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Sex differences in clot, vessel and tissue characteristics in patients with a large vessel occlusion treated with endovascular thrombectomy

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Abstract

Introduction: To improve our understanding of the relatively poor outcome after endovascular treatment (EVT) in women we assessed possible sex differences in baseline neuroimaging characteristics of acute ischemic stroke patients with large anterior vessel occlusion (LVO).

Patients and methods: We included all consecutive patients from the MR CLEAN Registry who underwent EVT between 2014 and 2017. On baseline non-contrast CT and CT angiography, we assessed clot location and clot burden score (CBS), vessel characteristics (presence of atherosclerosis, tortuosity, size, and collateral status), and tissue characteristics with the Alberta Stroke Program Early Computed Tomography Score (ASPECTS). Radiological outcome was assessed with the extended thrombolysis in cerebral infarction score (eTICI) and functional outcome with the modified Rankin Scale score (mRS) at 90 days. Sex-differences were assessed with multivariable regression analyses with adjustments for possible confounders.

Results: 3180 patients were included (median age 72 years, 48% women). Clots in women were less often located in the intracranial internal carotid artery (ICA) (25% vs 28%, odds ratio (OR) 0.85; 95% confidence interval: 0.73–1.00). CBS was similar between sexes (median 6, IQR 4–8). Intracranial (aOR 0.73; 95% CI: 0.62–0.87) and extracranial (aOR

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0.64;95% CI:0.43–0.95) atherosclerosis was less prevalent in women. Vessel tortuosity was more frequent in women in the cervical ICA (aOR 1.89;95% CI:1.39–2.57) and women more often had severe elongation of the aortic arch (aOR 1.38;95% CI:1.00–1.91). ICA radius was smaller in women (2.3 vs 2.5 mm, mean difference 0.22;95% CI:0.09–0.35) while M1 radius was essentially equal (1.6 vs 1.7 mm, mean difference 0.09;95% CI:–0.02–0.21). Women had better collateral status ($\geq 50\%$ filling in 62% vs 53% in men, aOR 1.48;95% CI:1.29–1.70). Finally, ASPECT scores were equal between women and men (median 9 in both sexes, IQR 8–10 vs 9–10). Reperfusion rates were similar between women and men (acOR 0.94;95% CI:0.83–1.07). However, women less often reached functional independence than men (34% vs 46%, aOR 0.68;95% CI:0.53–0.86).

Discussion and conclusion: On baseline imaging of this Dutch Registry, men and women with LVO mainly differ in vessel characteristics such as atherosclerotic burden, extracranial vessel tortuosity, and collateral status. These sex differences do not result in different reperfusion rates and are, therefore, not likely to explain the worse functional outcome in women after EVT.

Keywords

Sex differences, endovascular treatment, ischemic stroke, large vessel occlusion, neuroimaging

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Introduction

The lifetime prevalence of ischemic stroke is higher in women than in men and women more often have an unfavorable outcome.^{1–6} Endovascular treatment (EVT) is highly effective for anterior large vessel occlusion (LVO) related stroke and doubles the probability of favorable functional outcome.⁷ Most clinical trials reported similar treatment effects and functional outcomes after EVT in men and women, whereas several cohort and registry studies found poorer outcomes in women.^{8–12} Until now, reports on sex differences in large patient series treated in routine clinical practice are lacking.

Little is known about sex differences in neuroimaging characteristics in the acute phase of LVO stroke, in relation to outcome after EVT. Most data are derived from randomized trials or small cohorts of treated patients. Women have a higher incidence of atrial fibrillation, which might lead to larger clots in more proximal locations.^{13,14} Also, some studies suggest that women more often have intracranial atherosclerosis, which could impair endovascular access and increase the chance of distal emboli during the procedure.¹⁵ Furthermore, there might be sex differences in vessel tortuosity, vessel size, collateral status, or early ischemic tissue changes.^{16–20}

More insight in possible sex differences in neuroimaging features is needed to understand the possible poorer functional outcome after EVT in women and to optimize personalized treatment in patients with LVO. We aimed to investigate sex differences in clot, vessel and tissue characteristics and assessed radiological and functional outcome after EVT in a large real-world cohort of LVO patients.

Methods

Study design

The MR CLEAN (Multicenter Randomized Clinical Trial of Endovascular Treatment for Acute Ischemic Stroke in

The Netherlands) Registry is a national, prospective, open, multicenter, observational monitoring study in which all 16 stroke intervention centers in the Netherlands that perform EVT participated.²¹ The registration started after the final inclusion in the original MR CLEAN trial and was set up to study safety and effectiveness of EVT in routine clinical practice. Consecutive patients in the Netherlands were included in the MR CLEAN Registry from March 2014 until January 2019. Data of patients who were treated up to 1 November 2017 were processed and used in the current analysis.

Study population

Inclusion criteria were an age of 18 years and older, arterial puncture within 6.5 h of symptom onset, and LVO of the intracranial anterior circulation (intracranial carotid artery, anterior cerebral artery, or middle cerebral artery). EVT was defined as receiving arterial puncture with the aim to perform thrombectomy. The method of EVT was left to the discretion of the treating physician.

Baseline characteristics

For this analysis, the following baseline clinical characteristics were collected: age, cardiovascular risk factors, medication, pre-stroke modified Rankin Scale (mRS) score, admission National Institutes of Health Stroke Scale (NIHSS) score, blood pressure on admission, and treatment with intravenous thrombolysis (IVT) prior to EVT. Furthermore, time from stroke onset to hospital admission, to IVT, to groin puncture, and to reperfusion was recorded.

Neuroimaging characteristics

All patients underwent non-contrast brain CT followed by a CT angiography (CTA) or MRA and Digital Subtraction Angiography (DSA). The neuroimaging characteristics of

interest were divided into three main categories: (1) clot, (2) vessel, and (3) tissue characteristics. These categories were further divided into subcategories.

