

Collateral status and recanalization after endovascular treatment for acute ischemic stroke

Venema, S.M.U.; Dankbaar, J.W.; Wolff, L.; Es, A.C.G.M. van; Sprengers, M.; Lugt, A. van der; ... ; MR CLEAN Registry Investigators

Citation

Venema, S. M. U., Dankbaar, J. W., Wolff, L., Es, A. C. G. M. van, Sprengers, M., Lugt, A. van der, ... Worp, H. B. van der. (2022). Collateral status and recanalization after endovascular treatment for acute ischemic stroke. *Journal Of Neurointerventional Surgery*. doi:10.1136/neurintsurg-2021-018545

Version:Publisher's VersionLicense:Licensed under Article 25fa Copyright Act/Law (Amendment Taverne)Downloaded from:https://hdl.handle.net/1887/3512779

Note: To cite this publication please use the final published version (if applicable).

Original research

Collateral status and recanalization after endovascular treatment for acute ischemic stroke

Simone M Uniken Venema ^(b), ¹ Jan Willem Dankbaar, ² Lennard Wolff, ³ Adriaan C G M van Es, ⁴ Marieke Sprengers, ⁵ Aad van der Lugt, ³ Diederik W J Dippel ^(b), ⁶ H. Bart van der Worp, ¹ on behalf of the MR CLEAN Registry investigators

ABSTRACT

► Additional supplemental material is published online only. To view, please visit the journal online (http://dx.doi. org/10.1136/neurintsurg-2021-018545).

¹Department of Neurology and Neurosurgery, Brain Center, University Medical Centre Utrecht, Útrecht, The Netherlands ²Department of Radiology, University Medical Center Utrecht Imaging Division, Utrecht, The Netherlands ³Department of Radiology and Nuclear Medicine, Erasmus Medical Center, Rotterdam, Zuid-Holland, The Netherlands ⁴Department of Radiology, Leiden Universitair Medisch Centrum, Leiden, Zuid-Holland. The Netherlands ⁵Department of Radiology and Nuclear Medicine, Amsterdam UMC Locatie AMC, Amsterdam, North Holland, The Netherlands ⁶Department of Neurology, Erasmus Medical Center, Rotterdam, Zuid-Holland, The Netherlands

Correspondence to

Simone M Uniken Venema, University Medical Centre Utrecht Brain Centre, Utrecht, The Netherlands; s.m. unikenvenema@umcutrecht.nl

Received 14 December 2021 Accepted 2 May 2022

Check for updates

© Author(s) (or their employer(s)) 2022. No commercial re-use. See rights and permissions. Published by BMJ.

To cite: Uniken Venema SM, Dankbaar JW, Wolff L, et al. J NeuroIntervent Surg Epub ahead of print: [please include Day Month Year]. doi:10.1136/ neurintsurg-2021-018545 **Background** Successful recanalization and good collateral status are associated with good clinical outcomes after endovascular treatment (EVT) for acute ischemic stroke, but the relationships among them are unclear.

Objective To assess if collateral status is associated with recanalization after EVT and if collateral status modifies the association between successful recanalization and functional outcome.

Methods We retrospectively analyzed data from the MR CLEAN Registry, a multicenter prospective cohort study of patients with a proximal anterior occlusion who underwent EVT in the Netherlands. We determined collateral status with a previously validated four-point visual grading scale and defined successful recanalization as an extended Thrombolysis in Cerebral Infarction score \geq 2B. Functional outcome was determined using the modified Rankin Scale score at 90 days. We assessed, with multivariable logistic regression models, the associations between (1) collateral status and successful recanalization. (2) successful recanalization and functional outcome. (3) collateral status and functional outcome. An interaction of collateral status and successful recanalization was assessed. Subgroup analyses were performed for patients treated with intravenous thrombolvsis.

Results We included 2717 patients, of whom 1898 (70%) had successful recanalization. There was no relationship between collateral status and successful recanalization (adjusted common OR (95% CI) of grades 1, 2, and 3 vs 0: 1.19 (0.82 to 1.72), 1.20 (0.83 to 1.75), and 1.10 (0.74 to 1.63), respectively). Successful recanalization (acOR (95% CI): 2.15 (1.84 to 2.52)) and better collateral grades (acOR (95% CI) of grades 1, 2, and 3 vs 0: 2.12 (1.47 to 3.05), 3.46 (2.43 to 4.92), and 4.16 (2.89 to 5.99), respectively) were both associated with a shift towards better functional outcome, without an interaction between collateral status and successful recanalization. Results were similar for the subgroup of thrombolysed patients.

Conclusions Collateral status is not associated with the probability of successful recanalization after EVT and does not modify the association between successful recanalization and functional outcome.

INTRODUCTION

Successful recanalization^{1 2} and good collateral status^{3 4} are strongly associated with favorable clinical outcomes after endovascular treatment (EVT) for acute ischemic stroke. Some studies have suggested that good collateral status at baseline is associated with a higher probability of vessel recanalization and tissue reperfusion of the downstream territory after intravenous thrombolysis⁵ or EVT,⁷⁻⁹ including two recent studies with successful recanalization rates between 85% and 90%.^{10 11} Sufficient flow into collateral vessels may enhance the delivery of thrombolytics to the distal end of the clot⁷ and may prevent detached thrombus segments from occluding more distal vessels.¹⁰ Moreover, it has been suggested that collateral status modifies the relationship between successful reperfusion and functional outcome, in such a way that collateral status and successful reperfusion synergistically contribute to a better outcome.^{12–14} A more recent study did however not confirm this.¹¹

It therefore remains uncertain if, and to what extent, collateral status facilitates recanalization in current practice and if collateral status modifies the association between successful recanalization and functional outcome. We addressed these questions by analyzing data from the MR CLEAN Registry, a multicenter prospective cohort study of patients treated with EVT in routine clinical practice in the Netherlands.

