 (2)	2 (6 1)	showing defined clusters of endothelial cells smooth m
2b	56 (27.4)	10 (30.3)	cells, and subendothelial connective tissue at the margin of
20	18 (8.8)	7 (21.2)	clot. This pattern may suggest focal endothelial denudation
3	113 (55 4)	10 (30 3)	alteration of tunica media/adventitia.
S Not available	2 (1)	-	Our study has several limitations. Chemical and phy
	2 (1)		manipulation of clots might affect their internal organization

No of patients (%)

injury status

artery; ICA, internal carotid artery; PCA, posterior cerebral artery; ACA, anterior cerebral artery; mTICI Score, modified Thrombolysis in Cerebral Infarction Score

segment of posterior cerebral artery reached by devices, and a high number of passes were associated with the presence of vascular wall components.²⁰ We also found that VWI was associated with a significantly higher number of passes. Moreover, the recanalization outcome was poorer in VWI cases. However, the incidence of VWI in our study was lower and the occlusion was predominantly located in the M1 segment

Ischemic stroke

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CONCLUSIONS

Histological markers of vascular wall damage were identified in 12% of AIS thrombi. Their presence may indicate that VWI occurred during MT. There was no significant correlation between VWI and the MT device used. Different degrees of damage showed characteristic patterns of injury. VWI was associated with soft thrombus consistency, a higher number of procedural passes and poorer revascularization outcome. Taken together, our findings may improve our understanding of the factors that might affect MT effectiveness in order to achieve fast and complete recanalization without damaging the occluded vessel.

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Contributors OMM and MA contributed equally to this study. OMM, MA, and WB were involved in all stages of the manuscript from concept design to drafting the manuscript. SF, DD, and RK were responsible for collecting and recording the clinical and procedural information from patients and were involved in the histological analysis of clots. This study is the result of a multicenter registry requiring the collaboration of multiple neurointerventionalists, research fellows, and histopathologists and all authors meet the ICJME criteria for authorship. All authors reviewed, edited, and approved the final manuscript prior to submission.

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Competing interests RK reports NIH funding (R01 NS076491, R43 NS110114, and R44 NS107111), is a research consultant for Cerenovus, Insera Therapeutics LLC, Marblehead Medical LLC, Microvention Inc, MIVI Neuroscience Inc, Neurogami Medical Inc, and Triticum Inc, and has stock in Neurosigma Inc (money paid to institution). AJY receives research support from Medtronic, Cerenovus, Penumbra, and Stryker, and is a consultant for Penumbra, Cerenovus, Zoll Circulation, and Vesalio. He is on the Scientific Advisory Board of XCath and Nico-lab, and has equity interest in Insera Therapeutics LLC. JEDA declares competing interests in the form of employment (modest compensation) from Medtronic and Penumbra. AMD received honoraria from Medtronic for continuing medical education events. RGN declares competing interests in the form of Stryker (DAWN Trial (DWI or CTP Assessment With Clinical Mismatch in the Triage of Wake-Up and Late Presenting Strokes Undergoing Neurointervention With TREVO) principal investigator, no compensation; TREVO Registry Steering Committee, no compensation; TREVO-2 Trial

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Ischemic stroke

SUPPLEMENTARY MATERIAL

SUPPLEMENTARY TABLES:

Supplementary Table 1. Histological composition of retrieved thrombi.

	No injury	Vessel wall injury	Kruskal-Wallis test
Red blood cells (%)	41.78 ± 21.32	52.16 ± 16.88	N=237, H1=7.556, p=0.006*
White blood cells (%)	3.27 ± 2.19	3.48 ± 2.57	N=237, H1=0.027, p=0.869
Fibrin (%)	31.18 ± 16.25	29.15 ± 12.83	N=237, H1=0.148, p=0.701
Platelets/other (%)	23.12 ± 17.88	15.17 ± 14.6	N=237, H1=8.054, p=0.005*
Total ECA (mm ²)	62.75 ± 82.52	43.17 ± 36.28	N=228, H1=1.425, p=0.233

SUPPLEMENTARY FIGURES:



Supplementary Figure 1. Morphological features of thrombi in two representative cases. (A) Martius Scarlett Blue (MSB) staining identifies the main components of clots: red blood cells (yellow), fibrin (red) and platelets/other (light pink). Clot area in the square is showed at higher magnification in the immunostaining images (B, C). Small foci of CD34-positive endothelial cells (B) and smooth muscle actin (SMA)-positive cells (C, purple) were present inside a fragment suggesting early organization of the clot. Immunohistochemistry for CD34 (D) and von Willebrand Factor (E) demonstrates that CD34-positive endothelial cells were also positive for vWF (purple). Scale bar = 3 mm (A), 700 μ m (B, C) and 300 μ m (D, E).