Clot characteristics were subdivided into clot location and clot extent. Clot location was defined as the most proximal occlusion segment (intracranial internal carotid artery (ICA) / internal carotid artery terminus (ICA-T), middle cerebral artery segment 1 (M1), M2, other). Clot extent was measured on CTA by the clot burden score (CBS). This is a grading scale from 0 to 10, that quantifies the extent of the intracranial thrombus by assigning points for the presence of contrast on imaging within the intracranial ICA, M1, M2, and anterior cerebral artery (A1) vessel segments. A score of 10 indicates that no clot is present and a score of 0 indicates complete occlusion of all vessel segments of the proximal intracranial anterior circulation.²² We calculated the median CBS and the proportion of participants with a CBS < 7, which is related to poor functional outcome.²³

Vessel characteristics were subdivided into atherosclerosis including stenosis, vessel tortuosity, vessel size, and collateral status. Intracranial atherosclerosis and extracranial atherosclerosis, at the carotid bifurcation, was scored as absent or present on CTA. In addition, the proportion of patients with a cervical ICA stenosis of $\geq 70\%$ and $\geq 99\%$ was calculated using the North American Symptomatic Carotid Endarterectomy Trial (NASCET) criteria.²⁴ These cut-offs were chosen based on their relevance when performing EVT. Vessel tortuosity was assessed for supra-aortic arteries, cervical internal carotid arteries, and aortic arch in two ways: the proportion of patients with one or more angles larger than 90° was assessed for the supra-aortic and cervical ICA and the proportion of patients with severe elongation of the aortic arch (aortic arch type III, indicating that at least one of the supra aortic arteries arises below the level of the inner curvature of the aorta) was calculated.²⁵ Vessel size was measured by calculating the radius of the ICA and M1. Finally, collateral status was divided into absent collaterals (0% filling of the vessel downstream of the occlusion as compared to the contralateral hemisphere), poor collaterals ($\leq 50\%$ filling), moderate collaterals (51%–99% filling), and excellent collaterals (100% filling).²⁶

Tissue characteristics were measured using the Alberta Stroke Program Early Computed Tomography Score (ASPECTS) on non-contrast CT. The ASPECTS is a grading scale from 0 to 10 for which the vascular territory of the middle cerebral artery is divided in 10 regions. Points are subtracted in the regions with any evidence of early ischemic tissue changes.^{27,28} We calculated the median ASPECTS, different categories of ASPECTS, and the proportion of patients with an ASPECTS < 8, which is related to an increased probability of death or dependency.²⁷

Clot and tissue characteristics, presence of intracranial and extracranial atherosclerosis, and collateral status were evaluated by an independent core lab of the MR CLEAN

Registry, blinded to clinical findings.²¹ ICA stenosis and vessel tortuosity were evaluated by trained students under supervision of an experienced neuroradiologist.²⁵ The inter-rater agreement of the assessment of several of the characteristics has been previously described.^{21,25} To calculate the ICA and M1 radius the intracranial vessels were segmented from the CTA data using custom Convolutional Neural Network (CNN) software (StrokeViewer v2.1.22, NICO. LAB, Amsterdam, the Netherlands <http://www.strokeviewer.nico-lab.com>). The semi-automatic software iCafe (©2016–2020, The University of Washington, used with permission) was here used to extract the center line graph, to anatomically label arterial segments, and to determine the local vessel radius at each point.²⁹

Radiological and functional outcome

Radiological outcome was the extended thrombolysis in cerebral infarction (eTICI) score post-EVT, rated on the final DSA run. The eTICI is a grading scale from 0 to 3, and measures the extent of reperfusion in the downstream territory of the occluded vessel. Grade 0 indicates no reperfusion, with grade 1 there is a reduction in thrombus but with no reperfusion, grades 2A, B and C indicate respectively 1%–49%, 50%–89%, and 90%–99% reperfusion, and grade 3 indicates complete reperfusion.^{30,31} Successful reperfusion is defined as a score of 2B or higher, and excellent reperfusion as a score of 2C or higher. To reach these scores, DSA runs including both anteroposterior and lateral views needed to be completed after EVT. If only an anteroposterior view was present, the maximum score was 2A.

Functional outcome was the modified Rankin Scale (mRS) score at 90 days.³² The mRS is a grading scale ranging from 0 to 6 in which 0 indicates no symptoms and six indicates dead. Functional independence is defined as a score of 2 or lower. Furthermore we assessed early neurological outcome with the NIHSS at 24–48 h, and the occurrence of symptomatic intracranial hemorrhages, defined as a hemorrhage with a related decline of four points or more on the NIHSS, according to the Heidelberg criteria.³³

Missing data

If the baseline NIHSS score was missing, it was retrospectively scored with a standardized chart based on the neurological examination in the electronic patient record.³⁴ In case successful reperfusion was not achieved, time of the last contrast bolus was used as a proxy for the time of reperfusion. If the mRS was assessed within 30 days of symptom onset instead of at 90 days and the score was 0–5, it was considered invalid. These values were regarded as missing and replaced by mRS scores resulting from multiple imputations. Regression analyses were performed with imputed data except for part of the vessel characteristics (extracranial atherosclerosis including stenosis, vessel

tortuosity, and vessel size). Because the measurements of these characteristics were only available in a small selection of patients, regression analyses in this subset were executed without multiple imputations.

Statistical analysis

We performed descriptive statistics for baseline characteristics. Neuroimaging characteristics were compared between men and women using *t*-tests for continuous variables and chi-square tests for dichotomous variables. Multivariable regression analyses were performed to assess the association between sex and neuroimaging characteristics: binary logistic regression analyses to assess the association between sex and clot location, atherosclerosis and vessel tortuosity; linear regression analyses to assess the association between sex and CBS, vessel radius and ASPECTS; and ordinal logistic regression analyses to assess the association between sex and collateral status. No adjustments were made in the analyses of clot location and vessel radius, since we assumed it to be unlikely that these characteristics would be significantly influenced by confounders. Analyses of CBS were adjusted for age, onset-to-groin time and collateral status; analyses of vessel tortuosity were adjusted for age and hypertension; analyses of atherosclerosis were adjusted for age, smoking, hypertension, hypercholesterolemia and diabetes mellitus; and analyses of collateral status and ASPECTS were adjusted for age and onset-to-groin time. The association between sex and eTICI post-EVT, mRS, and NIHSS at 24–48 h was assessed using ordinal logistic regression (shift) analyses. Binary logistic regression analyses were performed to assess the association between sex and successful and excellent reperfusion, functional independence, mortality, and symptomatic intracranial hemorrhages. Analyses of eTICI post-EVT were adjusted for age, eTICI pre-EVT, CBS, and additional neuroimaging characteristics that appeared to be significantly different between men and women; analyses of mRS and NIHSS were adjusted for age, diabetes mellitus, pre-stroke mRS, NIHSS on admission (and for mRS NIHSS at 24–48 h), onset to groin time, eTICI post-EVT, and additional neuroimaging characteristics that appeared to be significantly different; and analyses of symptomatic intracranial hemorrhages were adjusted for age, hypertension, NIHSS on admission and NIHSS at 24–48 h. Unadjusted and adjusted (pooled) effect estimates were expressed as mean differences or (common) odds ratios (OR) with 95% confidence intervals. For all statistical analyses SPSS for Windows version 25.0.0 was used.