METHODS

Study protocol and patients

The Multicenter Randomized Controlled Trial of Endovascular Treatment for Acute Ischemic Stroke (MR CLEAN) Registry is a prospective, multicenter, observational study of patients with acute ischemic stroke treated with EVT in the Netherlands. The present retrospective study is based on data from patients with acute ischemic stroke included in the MR CLEAN Registry between March 16, 2014 and November 1, 2017 who fulfilled the following criteria: (1) age 18 years or older, (2) EVT initiated within 6.5 hours of symptom onset or last seen well, (3) availability of an extended Thrombolysis in Cerebral Infarction (eTICI) score at the end of EVT, as assessed on two-dimensional (anteroposterior and lateral) digital subtraction angiography (DSA) runs, (4) presence of a proximal arterial



1

occlusion in the anterior circulation at baseline (internal carotid artery (ICA, ICA-terminus), middle (M1, M2) cerebral artery), confirmed with CT angiography (CTA). EVT was performed as part of routine clinical care and the exact procedure following groin puncture and catheterization to the level of occlusion (ie, mechanical thrombectomy, thrombus aspiration, or both, with or without the use of a thrombolytic agent) was left at the discretion of the treating interventionalist.

The study protocol of the MR CLEAN Registry was approved by the Erasmus University Medical Center ethics committee and permission was granted to carry out the study as a registry (MEC-2014–235). The requirement for written informed consent was waived, but patients were provided with information on the study and were given the opportunity to refuse participation. Source data from patients cannot be distributed, but syntax and output files from statistical analyses can be made available on reasonable request to the corresponding author.

Assessment of clinical and imaging data

Clinical data were collected from electronic patient records at the recruiting centers by local study investigators. The score on the modified Rankin Scale (mRS), an ordinal disability scale ranging from 0 (no symptoms) to 6 (death), was assessed at 90 days (± 14 days) after stroke through a telephone interview. Neuroimaging data were centrally analyzed by an imaging core laboratory consisting of trained neuroradiologists and interventionalists, blinded to all clinical findings except the symptom side. Collateral status on baseline CTA was determined using a previously validated scoring method.¹⁵ This is a four-point visual grading scale, with 0 for absent collaterals (0% filling of the vascular territory downstream of the occlusion), 1 for poor collaterals (>0% and \leq 50% filling), 2 for moderate collaterals (>50% and <100% filling), and 3 for excellent collaterals (100% filling). Interobserver agreement for this method has previously been determined and was moderate (k=0.60).³ The extent of antegrade recanalization of the previously occluded vascular territory at the end of EVT was assessed on DSA using the eTICI score.¹⁶ A score of 2B or higher, indicating antegrade recanalization of more than half of the previously occluded target artery ischemic territory (eg, in two major divisions of the middle cerebral artery (and their territories), was considered successful recanalization. Time from symptom onset to the end of EVT was defined as the moment at which successful recanalization was achieved or, if not applicable, the moment the last contrast bolus was given.

Outcomes

The primary outcome was successful recanalization (eTICI \geq 2B). Secondary outcomes were complete recanalization (eTICI of 3), post-EVT eTICI score (as an ordinal outcome variable), spontaneous recanalization (eTICI \geq 2B prior to EVT on diagnostic DSA), first pass effect (defined as the achievement of nearcomplete or complete (eTICI 2C–3) recanalization after a single pass of the device, without rescue by intra-arterial thrombolytics) and the score on the mRS at 90 days post-stroke.

Statistical analysis

Baseline characteristics were compared between patients in whom successful recanalization was achieved (eTICI $\ge 2B$) versus those without successful recanalization (eTICI $\le 2A$) We assessed the relationship between collateral status and recanalization outcomes (successful recanalization, complete recanalization, ordinal eTICI score, spontaneous recanalization, and first pass effect) using multivariable binary or ordinal logistic regression adjusted for potential confounders. For ordinal outcome variables, we tested the proportional odds assumption and, if not violated, we used ordinal logistic regression. To assess the possibility of a differential effect of collateral status on recanalization by occlusion location, we added an interaction term (collateral status×occlusion location). If the interaction test was significant, we performed subgroup analyses by occlusion location.

We then assessed the relationship between successful recanalization and functional outcome, as assessed with the mRS score at 90 days, in a multivariable ordinal logistic regression model. We added an interaction term for collateral status (successful recanalization×collateral status). If the model containing the interaction terms performed significantly better than the model without, as assessed with a likelihood ratio test, we then repeated the analysis in subgroups of patients according to recanalization status. All multivariable models were adjusted for age, systolic blood pressure on admission, Alberta Stroke Program Early CT Score at baseline, extracranial carotid artery disease, occluded segment, and time from symptom onset to recanalization.

Subgroup analyses were performed for patients treated with intravenous alteplase prior to EVT. Sensitivity analyses were performed for the subgroup of patients undergoing intra-arterial thrombectomy. In all models, collateral status was analyzed as an ordinal independent variable, with absent collaterals (grade 0) as a reference category and effect estimates-adjusted common ORs (acOR) and 95% confidence intervals (CIs)-were calculated for collateral grades 1, 2, and 3 compared with grade 0. In sensitivity analyses, collateral status was analyzed as a numeric independent variable, and effect estimates were calculated for each one-point increase in collateral grade. All analyses were done using R software (version 1.3.1093). In order to obtain unbiased effect estimates, missing data (assumed to be missing at random) were imputed with multiple imputations (n=5) using the AregImpute function. All multivariable analyses were done using imputed data.

RESULTS

Patient characteristics

Of 3279 patients included in the MR CLEAN Registry, 2717 patients were included in the present study (online supplemental figure 1). Successful recanalization at the end of EVT was achieved in 1898 patients (70%). Those with successful recanalization were younger, had a lower systolic blood pressure on admission, less frequently had a history of atrial fibrillation, and used coumarins or direct oral anticoagulants less frequently (table 1). ICA or ICA-T occlusions were more frequent in those with unsuccessful recanalization, whereas those with successful recanalization more frequently had an M1 occlusion. Time from symptom onset to the end of EVT was shorter in those with successful recanalization. Finally, patients with successful recanalization more often underwent general anesthesia and were less frequently treated during shift hours. A total of 411 patients did not undergo intraarterial thrombectomy (IAT) due to the inability to obtain intracranial access (164 patients), spontaneous recanalization (244 patients) or for an unknown reason (3 patients). The remaining 2306 patients all underwent IAT, 1704 of whom (74%) had successful recanalization.