Results

Of the 3637 consecutive EVT patients, 3180 were included in our analysis, 1526 (48%) of whom were women (Figure 1). Women were 5 years older than men (median age 75 vs

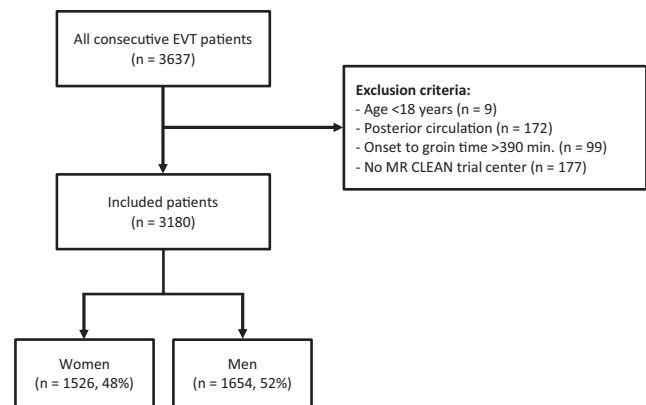


Figure 1. Flow chart of patient selection.

Extent of reperfusion: acOR 0.94; 95% CI 0.83–1.07*.

Successful reperfusion (eTICI ≥ 2B): aOR 0.94; 95% CI 0.81–1.10*.

Excellent reperfusion (eTICI ≥ 2C): aOR 0.90; 95% CI 0.78–1.05*.

*Adjusted for: age, eTICI pre-EVT, CBS, intracranial atherosclerosis, collateral score.

70 years, Table 1), more frequently had a history of hypertension and less often a history of smoking or myocardial infarction. Women were more often functionally dependent prior to the stroke (pre-stroke mRS ≤ 2 in 1259/1526 (85%) vs 1487/1654 (92%)). There was no difference in NIHSS score on admission (median 16 in both groups). The median time from stroke onset to hospital admission (57 vs 50 min) and to groin puncture (200 vs 189 min) was longer in women, and procedural time was somewhat shorter (60 vs 62 min). The overall time from stroke onset to reperfusion or last contrast bolus was longer in women (255 vs 246 min).

Neuroimaging characteristics

There were no statistically significant differences in clot location, although in women the clot tended to be less often located in the ICA (360/1458 (25%) vs 435/1566 (28%), OR 0.85; 95% CI 0.73–1.00) and more often in the M1 segment (874/1458 (60%) vs 890/1566 (57%), OR 1.14; 95% CI 0.98–1.31) than in men (Tables 2–4). No difference was seen in CBS (median 6; IQR 4–8 in both groups).

The prevalence of intracranial atherosclerosis was equal in women and men (877/1462 vs 959/1588, both 60%). However, after adjustment for age, smoking, hypertension, hypercholesterolemia and diabetes mellitus, intracranial atherosclerosis was less frequent in women (adjusted OR (aOR) 0.73; 95% CI 0.62–0.87). Women also less often had extracranial atherosclerosis (275/398 (69%) vs 306/421 (73%), aOR 0.64; 95% CI 0.43–0.95), an ICA stenosis ≥ 70% (90/679 (13%) vs 203/471 (27%), aOR 0.34; 95% CI 0.25–0.47), or ≥ 99% (70/679 (10%) vs 183/741 (25%), aOR 0.27; 95% CI 0.19–0.39). Women more often had one or more angles ≥ 90° in the cervical ICA (226/366 (62%) vs 155/344 (45%), aOR 1.89; 95% CI 1.39–2.57), and more often an aortic arch type III (129/537 (24%) vs 88/570

Table 1. Baseline characteristics of the patients according to sex.

Baseline characteristics	All (n = 3180)	Women (n = 1526)	Men (n = 1654)
Median age (IQR) – yr	72 (61–81)	75 (63–84)	70 (60–77)
Cardiovascular risk factors - no. (%)			
Hypertension	1633 (52)	851 (57)	782 (48)
Hypercholesterolemia	938 (31)	435 (30)	503 (32)
Diabetes mellitus	510 (16)	256 (17)	254 (16)
Atrial fibrillation	756 (24)	383 (26)	373 (23)
Myocardial infarction	441 (14)	161 (11)	280 (17)
Peripheral artery disease	292 (9)	121 (8)	171 (11)
Previous stroke	531 (17)	264 (18)	267 (16)
Current smoking	677 (28)	294 (25)	383 (31)
Medication use on admission - no. (%)			
Antiplatelet	982 (31)	439 (29)	543 (33)
Coumarin	409 (13)	200 (13)	209 (13)
DOAC	104 (3)	47 (3)	57 (4)
Heparin	98 (3)	50 (3)	48 (3)
Statin	1099 (35)	501 (34)	598 (37)
Pre-stroke mRS - no. (%)			
Pre-stroke mRS 0	2107 (68)	926 (62)	1181 (73)
Pre-stroke mRS 1 and 2	639 (21)	333 (22)	306 (19)
Pre-stroke mRS 3	206 (7)	138 (9)	68 (4)
Mean systolic blood pressure on admission \pm SD - mmHg	150 \pm 25	151 \pm 26	149 \pm 24
Median NIHSS score on admission (IQR)	16 (11–19)	16 (11–19)	16 (11–20)
Intravenous thrombolysis treatment - no. (%)	2427 (77)	1145 (75)	1282 (78)
Median time (IQR) - mins.			
From stroke onset to hospital admission	53 (36–90)	57 (37–95)	50 (35–80)
From hospital admission to intravenous thrombolysis	24 (18–33)	25 (19–35)	23 (18–30)
From stroke onset to groin puncture	195 (150–250)	200 (155–260)	189 (145–240)
From start to end of procedure*	60 (40–85)	60 (40–85)	62 (41–89)
From onset of stroke to recanalization/last contrast bolus	250 (199–311)	255 (202–317)	246 (195–305)

Baseline characteristics, with the exception of the median time from start to end of procedure, were described in all 3180 patients.

*Patients were excluded in whom EVT was not performed due to recovery of symptoms before start of the procedure or reperfusion of the occluded vessel, established during digital subtraction angiography.

IQR: interquartile range; DOAC: direct oral anticoagulant; mRS: modified Rankin Scale; SD: standard deviation; NIHSS: National Institutes of Health Stroke Scale.

Table 2. Radiological parameters compared between women and men.

Neuroimaging characteristics	Women	Men	p Value
Clot characteristics			
Clot location			
Intracranial ICA / ICA-T – no. (%)	360/1458 (25)	435/1566 (28)	0.05
M1 – no. (%)	874/1458 (60)	890/1566 (57)	0.08
M2 – no. (%)	214/1458 (15)	227/1566 (15)	0.89
Other: M3/anterior – no. (%)	10/1458 (1)	14/1566 (1)	0.52
Clot extent			
Median clot burden score (IQR)	6 (4–8)	6 (4–8)	0.71
Clot burden score < 7 – no. (%)	721/1187 (61)	729/1247 (59)	0.25
Vessel characteristics			
Vessel aspects			
Presence of intracranial atherosclerosis – no. (%)	877/1462 (60)	959/1588 (60)	0.82
Presence of extracranial atherosclerosis (ICA) – no. (%)	275/398 (69)	306/421 (73)	0.26
ICA stenosis \geq 70% – no. (%)	90/679 (13)	203/741 (27)	<0.001
ICA stenosis \geq 99% – no. (%)	70/679 (10)	183/741 (25)	<0.001

(Continued)

Table 2. (Continued)

Neuroimaging characteristics	Women	Men	p Value
Anatomical aspects			
≥ 1 angles > 90° in supra-aortic arteries – no. (%)	175/410 (43)	163/453 (36)	0.04
≥ 1 angles > 90° in cervical internal carotid arteries – no. (%)	226/366 (62)	155/344 (45)	<0.001
Aortic arch type III – no. (%)	129/537 (24)	88/570 (15)	<0.001
Median radius ICA – mm. (IQR) [#]	2.3 (2.1–2.6)	2.5 (2.3–2.8)	0.002
Median radius M1 – mm. (IQR) [#]	1.6 (1.4–1.8)	1.7 (1.5–1.9)	0.11
Collateral status			
Absent collaterals – no. (%)	69/1434 (5)	117/1544 (8)	0.002
Filling < 50% of occluded area – no. (%)	470/1434 (33)	602/1544 (39)	<0.001
Filling > 50% but less than 100% – no. (%)	593/1434 (41)	561/1544 (36)	0.005
Filling 100% of occluded area – no. (%)	302/1434 (21)	264/1544 (17)	0.006
Tissue characteristics			
Median ASPECTS (IQR),*	9 (8–10)	9 (7–10)	0.05
ASPECTS 0–4 – no. (%),*	65/1366 (5)	72/1499 (5)	0.98
ASPECTS 5–7 – no. (%),*	266/1366 (20)	321/1499 (21)	0.20
ASPECTS 8–10 – no. (%),*	1035/1366 (76)	1106/1499 (74)	0.22
ASPECTS < 8 – no. (%),*	331/1366 (24)	393/1499 (26)	0.22

ICA: internal carotid artery; ICA-T: internal carotid artery terminal; M1/2/3: middle cerebral artery segment 1/2/3; IQR: interquartile range; ASPECTS: Alberta stroke program early CT score.

[#]ICA and M1 radius available in 1223/180 patients.

*Cases with an anterior cerebral artery segment 1 or 2 occlusion excluded.

Table 3. Unadjusted and adjusted effect estimates of sex on neuroimaging characteristics (imputed data).

Neuroimaging characteristics	Effect variable	Unadjusted value (95% CI)	Adjusted value (95% CI)
Clot location			
Intracranial ICA / ICA-T	OR (95% CI)	0.85 (0.73–1.00)	X ^a
M1	OR (95% CI)	1.14 (0.98–1.31)	X ^a
M2	OR (95% CI)	1.02 (0.83–1.24)	X ^a
Other: M3 / anterior	OR (95% CI)	0.77 (0.34–1.73)	X ^a
Clot burden score	β (95% CI)	−0.05 (−0.25 to 0.15)	0.10 (−0.09 to 0.30) ^b
Presence of intracranial atherosclerosis	OR (95% CI)	0.98 (0.84–1.13)	0.73 (0.62–0.87) ^c
Collateral status	cOR (95% CI)	1.40 (1.23–1.60)	1.48 (1.29–1.70) ^d
ASPECTS,*	β (95% CI)	−0.14 (−0.31 to 0.03)	−0.14 (−0.31 to 0.02) ^d

ICA: internal carotid artery; ICA-T: internal carotid artery terminal; M1/2/3: middle cerebral artery segment 1/2/3; ASPECTS: Alberta stroke program early CT score; OR: odds ratio; CI: confidence interval; β: mean difference; cOR: common odds ratio.

^aAdjustments not desirable methodologically.

^bAdjusted for age, time to groin, and collateral status.

^cAdjusted for age, smoking, hypertension, hypercholesterolemia and diabetes mellitus.

^dAdjusted for age and time to groin.

*Cases with an anterior cerebral artery segment 1 or 2 occlusion excluded.

Table 4. Unadjusted and adjusted effect estimates of sex on neuroimaging characteristics (unimputed data).

Neuroimaging characteristics	Effect variable	Unadjusted value (95% CI)	Adjusted value (95% CI)
Presence of extracranial atherosclerosis (ICA)	OR (95% CI)	0.84 (0.60–1.18)	0.64 (0.43–0.95) ^a
ICA stenosis ≥70%	OR (95% CI)	0.33 (0.24–0.46)	0.34 (0.25–0.47) ^a
ICA stenosis ≥99%	OR (95% CI)	0.27 (0.19–0.38)	0.27 (0.19–0.39) ^a
≥ 1 angles >90° in supra-aortic arteries	OR (95% CI)	1.30 (0.98–1.71)	1.11 (0.82–1.49) ^b
≥ 1 angles >90° in cervical internal carotid arteries	OR (95% CI)	1.98 (1.46–2.67)	1.89 (1.39–2.57) ^b
Aortic arch type III	OR (95% CI)	1.75 (1.29–2.38)	1.38 (1.00–1.91) ^b
Radius ICA	β (95% CI)	0.22 (0.09–0.35)	X ^c
Radius M1	β (95% CI)	0.09 (−0.02–0.21)	X ^c

ICA: internal carotid artery; OR: odds ratio; CI: confidence interval; β: mean difference.