Relationship between collaterals and recanalization

In univariable analysis, there was no statistically significant association between collateral grades and the probability of successful recanalization (online supplemental table 1), and

	Unsuccessful recanalization* (n=819)	Successful recanalization† (n=1898)	P value
Age, median (IQR)	74 (63–82)	71 (61–80)	<0.01
Men, n (%)	417 (50.9)	1000 (52.7)	0.42
NIHSS, median (IQR)	16 (12–20)	16 (11–19)	0.31
SBP, mm Hg; mean (SD)	152 (26)	149 (24)	0.01
Glucose, mmol/L; median (IQR)	6.8 (5.9–8.2)	6.8 (5.9–8.0)	0.72
Diabetes mellitus, n (%)	131 (16.0)	306 (16.2)	0.99
Hypertension, n (%)	425 (52.9)	958 (51.6)	0.56
Atrial fibrillation, n (%)	218 (27.0)	423 (22.6)	0.02
Myocardial infarction, n (%)	98 (12.2)	269 (14.5)	0.14
Peripheral artery disease, n (%)	78 (9.8)	164 (8.8)	0.45
Ischemic stroke, n (%)	148 (18.2)	307 (16.3)	0.25
Hypercholesterolemia, n (%)	227 (29.1)	566 (31.1)	0.32
Pre-stroke mRS score ≤2, n (%)	691 (86.5)	1652 (89.1)	0.07
Statin use, n (%)	278 (34.7)	680 (36.7)	0.33
Antiplatelet use, n (%)	250 (30.9)	594 (31.7)	0.70
Coumarin use or DOAC, n (%)	150 (18.5)	283 (15.1)	0.03
Antihypertensive use, n (%)	433 (54.0)	1005 (54.0)	1.00
Current smoking, n (%)	170 (26.3)	413 (28.4)	0.33
Left hemisphere, n (%)	391 (47.8)	881 (46.4)	0.11
Occluded segment, n (%)			0.01
ICA	53 (6.8)	81 (4.4)	
ICA-T	177 (22.8)	366 (20.0)	
M1	430 (55.3)	1118 (61.2)	
M2	118 (15.2)	262 (14.3)	
Symptomatic carotid stenosis, n (%)	,		0.26
<50% stenosis at carotid bifurcation	564 (78.9)	1367 (81.3)	0.20
≥50% stenosis at carotid bifurcation	66 (9.2)	152 (9.0)	
Occlusion at carotid bifurcation	85 (11.9)	163 (9.7)	
ASPECTS on NCCT, median (IQR)	9 (7–10)	9 (8–10)	0.86
Collaterals, n (%)		5 (6 . 6)	0.23
Grade 0	53 (6.9)	101 (5.6)	0.20
Grade 1	272 (35.5)	647 (36.2)	
Grade 2	288 (37.6)	707 (39.5)	
Grade 3	153 (20.0)	334 (18.7)	
Intravenous alteplase, n (%)	618 (75.8)	1461 (77.3)	0.45
Time from onset to ER, min; median (IQR)	55 (40–101)	57 (37–95)	0.45
Time from onset to groin puncture, min; median (IQR)	196 (150–262)	190 (150–245)	0.47
Time from onset to successful recanalization, min; median (IQR)	270 (216–334)	240 (191–300)	<0.00
General anesthesia, n (%)	167 (21.9)	524 (29.1)	< 0.01
Treatment during shift hours, n (%)	540 (65.9)	1180 (62.2)	0.06
*Post-EVT eTICI 0, 1, or 2A:	<i>(כ.כ</i>) סדכ	1100 (02.2)	0.00

*Post-EVT eTICI 0, 1, or 2A;

†Post-EVT eTICI 2B, 2C, or 3.

ASPECTS, Alberta Stroke Program Early CT Score; DOAC, direct oral anticoagulant; ER, emergency room; eTICI, extended Thrombolysis in Cerebral Infarction; EVT, endovascular treatment; ICA, internal carotid artery; ICA-T, internal carotid artery terminus; M1/M2, middle cerebral artery; mRS, modified Rankin Scale; NCCT, non-contrast CT; NIHSS, National Institutes of Health Stroke Scale; SBP, systolic blood pressure.

this did not change after adjusting for potential confounders (table 2). The analyses for secondary outcomes showed consistent results (online supplemental table 1, table 2). Excellent collaterals (grade 3) were borderline associated with complete recanalization (eTICI 3) after EVT (acOR 1.49, 95% CI 0.99 to 2.25). Results were similar for the

subgroup of 2079 patients treated with intravenous alteplase prior to EVT and in the sensitivity analysis of 2306 patients who underwent IAT (online supplemental table 1, table 3). Results were also similar in sensitivity analyses with collateral grade as a numeric independent variable (online supplemental table 2).

Table 2 Multivariable associations of collateral grade with recanalization in all patients					
	Collateral grade	n/N	acOR (95% CI)	P value	
Primary outcome					
Successful recanalization*	0	101/154	Reference	-	
	1	647/919	1.19 (0.82 to 1.72)	0.37	
	2	707/995	1.20 (0.83 to 1.75)	0.33	
	3	334/487	1.10 (0.74 to 1.63)	0.64	
Secondary outcomes					
Complete recanalization†	0	42/154	Reference	-	
	1	292/919	1.21 (0.82 to 1.78)	0.35	
	2	323/995	1.25 (0.85 to 1.84)	0.26	
	3	174/487	1.49 (0.99 to 2.25)	0.05	
eTICI score (ordinal)	0	N/A	Reference	-	
	1	N/A	1.14 (0.84 to 1.56)	0.4	
	2	N/A	1.19 (0.87 to 1.61)	0.27	
	3	N/A	1.22 (0.88 to 1.68)	0.23	
Spontaneous recanalization‡	0	12/154	Reference	-	
	1	68/919	0.73 (0.37 to 1.41)	0.34	
	2	89/995	0.82 (0.43 to 1.57)	0.55	
	3	51/487	0.91 (0.46 to 1.82)	0.79	
First pass effect	0	26/116	Reference	-	
	1	173/692	0.88 (0.55 to 1.41)	0.6	
	2	184/751	0.91 (0.58 to 1.43)	0.68	
	3	103/357	1.21 (0.69 to 2.11)	0.5	

Collateral grade is treated as an ordered predictor variable in all analyses. Binary or ordinal logistic regression models were used. Effect estimates are adjusted common ORs for collateral grade 1 compared with 0, 2 compared with 0, and 3 compared with 0 (grade 0 being the reference category). Analyses were adjusted for age, blood pressure on admission, Alberta Stroke Program Early CT Score at baseline, extracranial carotid artery disease, occluded segment, and time from onset to recanalization (or last contrast bolus). *eTICI >2B after EVT.

teTICI 3 after EVT.