^aAdjusted for age, smoking, hypertension, hypercholesterolemia and diabetes mellitus.

^bAdjusted for age and hypertension.

^cAdjustments not desirable methodologically.



Figure 2. (a) Distribution of eTICI after EVT.

Extent of reperfusion: aOR 0.94; 95%CI 0.83 to 1.07*.

Successful reperfusion (eTICI \geq 2B): aOR 0.94; 95%CI 0.81 to 1.10*.

Excellent reperfusion (eTICI \geq 2C): aOR 0.90; 95%CI 0.78 to 1.05*.

*Adjusted for: age, eTICI pre-EVT, CBS, intracranial atherosclerosis, collateral score.

(b) distribution of mRS at 90 days.

Distribution of mRS scores: aOR 0.81; 95%CI 0.69 to 0.94*.

Functional independence (mRS \leq 2): aOR 0.68; 95%CI 0.53 to 0.86*.

Mortality (mRS 6): aOR 0.76; 95%CI 0.59 to 0.99*.

*Adjusted for: age, diabetes mellitus, pre-stroke mRS, baseline NIHSS, follow-up NIHSS, onset to groin time, eTICI post-EVT, intracranial atherosclerosis, collateral score.

(15%), aOR 1.38; 95% CI 1.00–1.91). Vessel tortuosity in the supra-aortic arteries was essentially equal (175/410 (43%) vs 163/453 (36%), aOR 1.11; 95% CI 0.82–1.49). The ICA and M1 radius, data available in 122/3180 patients, were smaller in women, but differences were only minimal (median radius ICA: 2.3 mm (IQR 2.1–2.6 mm) vs 2.5 mm (IQR 2.3–2.8 mm), mean difference 0.22; 95% CI 0.09–0.35, median radius M1: 1.6 mm (IQR 1.4–1.8 mm) vs 1.7 mm (IQR 1.5–1.9 mm), mean difference 0.09; 95% CI –0.02 to 0.21). Absence of collaterals or less than 50% filling was less often seen in women (539/1434 (38%) vs 719/1544 (47%) and, overall, women had better collateral status (adjusted common OR (aOR) 1.46; 95% CI 1.27–1.67).

Early tissue changes were essentially similar with a median ASPECTS of nine in both groups and an ASPECTS $<$ 8 in 331/1366 (24%) versus 393/1499 (26%), adjusted mean difference –0.13; 95% CI –0.30 to 0.02.

Radiological and functional outcome

The difference in proportions between the various eTICI and mRS scores are presented in Figure 2(a) and (b). The

extent of reperfusion (eTICI) was similar in women and men (aOR 0.94; 95% CI 0.83–1.07). Successful reperfusion (eTICI \geq 2B) was reached equally often in both groups (61% vs 63%, aOR 0.94; 95% CI 0.81–1.10) as was excellent reperfusion (eTICI \geq 2C) (38% vs 40%, aOR 0.90; 95% CI 0.78–1.05). There was a substantial shift toward a worse functional outcome in women versus men in the distribution of mRS scores at 90 days (aOR 0.81; 95% CI 0.69–0.94) and women less often reached functional independence (34% vs 46%, aOR 0.68; 95% CI 0.53–0.86). Mortality was slightly higher in women compared with men (32% vs 27%, aOR 0.76; 95% CI 0.53–0.99). Median NIHSS after 24–48 h (early neurological outcome) was 11 (IQR 4–18) in women versus 9 (IQR 4–16) in men, aOR 1.08; 95% CI 0.95–1.24. Symptomatic intracranial hemorrhage rates were similar (7% vs 5% (aOR 1.22; 95% CI 1.00–1.48)).

Discussion

In this large, real-world, cohort of patients with acute anterior LVO stroke, performed in the Netherlands, we found sex differences in vessel characteristics. Women less often had intra- and extracranial atherosclerosis and had a better

collateral status, whereas vessel tortuosity was less often seen in men. Clot and tissue characteristics were the same or only minimally different in men and women with LVO. Despite sex differences in vessel characteristics, eTICI scores were similar in both sexes. Nevertheless, at 90 days, women had a worse functional outcome.

Women in our study tended to less often have an occlusion in the intracranial ICA and more often an occlusion of the M1 segment than men. This difference was also found in previous smaller studies, but never reached statistical significance (Table 5). Even in our large sample, statistical significance was not reached, implicating that if sex differences in clot location are present they are probably only minimal and will not have a substantial effect on treatment success or outcome. The finding that women had less intracranial atherosclerosis in our study compared to men contradicts with a Chinese study of 2864 patients with an acute ischemic stroke that found a higher frequency of intracranial atherosclerosis in women.¹⁵ A recent analysis of the DUST study group showed comparable intracranial atherosclerosis presence in a general ischemic stroke population of 1397 Dutch men and women.³⁵ Differences between the current study and the previous studies might be explained by differences in ethnic background or in stroke population. We found that women more often had tortuous vessels than men. Both the presence of atherosclerosis and vessel tortuosity may increase the technical difficulty of EVT and therefore prolong procedural duration, which may negatively affect functional outcome.³⁶ In our study, procedural times were equal between sexes. However, the number of patients in whom vessel characteristics were complete was relatively small so we refrained from performing a subanalysis investigating a direct relationship between vessel characteristics and procedural time. Finally, in line with a recent meta-analysis and a sub-analysis of the DEFUSE three trial women had a better collateral status.^{37,38} This better collateral circulation in women could lead to better protection against cerebral ischemia. In addition, early ischemic tissue changes as measured with the ASPECT score were comparable between sexes, but unfortunately no data were available on final infarct volumes. The similar ASPECTS in our study is in accordance with previous studies as presented in Table 5.

In our study, women less often reached functional independence after 90 days. In contrast, in most RCTs, functional outcomes were similar in men and women treated with EVT.^{8,9,39} A subgroup analysis of 1762 patients from seven RCTs found that functional independence was reached in 39% of men and women 90 days after EVT.³⁷ The difference with our study might be selection of a healthier and younger study population in the RCTs. In our study, both men and women were 5–7 years older than the RCT populations. The worse outcome in women after EVT in our study is also in contrast with two previous small observational studies, that again included a younger study

population compared with our study, but in line with another small observational study that found relatively poor functional outcome after 90 days in women (Table 5).^{9,39,40}

Our study has limitations. Because our analyses were exploratory, we did not correct for multiple testing. Results should thus be interpreted with caution due to the potential risk of type I errors. Although we corrected for possible confounders we cannot exclude that residual confounding occurred. Also, because the MR CLEAN Registry was not a randomized trial and did not include a non-treated control group, we were not able to investigate sex differences in treatment effect.^{8,37} Furthermore, because the MR CLEAN Registry only included patients eligible for EVT, we cannot exclude a selection bias in case there may be sex differences in neuroimaging characteristics leading to abstaining from EVT. Strengths of our study are the prospective study design and the large number of patients studied in a real-world setting.

Why women in our study, a real-world, non-trial setting, do worse than men after EVT remains unclear. In accordance with earlier findings, the time from stroke onset to hospital admission and in-hospital delays were slightly longer in women than in men. This might be because older women more often live alone than men, leading to larger prehospital delays.¹⁴ However, EVT treatment times were slightly shorter in women and the overall difference in onset to recanalization times was less than 10 min. Differences in outcome can also develop as a result of other in-hospital treatments, such as IVT or stroke-unit care. Both better and worse functional outcomes have been reported in women receiving IVT.^{43,44} In our cohort there were slight differences in the proportion of women receiving IVT compared with men (75% vs 78%) and time from hospital admission to intravenous thrombolysis was on average 2 min longer in women, but it is unlikely that these small differences account for the worse outcome in women. Moreover, a large prospective registry in Austria did not show a difference in quality of stroke unit care between women and men.⁴³ Another likely explanation is higher frailty in women. Women have a higher life expectancy and more often suffer from stroke at a later age compared with men.^{13,44,45} Consistent with previous studies, women in our study were more often functionally dependent before their stroke.⁴⁶ Although we adjusted for age and pre-stroke mRS, we cannot exclude that women had more additional co-morbidity. This frailty and co-morbidity probably increases the risk of complications or impaired recovery after discharge from the hospital. These social determinants of health might become clearer in a real-world versus a controlled research setting. This hypothesis is in line with two previous studies that found similar outcomes at discharge but impaired functional outcome at 90 days in women compared with men and supported by the similar NIHSS in men and women at 24 h in our study.^{9,11} Data on outcome at discharge were

Table 5. Overview of studies comparing sex differences in neuro-imaging characteristics in EVT treated patients.

Study characteristics		Radiological characteristics (women vs men)				Radiological outcome		Functional outcome	
Study	Patients	Women	Occlusion location	Collaterals	Mean diameter (mm)	ASPECTS	mTICI	mRS	mRS 0–2
Madsen et al. ⁹	279 All EVT	145 (52%)	ICA: 14%vs 18%# MI: 63%vs 61%#	X	X	X	X	X	aOR 0.37; 95% CI 0.16–0.87#
Fifi et al. ¹¹	285 All EVT	139 (49%)	ICA: 19%vs 21%# MI: 63%vs 56%#	>50% filling: 49%vs 48%#	X	8vs 8#	X	median score: 3vs 1 (p=0.04) shift: aOR 0.49; 95% CI 0.25–0.96# median score: 3vs 3 (p=0.24)	46%vs 60% (p=0.05)# aOR 0.41; 95% CI 0.16–1.00
Davison et al. ¹⁷	92 All EVT	42 (46%)	ICA: 24%vs 26%# MI: 50%vs 44%#	X	ICA: 2.81vs 3.08 (p=0.01)# MCA: 2.18vs 2.47 (p<0.0001)#		mTICI 2B or 3: 88%vs 84% (p=0.57)#		29%vs 44% (p=0.13)#
Demeestere et al. ¹⁸	198 All EVT	91 (46%)	X	X	X	7vs 7#	mTICI 2B or 3: 84%vs 87% (p=0.30)#	shift: acOR 1.79; 95% CI 1.04–3.08#	aOR 1.40; 95% CI 0.69–2.86#
Chalos et al. ³⁷	1762 EVT vs. control	833 (47%)	ICA: 26%vs 27%~ MI: 65%vs 65%~	>50% filling: 88%vs 80%~	X	8vs 8~	mTICI 2B or 3: 75%vs 76%~	shift: aOR 2.13; 95% CI 1.47–3.07* shift: aOR 2.16; 95% CI 1.59–2.96^ score: 3.5vs 2 (p=0.046)#	aOR 2.62; 95% CI 1.89–3.64* aOR 2.22; 95% CI 1.53–3.22^ X
Dula et al. ³⁸	182 EVT vs. control	92 (51%)	ICA: 32%vs 42%~ MI: 66%vs 58%~	>50% filling: 75%vs 67% (p=0.290)~	X	8vs 8~	X		
Carvalho et al. ³⁹	145 All EVT	81 (56%)	ICA: 30%vs 28%# MI: 56%vs 52%#	X	X	8vs 8#	mTICI 2B or 3: 85%vs 92% (p=0.19)#	×aOR 0.70; 95% CI 0.33–1.50#	
Sheth et al. ⁴⁰	389 All EVT	214 (55%)	ICA: 18%vs 20%# MI: 69%vs 64%#	X	X	8vs 8#	mTICI 2B or 3: 87%vs 83% (p=0.37)#	X	aOR 1.01; 95% CI 0.64–1.59#
Regenhardt et al. ⁴⁸	381 All EVT	193 (51%)	ICA: 18%vs 19%# MI: 67%vs 71%	X	X	X	mTICI 2B or 3: 78%vs 78% (p=1.00)#	X	44%vs 39% (p=0.377)#
Deb-Chatterji et al. ⁴⁹	2316 All EVT	1170 (51%)	X	X	X	9vs 9#	mTICI 2B or 3: 83%vs 83% (p=0.92)#	score: 4vs 3 (p=0.001)#	33%vs 41% (p<0.001)#
Casetta et al. ⁵⁰	3422 All EVT	1801 (53%)	ICA: 31%vs 40%# MI: 69%vs 61%#	X	X	X	mTICI 2B or 3: aOR 1.18 (1.03–1.38)#	shift: aOR 0.94 (0.82–1.06)#	aOR 1.19 (1.02–1.38)#
Sun et al. ⁵¹	3121 All EVT	1321 (37%)	ICA: 30%vs 34%# MI: 56%vs 56%	X	X	9vs 9#		median score: 3vs 3 (p=0.412)# shift: aOR 0.90 (0.70–1.16)#	39%vs 49% (p=0.970)# aOR 0.99 (0.73–1.35)#