‡eTICI 2B-3 on diagnostic DSA.

eTICI, extended Thrombolysis in Cerebral Infarction; EVT, endovascular treatment.

Relationship between recanalization and functional outcome

In multivariable analysis, both successful recanalization (acOR 2.15, 95% CI 1.84 to 2.52) and better collateral grades (grade 1 vs 0, acOR 2.12, 95% CI 1.47 to 3.05; grade 2 vs 0, acOR 3.46, 95% CI 2.43 to 4.92; grade 3 vs 0, acOR 4.16, 95% CI 2.89 to 5.99) were independently associated with a shift towards good functional outcome. Patients with unsuccessful recanalization and absent (grade 0) collaterals had the poorest outcomes (6% functional independence), whereas the best outcomes were achieved by those with successful recanalization and excellent (grade 3) collaterals (61% functional independence). There was no interaction between collateral grades and successful recanalization (likelihood ratio test p=0.78). Figure 1 depicts the distribution of mRS scores according to collateral grades and recanalization status after EVT.

Results for the subgroup analysis of patients treated with intravenous alteplase were similar: successful recanalization (acOR 2.24, 95% CI 1.87 to 1.47) and better collateral grades (grade 1 vs 0, acOR 2.06, 95% CI 1.37 to 3.09; grade 2 vs 0, acOR 3.34, 95% CI 2.23 to 4.99; grade 3 vs 0, acOR 3.92, 95% CI 2.59 to 5.93) were independently associated with a shift towards favorable functional outcome and there was no interaction between those variables. Similar results were also found for patients undergoing IAT (acOR for collateral grade 1 vs 0, 2.46, 95% CI 1.72 to 3.53; grade 2 vs 0, acOR 4.09, 95% CI 2.84 to 5.89; grade 3 vs 0, acOR 4.61, 95% CI 3.10 to 6.85; acOR for

successful recanalization, 1.99, 95% CI 1.67 to 2.37, without a statistical interaction).

Results were similar when collateral status was analyzed as a numeric independent variable in all patients (acOR for a shift towards good functional outcome: 1.51, 95% CI 1.38 to 1.65 per one-point increase in collateral grade; 2.18, 95% CI 1.84 to 2.58 for successful recanalization), in the subgroup of patients treated with intravenous thrombolysis (cOR for a shift towards good functional outcome: 1.48, 95% CI 1.34 to 1.65 per onepoint increase in collateral grade; 2.28, 95% CI 1.88 to 2.77 for successful recanalization), and in the subgroup of patients undergoing IAT (cOR for a shift towards good functional outcome: 1.52, 95% CI 1.37 to 1.68 per one-point increase in collateral grade; 2.01, 95% CI 1.66 to 2.43 for successful recanalization), without interactions between collateral status and successful recanalization.

DISCUSSION

In this large cohort of patients with large vessel occlusion treated with EVT, there was no relation between collateral status at baseline and the probability of spontaneous recanalization before EVT or successful recanalization after EVT. Higher collateral grades and successful recanalization were each associated with better functional outcome, but there was no statistical interaction between these variables. This suggests that achieving successful recanalization after EVT is similarly beneficial for patients with

Table 3	Multivariable associations of collateral grade with recanalization in patients who were treated with intravenous thrombolysis (n=2079)
and patie	ents who underwent intra-arterial treatment (n=2306)

	IVT subgroup			IAT subgroup			
	Collateral grade	n/N	acOR (95% CI)	P value	n/N	acOR (95% CI)	P value
Successful recanalization*	0	72/109	Reference	-	89/131	Reference	-
	1	515/712	1.26 (0.80 to 1.98)	0.32	596/796	1.36 (0.90 to 2.05)	0.14
	2	537/758	1.17 (0.74 to 1.83)	0.5	636/847	1.38 (0.91 to 2.10)	0.13
	3	250/317	0.99 (0.62 to 1.59)	0.97	295/411	1.23 (0.78 to 1.92)	0.37
Complete recanalization†	0	29/109	Reference	_	41/131	Reference	_
	1	241/712	1.36 (0.86 to 2.17)	0.19	280/796	1.12 (0.75 to 1.67)	0.58
	2	240/758	1.26 (0.79 to 1.99)	0.33	315/847	1.23 (0.83 to 1.83)	0.31
	3	128/371	1.43 (0.89 to 2.32)	0.14	165/411	1.45 (0.94 to 2.22)	0.09
eTICI score (ordinal)	0	N/A	Reference	_	N/A	Reference	_
	1		1.24 (0.86 to 1.79)	0.25		1.16 (0.84 to 1.62)	0.36
	2		1.14 (0.80 to 1.64)	0.46		1.26 (0.90 to 1.76)	0.17
	3		1.11 (0.76 to 1.63)	0.58		1.27 (0.89 to 1.82)	0.19
Spontaneous recanalization‡	0	12/109	Reference	_	N/A	N/A	N/A
	1	60/712	0.61 (0.31 to 1.19)	0.14			
	2	79/758	0.73 (0.37 to 1.41)	0.35			
	3	44/371	0.78 (0.39 to 1.57)	0.49			
First pass effect	0	20/79	Reference	_	26/116	Reference	_
	1	144/533	0.81 (0.47 to 1.41)	0.46	173/692	0.99 (0.58 to 1.69)	0.96
	2	142/551	0.78 (0.46 to 1.32)	0.36	184/751	1.00 (0.59 to 1.69)	0.99
	3	82/263	1.09 (0.57 to 2.10)	0.78	103/357	1.29 (0.70 to 2.37)	0.42

Collateral grade is treated as an ordered predictor variable in all analyses. Binary or ordinal logistic regression models were used. Effect estimates are adjusted common ORs for collateral grade 1 compared with 0, 2 compared with 0, and 3 compared with 0 (grade 0 being the reference category). Analyses were adjusted for age, blood pressure on admission, Alberta Stroke Program Early CT Score at baseline, extracranial carotid artery disease, occluded segment, and time from onset to recanalization (or last contrast bolus) *eTICI \geq 2B after EVT.