EVT: endovascular treatment; ICA: internal carotid artery; M1/2/3: middle cerebral artery segment 1; ASPECTS: Alberta Stroke Program Early CT Score; MCA: middle cerebral artery; mRS: modified Rankin Scale; aOR: adjusted odds ratio; acOR: adjusted common odds ratio; mTICI: modified thrombolysis in cerebral infarction.
 ~EVT and control women versus men. #EVT women versus men. *Women EVT versus control. ^Men EVT versus control.

unfortunately not available. Impaired recovery after discharge from the hospital could be caused by differences between men and women during inpatient rehabilitation. However, a recent Canadian study comparing 9459 women and 10,684 men admitted to inpatient rehabilitation after stroke did not show a difference in length of stay, and women were more likely to be discharged home compared to men.⁴⁹ In conclusion, substantial sex differences in vessel characteristics exist in a large population of LVO patients in a real-world setting in the Netherlands. However, these sex differences do not affect radiological outcome and are therefore unlikely to explain the worse functional outcome in women. Our data, therefore, provide no argument that, based on neuroimaging characteristics, decisions on treatment with EVT should be taken differently in men and women with LVO.

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Ethical approval

The MR CLEAN Registry was approved by the medical ethics committee of the Erasmus University Medical Center, Rotterdam, the Netherlands (MEC-2014-235).

Informed consent

Informed consent was not sought for the present study. As this is a registry study the medical ethics committee waived the need for obtaining informed consent.






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AM, MW and MJHW researched literature and conceived the study. GH, HO and PK gained data for the study. HO and HL were involved in data analysis. AM wrote the first draft of the manuscript. All authors reviewed and edited the manuscript and approved the final version of the manuscript.

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References

1. Reeves MJ, Bushnell CD, Howard G, et al. Sex differences in stroke: epidemiology, clinical presentation, medical care, and outcomes. *Lancet Neurol* 2008; 7: 915–926.
2. Gall SL, Tran PL, Martin K, et al. Sex differences in long-term outcomes after stroke: functional outcomes, handicap, and quality of life. *Stroke* 2012; 43: 1982–1987.
3. Phan HT, Blizzard CL, Reeves MJ, et al. Factors contributing to sex differences in functional outcomes and participation after stroke. *Neurology* 2018; 90: e1945–e1953.
4. Bushnell C, Howard VJ, Lisabeth L, et al. Sex differences in the evaluation and treatment of acute ischaemic stroke. *Lancet Neurol* 2018; 17: 641–650.
5. Silva GS, Lima FO, Camargo EC, et al. Gender differences in outcomes after ischemic stroke: role of ischemic lesion volume and intracranial large-artery occlusion. *Cerebrovasc Dis* 2010; 30: 470–475.
6. Gargano JW and Reeves MJ. Sex differences in stroke recovery and stroke-specific quality of life: results from a state-wide stroke registry. *Stroke* 2007; 38: 2541–2548.
7. Goyal M, Menon BK, van Zwam WH, et al. Endovascular thrombectomy after large-vessel ischaemic stroke: a meta-analysis of individual patient data from five randomised trials. *Lancet* 2016; 387: 1723–1731.
8. de Ridder IR, Fransen PS, Beumer D, et al. Is intra-arterial treatment for acute ischemic stroke less effective in women than in men? *Interv Neurol* 2016; 5: 174–178.
9. Madsen TE, DeCrose-Movson E, Hemendinger M, et al. Sex differences in 90-day outcomes after mechanical thrombectomy for acute ischemic stroke. *J Neurointerv Surg* 2019; 11: 221–225.
10. Uchida K, Yoshimura S, Sakai N, et al. Sex differences in management and outcomes of acute ischemic stroke with large vessel occlusion. *Stroke* 2019; 50: 1915–1918.
11. Fifi JT, Nguyen TN, Song S, et al. Sex differences in endovascular thrombectomy outcomes in large vessel occlusion: a propensity-matched analysis from the SELECT study. *J Neurointerv Surg* 2023; 15: 105–112.
12. Dmytriw AA, Ku JC, Yang VXD, et al. Do outcomes between women and men differ after endovascular thrombectomy? A meta-analysis. *AJNR Am J Neuroradiol* 2021; 42: 910–915.