†eTICI 3 after EVT.

‡eTICI 2B-3 on diagnostic DSA.

eTICI, extended Thrombolysis in Cerebral Infarction; EVT, endovascular treatment; IAT, intra-arterial treatment; IVT, intravenous thrombolysis.

different collateral grades at baseline. As expected, the best outcomes were achieved by those with the highest collateral grade who were successfully recanalized during EVT (61% functional independence).

Our findings contradict several previous studies, including two studies with recruitment periods similar to ours,^{10 11} that found an association between good collateral status and better tissue reperfusion or vessel recanalization after IV thrombolysis (IVT) alone^{5 6 17} or EVT with or without prior IVT.^{7-11 18} We tested the hypothesis that the collateral circulation may facilitate the delivery of thrombolytics to the distal end of the clot, thereby enhancing recanalization, but we also found no association in the subgroup of patients treated with IVT. The absence of an association between collateral status and recanalization appeared to be independent of occlusion location. Most previous studies differed from ours in their dichotomization of collateral status into good versus poor, whereas we analyzed collateral status with the full ordinal scale in order to be able to detect a trend. Moreover, some studies used different imaging modalities and a local, rather than central, assessment of imaging variables. Nonetheless, we cannot fully explain the absence of a previously established relationship between collateral status and the probability of achieving successful recanalization during EVT.

Sufficient flow into collateral vessels can preserve viability of the downstream territory until recanalization occurs, so it has been hypothesized that the effect of collateral status on outcome

is greater for recanalized patients than for non-recanalized patients. Vice versa, achieving successful recanalization may be more beneficial for those with a good collateral status than for those with a poor collateral status. This hypothesis was confirmed in the Interventional Management of Stroke III trial, where better collateral grades were strongly associated with a good functional outcome only in patients with recanalization (modified TICI ≥ 2), but not in those without recanalization.⁷ Several additional studies suggested a similar effect modification.9 12-14 19 By contrast, we found no interaction between collateral grades and successful recanalization in the analysis for functional outcome. This suggests that the effect of successful recanalization on functional outcome is similar for all collateral grades and there is no synergistic effect of the two. Our finding is in line with several previous studies,^{11 20 21} including a recent analysis from 2020 patients in the multicenter Endovascular Treatment in Ischemic Stroke (ETIS) Registry, the previously mentioned ETIS study.¹¹ The comparability of the studies is limited because of the use of various different grading scales for collateral status.

Of note, our rate of successful recanalization after EVT of 70% was lower than that in most previous studies, with recent studies reporting rates as high as 85% to 90%.^{10 11 22 23} In contrast to most registries, patients eligible for EVT were included consecutively in our registry so our cohort represents an unselected group of EVT-eligible patients, which could explain the difference.

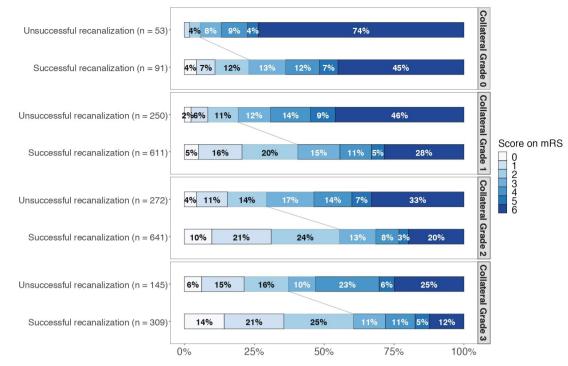


Figure 1 Distribution of modified Rankin Scale scores at 90 days according to collateral grades and recanalization status post-EVT. Lines indicate the difference between functional independence (mRS score \leq 2) and no functional independence (mRS score \geq 3). The benefit of successful recanalization on the mRS score is approximately equal in all collateral grade groups (as indicated by the parallel lines), suggesting that no effect modification is present. mRS, modified Rankin Scale.

The main strengths of this study are the large sample size and the fact that imaging data were prospectively collected and analyzed centrally by a core laboratory of trained neuroradiologists. Limitations include the use of single-phase CTA to assess collateral status, which could have led to misclassification of collateral status in cases of early or late imaging acquisition.²⁴ However, differential misclassification is unlikely and therefore it is doubtful that it influenced our results. Second, we measured only vessel recanalization-the reinstatement of antegrade flow in the affected vascular territory-using angiography. Since vessel recanalization does not always result in subsequent tissue reperfusion, the latter being assessed with perfusion imaging, we cannot exclude the possibility that the results of this study might have been different if we had measured reperfusion. Nonetheless, in a previous study, the association between recanalization and reperfusion was similar regardless of collateral status.²⁵ Third, the primary analyses also included patients in whom no IAT attempt was made due to the inability to obtain intracranial access or spontaneous recanalization. However, sensitivity analyses of patients who did undergo IAT showed similar results, mitigating the concern that this could explain the neutral findings. Fourth, collateral status was analyzed both as an ordinal and as a numeric variable in our study, which differs from most previous studies that dichotomized collateral status and therefore direct comparisons cannot be made. However, our approach prevented the loss of information that results from merging collateral grade categories.

CONCLUSIONS

We found no evidence that better collateral status at baseline facilitates vessel recanalization after EVT. Successful recanalization and better collateral grades were independently associated with improved functional outcome, but there was no statistical interaction between them. Therefore, the effect of successful recanalization on outcome appears to be independent of baseline collateral grade.

Acknowledgements We would like to thank the MR CLEAN Registry investigators; a complete list of the MR CLEAN Registry investigators is provided in the Appendix.