13. Appellos P, Stegmayr B and Terént A. Sex differences in stroke epidemiology: a systematic review. *Stroke* 2009; 40: 1082–1090.
14. Reeves MJ and Lisabeth LD. The confounding issue of sex and stroke. *Neurology* 2010; 74: 947–948.
15. Pu Y, Liu L, Wang Y, et al. Geographic and sex difference in the distribution of intracranial atherosclerosis in China. *Stroke* 2013; 44: 2109–2114.
16. Faber JE, Moore SM, Lucitti JL, et al. Sex differences in the cerebral collateral circulation. *Transl Stroke Res* 2017; 8: 273–283.
17. Davison MA, Ouyang B, Keppetipola KM, et al. Arterial diameter and the gender disparity in stroke thrombectomy outcomes. *J Neurointerv Surg* 2018; 10: 949–952.
18. Demeestere J, Christensen S, Mlynash M, et al. Effect of sex on clinical outcome and imaging after endovascular treatment of large-vessel ischemic stroke. *J Stroke Cerebrovasc Dis* 2021; 30(2): 1–6.
19. Kaymaz ZO, Nikoubashman O, Brockmann MA, et al. Influence of carotid tortuosity on internal carotid artery access time in the treatment of acute ischemic stroke. *Interv Neuroradiol* 2017; 23: 583–588.
20. Snelling BM, Sur S, Shah SS, et al. Unfavorable vascular anatomy is associated with increased revascularization time and worse outcome in anterior circulation thrombectomy. *World Neurosurg* 2018; 120: e976–e983.
21. Jansen IGH, Mulder MJHL and Goldhoorn RB. Endovascular treatment for acute ischaemic stroke in routine clinical practice: prospective, observational cohort study (MR CLEAN Registry). *BMJ* 2018; 360: k949.
22. Puetz V, Dzialowski I, Hill MD, et al. Intracranial thrombus extent predicts clinical outcome, final infarct size and hemorrhagic transformation in ischemic stroke: the clot burden score. *Int J Stroke* 2008; 3: 230–236.
23. Treurniet KM, Yoo AJ, Berkhemer OA, et al. Clot burden score on baseline computerized tomographic angiography and intra-arterial treatment effect in acute ischemic stroke. *Stroke* 2016; 47: 2972–2978.
24. Barnett HJM, Taylor DW, Haynes RB, et al. Beneficial effect of carotid endarterectomy in symptomatic patients with high-grade carotid stenosis. *New Engl J Med* 1991; 325: 445–453.
25. Holswilder G, Stuart MP, Dompeling T, et al. The prognostic value of extracranial vascular characteristics on procedural duration and revascularization success in endovascularly treated acute ischemic stroke patients. *Eur Stroke J* 2022; 7: 48–56.
26. Tan IY, Demchuk AM, Hopyan J, et al. CT angiography clot burden score and collateral score: correlation with clinical and radiologic outcomes in acute middle cerebral artery infarct. *AJNR Am J Neuroradiol* 2009; 30: 525–531.
27. Barber PA, Demchuk AM, Zhang J, et al. Validity and reliability of a quantitative computed tomography score in predicting outcome of hyperacute stroke before thrombolytic therapy. *Lancet* 2000; 355: 1670–1674.
28. Pexman JH, Barber PA, Hill MD, et al. Use of the Alberta Stroke Program Early CT Score (ASPECTS) for assessing CT scans in patients with acute stroke. *AJNR Am J Neuroradiol* 2001; 22: 1534–1542.
29. Chen L, Mossa-Basha M, Balu N, et al. Development of a quantitative intracranial vascular features extraction tool on 3D MRA using semiautomated open-curve active contour vessel tracing. *Magn Reson Med* 2018; 79: 3229–3238.
30. Higashida RT, Furlan AJ, Roberts H, et al. Trial design and reporting standards for intra-arterial cerebral thrombolysis for acute ischemic stroke. *Stroke* 2003; 34: e109–e137.
31. Dekker L, Geraedts VJ, Hund H, et al. Importance of reperfusion status after intra-arterial thrombectomy for prediction of outcome in anterior circulation large vessel stroke. *Interv Neurol* 2018; 7: 137–147.
32. van Swieten JC, Koudstaal PJ, Visser MC, et al. Interobserver agreement for the assessment of handicap in stroke patients. *Stroke* 1988; 19: 604–607.
33. vonKummer R, Broderick JP, Campbell BC, et al. The Heidelberg bleeding classification: classification of bleeding events after ischemic stroke and reperfusion therapy. *Stroke* 2015; 46: 2981–2986.
34. Williams LS, Yilmaz EY and Lopez-Yunez AM. Retrospective assessment of initial stroke severity with the NIH Stroke Scale. *Stroke* 2000; 31: 858–862.
35. Voigt S, van Os H, van Walderveen M, et al. Sex differences in intracranial and extracranial atherosclerosis in patients with acute ischemic stroke. *Int J Stroke* 2021; 16: 385–391.
36. Leker RR, Kasner SE, El Hasan HA, et al. Impact of carotid tortuosity on outcome after endovascular thrombectomy. *Neurosci Sci* 2021; 42: 2347–2351.
37. Chalos V, de Ridder IR, Lingsma HF, et al. Does sex modify the effect of endovascular treatment for ischemic stroke? *Stroke* 2019; 50: 2413–2419.
38. Dula AN, Mlynash M, Zuck ND, et al. Neuroimaging in ischemic stroke is different between men and women in the DEFUSE 3 cohort. *Stroke* 2020; 51: 481–488.
39. Carvalho A, Cunha A, Gregório T, et al. Is the efficacy of endovascular treatment for acute ischemic stroke sex-related. *Interv Neurol* 2018; 7: 42–47.
40. Sheth SA, Lee S, Warach SJ, et al. Sex differences in outcome after endovascular stroke therapy for acute ischemic stroke. *Stroke* 2019; 50: 2420–2427.
41. Kent DM, Price LL, Ringleb P, et al. Sex-based differences in response to recombinant tissue plasminogen activator in acute ischemic stroke: a pooled analysis of randomized clinical trials. *Stroke* 2005; 36: 62–65.
42. Spaander FH, Zinkstok SM, Baharoglu IM, et al. Sex differences and functional outcome after intravenous thrombolysis. *Stroke* 2017; 48: 699–703.
43. Gatttringer T, Ferrari J, Knoflach M, et al. Sex-related differences of acute stroke unit care: results from the Austrian stroke unit registry. *Stroke* 2014; 45: 1632–1638.
44. Gall SL, Donnan G, Dewey HM, et al. Sex differences in presentation, severity, and management of stroke in a population-based study. *Neurology* 2010; 74: 975–981.
45. Phan HT, Blizzard CL, Reeves MJ, et al. Sex differences in long-term mortality after stroke in the INSTRUCT (INternational STroke oUtcomes sTudy): a meta-analysis of individual participant data. *Circ Cardiovasc Qual Outcomes* 2017; 10(2): 116. DOI: 10.1161/CIRCOUTCOMES.116.003436

46. Di Carlo A, Lamassa M, Baldereschi M, et al. Sex differences in the clinical presentation, resource use, and 3-month outcome of acute stroke in Europe: data from a multicenter multinational hospital-based registry. *Stroke* 2003; 34: 1114–1119.
47. MacDonald SL, Hall RE, Bell CM, et al. Sex differences in the outcomes of adults admitted to inpatient rehabilitation after stroke. *PM R* 2022; 14: 779–785.
48. Regenhardt RW, Turner AC, Hirsch JA, et al. Sex-specific differences in presentations and determinants of outcomes after endovascular thrombectomy for large vessel occlusion stroke. *J Neurol* 2022; 269: 307–315.
49. Deb-Chatterji M, Schlemm E, Flottmann F, et al. Sex differences in outcome after thrombectomy for acute ischemic stroke are explained by confounding factors. *Clin Neuroradiol* 2021; 31: 1101–1109.
50. Casetta I, Fainardi E, Pracucci G, et al. Sex differences in outcome after thrombectomy for acute ischemic stroke. A propensity score-matched study. *Eur Stroke J* 2022; 7: 151–157.
51. Sun D, Raynald R, Huo X, et al. Sex-related differences in outcomes of endovascular treatment for anterior circulation large vessel occlusion. *Stroke* 2023; 54: 327–336.