Collaborators MR CLEAN Registry investigators-Executive committee: Diederik W.J. Dippel (Department of Neurology); Aad van der Lugt (Radiology); Charles B.L.M. Majoie (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Yvo B.W.E.M. Roos (Neurology); Robert J. van Oostenbrugge (Amsterdam UMC, University of Amsterdam, Amsterdam; Department of Neurology); Wim H. van Zwam (Radiology); Jelis Boiten (Neurology); Jan Albert Vos (Radiology) Study coordinators: Ivo G.H. Jansen (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Maxim J.H.L. Mulder (Department of Neurology), (Radiology); Robert- Jan B. Goldhoorn (Amsterdam UMC, University of Amsterdam, Amsterdam; Department of Neurology), (Radiology); Kars C.J. Compagne (Radiology); Manon Kappelhof (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Josje Brouwer (Neurology); Sanne J. den Hartog (Department of Neurology), (Radiology), (Public Health); Wouter H. Hinsenveld (Amsterdam UMC, University of Amsterdam, Amsterdam; Department of Neurology), (Radiology); Local principal investigators: Diederik W.J. Dippel (Department of Neurology); Bob Roozenbeek (Department of Neurology); Aad van der Lugt (Radiology); Adriaan C.G.M. van Es (Radiology); Charles B.L.M. Majoie (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Yvo B.W.E.M. Roos (Neurology); Bart J. Emmer (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Jonathan M. Coutinho (Neurology); Wouter J. Schonewille (Maastricht University Medical Center and Cardiovascular Research Institute Maastricht (CARIM); Department of Neurology); Jan Albert Vos (Radiology); Marieke J.H. Wermer (Sint Antonius Hospital, Nieuwegein; Department of Neurology); Marianne A.A. van Walderveen (Radiology); Julie Staals (Amsterdam UMC, University of Amsterdam, Amsterdam; Department of Neurology); Robert J. van Oostenbrugge (Amsterdam UMC, University of Amsterdam, Amsterdam; Department of Neurology); Wim H. van Zwam (Radiology); Jeannette Hofmeijer (Leiden University Medical Center; Department of Neurology); Jasper M. Martens (Radiology); Geert J. Lycklama à Nijeholt (Rijnstate Hospital, Arnhem; Department of Radiology); Jelis Boiten (Neurology); Sebastiaan F. de Bruijn

Neuroimaging

(Haaglanden MC, the Hague; Department of Neurology); Lukas C. van Dijk1 (Radiology): H. Bart van der Worp (HAGA Hospital, the Hague: Department of Neurology); Rob H. Lo (Radiology); Ewoud J. van Dijk (University Medical Center Utrecht; Department of Neurology); Hieronymus D. Boogaarts (Neurosurgery); J. de Vries (Elisabeth-TweeSteden ziekenhuis, Tilburg; Department of Neurology); Paul L.M. de Kort (Radboud University Medical Center, Nijmegen; Department of Neurology); Julia van Tuiil (Radboud University Medical Center, Niimegen: Department of Neurology); Jo P. Peluso (Radiology); Puck Fransen (Elisabeth-TweeSteden ziekenhuis, Tilburg; Department of Neurology); Jan S.P. van den Berg (Elisabeth-TweeSteden ziekenhuis, Tilburg; Department of Neurology); Boudewijn A.A.M. van Hasselt (Radiology); Leo A.M. Aerden (Isala Klinieken, Zwolle; Department of Neurology); René J. Dallinga (Radiology); Maarten Uyttenboogaart (Reinier de Graaf Gasthuis, Delft; Department of Neurology); Omid Eschgi (Radiology); Reinoud P.H. Bokkers (Radiology); Tobien H.C.M.L. Schreuder (University Medical Center Groningen; Department of Neurology); Roel J.J. Heijboer (Radiology); Koos Keizer (Atrium Medical Center, Heerlen; Department of Neurology); Lonneke S.F. Yo (Radiology); Heleen M. den Hertog (Elisabeth-TweeSteden ziekenhuis, Tilburg: Department of Neurology); Tomas Bulut (Radiology); Paul J.A.M. Brouwers (Catharina Hospital, Eindhoven; Department of Neurology). Imaging assessment committee: Charles B.L.M. Majoie (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine)(chair); Wim H. van Zwam (Radiology); Aad van der Lugt (Radiology): Geert J. Lycklama à Nijeholt (Rijnstate Hospital, Arnhem: Department of Radiology); Marianne A.A. van Walderveen (Radiology); Marieke E.S. Sprengers (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Sjoerd F.M. Jenniskens (Radiology); René van den Berg (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Albert J. Yoo (Department of Radiology, Texas Stroke Institute, Texas, United States of America); Ludo F.M. Beenen (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Alida A. Postma (Radiology); Stefan D. Roosendaal (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Bas F.W. van der Kallen (Rijnstate Hospital, Arnhem; Department of Radiology): Ido R. van den Wiingaard (Riinstate Hospital, Arnhem: Department of Radiology); Adriaan C.G.M. van Es (Radiology); Bart J. Emmer, (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Jasper M. Martens (Radiology); Lonneke S.F. Yo (Radiology); Jan Albert Vos (Radiology); Joost Bot (Medisch Spectrum Twente, Enschede; Department of Radiology), Pieter-Jan van Doormaal (Radiology): Anton Meijer (Radiology): Elvas Gharig (Rijnstate Hospital, Arnhem; Department of Radiology); Reinoud P.H. Bokkers (Radiology); Marc P. van Proosdij (Amsterdam UMC, Vrije Universiteit van Amsterdam, Amsterdam; Department of Radiology);G. Menno Krietemeijer (Radiology);Jo P. Peluso (Radiology); Hieronymus D. Boogaarts (Neurosurgery); Rob Lo (Radiology); Wouter Dinkelaar (Radiology); Auke P.A. Appelman (Radiology); Bas Hammer1 (Radiology); Sjoert Pegge (Radiology); Anouk van der Hoorn (Radiology); Saman Vinke (Neurosurgery). Writing committee: Diederik W.J. Dippel (Department of Neurology) (chair); Aad van der Lugt (Radiology); Charles B.L.M. Majoie (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Yvo B.W.E.M. Roos (Neurology); Robert J. van Oostenbrugge (Amsterdam UMC, University of Amsterdam, Amsterdam; Department of Neurology); Wim H. van Zwam (Radiology); Geert J. Lycklama à Nijeholt (Rijnstate Hospital, Arnhem; Department of Radiology); Jelis Boiten (Neurology); Jan Albert Vos (Radiology); Wouter J. Schonewille (Maastricht University Medical Center and Cardiovascular Research Institute Maastricht (CARIM); Department of Neurology); Jeannette Hofmeijer (Leiden University Medical Center; Department of Neurology); Jasper M. Martens (Radiology); H. Bart van der Worp (HAGA Hospital, the Hague; Department of Neurology); Rob H. Lo (Radiology) Adverse event committee: Robert J. van Oostenbrugge (Amsterdam UMC, University of Amsterdam, Amsterdam; Department of Neurology) (chair); Jeannette Hofmeijer (Leiden University Medical Center; Department of Neurology); H. Zwenneke Flach (Radiology); Trial methodologist: Hester F. Lingsma (Public Health) Research nurses / local trial coordinators: Naziha el Ghannouti (Department of Neurology); Martin Sterrenberg1; Wilma Pellikaan (Maastricht University Medical Center and Cardiovascular Research Institute Maastricht (CARIM); Department of Neurology); Rita Sprengers (Neurology); Marjan Elfrink (Leiden University Medical Center; Department of Neurology); Michelle Simons (Leiden University Medical Center; Department of Neurology); Marjolein Vossers (Radiology); Joke de Meris (Neurology); Tamara Vermeulen (Neurology); Annet Geerlings (University Medical Center Utrecht; Department of Neurology); Gina van Vemde (Elisabeth-TweeSteden ziekenhuis, Tilburg; Department of Neurology); Tiny Simons (University Medical Center Groningen; Department of Neurology); Gert Messchendorp (Reinier de Graaf Gasthuis, Delft; Department of Neurology);Nynke Nicolaij (Reinier de Graaf Gasthuis, Delft; Department of Neurology);Hester

Bongenaar (Atrium Medical Center, Heerlen; Department of Neurology):Karin Bodde (Isala Klinieken, Zwolle; Department of Neurology);Sandra Kleijn (Catharina Hospital, Eindhoven; Department of Neurology); Jasmijn Lodico (Catharina Hospital, Eindhoven; Department of Neurology); Hanneke Droste (Catharina Hospital, Eindhoven; Department of Neurology); Maureen Wollaert (Amsterdam UMC, University of Amsterdam, Amsterdam; Department of Neurology); Sabrina Verheesen (Amsterdam UMC, University of Amsterdam, Amsterdam; Department of Neurology); D. Jeurrissen (Amsterdam UMC, University of Amsterdam, Amsterdam; Department of Neurology); Erna Bos (Sint Antonius Hospital, Nieuwegein; Department of Neurology); Yvonne Drabbe (Haaglanden MC, the Hague; Department of Neurology); Michelle Sandiman (Haaglanden MC, the Hague; Department of Neurology); Nicoline Aaldering (Leiden University Medical Center; Department of Neurology); Berber Zweedijk (HAGA Hospital, the Hague; Department of Neurology); Jocova Vervoort (Radboud University Medical Center, Nijmegen; Department of Neurology); Eva Ponjee (Elisabeth-TweeSteden ziekenhuis, Tilburg; Department of Neurology); Sharon Romviel (University Medical Center Utrecht; Department of Neurology); Karin Kanselaar (University Medical Center Utrecht; Department of Neurology); Denn Barning (Radiology), PhD / Medical students: Esmee Venema (Public Health): Vicky Chalos(Department of Neurology), (Public Health); Ralph R. Geuskens (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Tim van Straaten (University Medical Center Utrecht; Department of Neurology); Saliha Ergezen (Department of Neurology); Roger R.M. Harmsma(Department of Neurology); Daan Muijres(Department of Neurology); Anouk de Jong (Department of Neurology); Olvert A. Berkhemer (Department of Neurology), (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine), (Radiology); Anna M.M. Boers (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine), (Biomedical Engineering & Physics); J. Huguet (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); P.F.C. Groot (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Marieke A. Mens (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Katinka R. van Kranendonk (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Kilian M. Treurniet (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Manon L. Tolhuisen (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine), (Biomedical Engineering & Physics); Heitor Alves (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Annick J. Weterings (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine), Eleonora L.F. Kirkels (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine), Eva J.H.F. Voogd (Leiden University Medical Center; Department of Neurology); Lieve M. Schupp (Erasmus MC University Medical Center; Department of Radiology and Nuclear Medicine); Sabine L. Collette (Reinier de Graaf Gasthuis, Delft; Department of Neurology), (Radiology); Adrien E.D. Groot (Neurology); Natalie E. LeCouffe (Neurology); Praneeta R. Konduri (Biomedical Engineering & Physics); Haryadi Prasetya (Biomedical Engineering & Physics); Nerea Arrarte-Terreros (Biomedical Engineering & Physics); Lucas A. Ramos (Biomedical Engineering & Physics).

Contributors Simone M. Uniken Venema: study concept, statistical analysis, interpretation of the results, drafting of the manuscript. Jan Willem Dankbaar, Lennard Wolff, Adriaan C. G. M. van Es, Marieke Sprengers, Aad van der Lugt, Diederik W. J. Dippel: critical revision of the manuscript. H. Bart van der Worp: study concept, interpretation of the results, critical revision of the manuscript. H. Bart van der Worp is the guarantor of this work.

Funding The MR CLEAN Registry was partly funded by TWIN (Toegepast Wetenschappelijk Instituut voor Neuromodulatie) Foundation, Erasmus MC Medical Center, Maastricht University Medical Center, and Academic Medical Center Amsterdam. Simone M. Uniken Venema performed this work as coordinator of the trial MR ASAP, which is funded by Netherlands Cardiovascular Research Initiative, an initiative from the Dutch Heart Foundation.

Competing interests Simone M. Uniken Venema, Jan Willem Dankbaar, Lennard Wolff, Adriaan C. G. M. van Es, Marieke Sprengers report no disclosures; Diederik W. J. Dippel: reports funding from the Dutch Heart Foundation, Brain Foundation Netherlands, The Netherlands Organisation for Health Research and Development, Health Holland Top Sector Life Sciences and Health, and unrestricted grants from Penumbra Inc., Stryker, Medtronic, Cerenovus and Thrombolytic Science, LLC for research, all paid to his institution; Aad van der Lugt: reports funding from the Dutch Heart Foundation, Brain Foundation Netherlands, The Netherlands Organisation for Health Research and Development, Health Holland Top Sector Life Sciences and Health, and unrestricted grants from Penumbra Inc., Stryker, Medtronic, Cerenovus, and Thrombolytic Science, LLC for research, all paid to his institution; H. Bart van der Worp: received fees for consultation for Bayer and LivaNova, all paid to his

Neuroimaging

institution.

Patient consent for publication Not applicable.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data are available upon reasonable request. Source data from patients cannot be distributed, but syntax and output files from statistical analyses can be made available upon reasonable request to the corresponding author.

Supplemental material This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation and adaptation or otherwise.

ORCID iDs

Simone M Uniken Venema http://orcid.org/0000-0002-5374-3858 Diederik W J Dippel http://orcid.org/0000-0002-9234-3515

REFERENCES

- LeCouffe NE, Kappelhof M, Treurniet KM, et al. 2B, 2C, or 3: what should be the angiographic target for endovascular treatment in ischemic stroke? *Stroke* 2020;51:1790–6.
- 2 Liebeskind DS, Bracard S, Guillemin F, et al. eTICI reperfusion: defining success in endovascular stroke therapy. J Neurointerv Surg 2019;11:433–8.
- 3 Berkhemer OA, Jansen IGH, Beumer D, et al. Collateral status on baseline computed tomographic angiography and intra-arterial treatment effect in patients with proximal anterior circulation stroke. Stroke 2016;47:768–76.
- 4 Venema E, Mulder MJHL, Roozenbeek B, et al. Selection of patients for intra-arterial treatment for acute ischaemic stroke: development and validation of a clinical decision tool in two randomised trials. BMJ 2017;357:j1710.
- 5 Nicoli F, Lafaye de Micheaux P, Girard N. Perfusion-weighted imaging-derived collateral flow index is a predictor of MCA M1 recanalization after IV. thrombolysis. *AJNR Am J Neuroradiol* 2013;34:107–14.
- 6 Seners P, Roca P, Legrand L, *et al*. Better collaterals are independently associated with post-thrombolysis recanalization before thrombectomy. *Stroke* 2019;50:867–72.
- 7 Liebeskind DS, Tomsick TA, Foster LD, et al. Collaterals at angiography and outcomes in the Interventional Management of Stroke (IMS) III trial. Stroke 2014;45:759–64.
- 8 Liebeskind DS, Jahan R, Nogueira RG, et al. Impact of collaterals on successful revascularization in Solitaire FR with the intention for thrombectomy. *Stroke* 2014;45:2036–40.
- 9 Bang OY, Saver JL, Kim SJ, et al. Collateral flow predicts response to endovascular therapy for acute ischemic stroke. *Stroke* 2011;42:693–9.

- 10 García-Tornel Álvaro, Ciolli L, Rubiera M, et al. Leptomeningeal collateral flow modifies endovascular treatment efficacy on large-vessel occlusion strokes. Stroke 2021;52:STROKEAHA120031338.
- 11 Anadani M, Finitsis S, Clarençon F, et al. Collateral status reperfusion and outcomes after endovascular therapy: insight from the Endovascular Treatment in Ischemic Stroke (ETIS) Registry. J Neurointerv Surg 2021;53. doi:10.1136/ neurintsurg-2021-017553. [Epub ahead of print: 17 Jun 2021].
- 12 Gersing AS, Schwaiger BJ, Kleine JF, et al. Clinical outcome predicted by collaterals depends on technical success of mechanical thrombectomy in middle cerebral artery occlusion. J Stroke Cerebrovasc Dis 2017;26:801–8.
- 13 Mangiafico S, Saia V, Nencini P, et al. Effect of the interaction between recanalization and collateral circulation on functional outcome in acute ischaemic stroke. *Interv Neuroradiol* 2014;20:704–14.
- 14 Nambiar V, Sohn SI, Almekhlafi MA, et al. CTA collateral status and response to recanalization in patients with acute ischemic stroke. Am J Neuroradiol 2014;35:884–90.
- 15 Tan IYL, 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;30:525–31.
- 16 Noser EA, Shaltoni HM, Hall CE, et al. Aggressive mechanical clot disruption: a safe adjunct to thrombolytic therapy in acute stroke? Stroke 2005;36:292–6.
- 17 Zhang S, Zhang X, Yan S, *et al*. The velocity of collateral filling predicts recanalization in acute ischemic stroke after intravenous thrombolysis. *Sci Rep* 2016;6:1–7.
- 18 Leng X, Fang H, Leung TWH, et al. Impact of collaterals on the efficacy and safety of endovascular treatment in acute ischaemic stroke: a systematic review and metaanalysis. J Neurol Neurosurg Psychiatry 2016;87:537–44.
- 19 Miteff F, Levi CR, Bateman GA, et al. The independent predictive utility of computed tomography angiographic collateral status in acute ischaemic stroke. Brain 2009;132:2231–8.
- 20 Marks MP, Lansberg MG, Mlynash M, et al. Effect of collateral blood flow on patients undergoing endovascular therapy for acute ischemic stroke. Stroke 2014;45:1035–9.
- 21 Gerber JC, Petrova M, Krukowski P, et al. Collateral state and the effect of endovascular reperfusion therapy on clinical outcome in ischemic stroke patients. *Brain Behav* 2016;6:1–9.
- 22 Zaidat OO, Fifi JT, Nanda A, *et al.* Endovascular treatment of acute ischemic stroke with the Penumbra system in routine practice: complete registry results. *Stroke* 2022;53:769–78.
- 23 Mueller-Kronast NH, Zaidat OO, Froehler MT, et al. Systematic evaluation of patients treated with Neurothrombectomy devices for acute ischemic stroke: primary results of the STRATIS registry. Stroke 2017;48:2760–8.
- 24 Menon BK, Smith EE, Modi J, et al. Regional leptomeningeal score on CT angiography predicts clinical and imaging outcomes in patients with acute anterior circulation occlusions. AJNR Am J Neuroradiol 2011;32:1640–5.
- 25 Villringer K, Zimny S, Galinovic I, et al. The association between recanalization, collateral flow, and reperfusion in acute stroke patients: a dynamic susceptibility contrast MRI study. Front Neurol 2019;10:1–8.