

ORIGINAL RESEARCH

Mechanical thrombectomy for basilar artery occlusion: efficacy, outcomes, and futile recanalization in comparison with the anterior circulation

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ABSTRACT

currently not evidence-based.

large vessel occlusion (ACLVO) MT.

as adjusted Odds Ratios, aOR and 95% confidence intervals, 95%-CI) outcome comparisons between BAO (N=165) and ACLVO (N=1574) were performed. The primary outcome was favorable outcome at 90 days (modified Rankin Scale, mRS 0-2). Secondary outcome included mortality, symptomatic intracranial hemorrhage (sICH) and FR. The relative merits of achieving successful recanalization between ACLVO and BAO were evaluated with interaction terms.

Background Performing mechanical thrombectomy

(MT) in patients with basilar artery occlusion (BAO) is

Objective To compare patients' outcome, relative

merits of achieving recanalization, and predictors of futile

recanalization (FR) between BAO and anterior circulation

Methods In the multicenter BEYOND-SWIFT registry

(NCT03496064), univariate and multivariate (displayed

Results MT in BAO was more often technically effective and equally safe in regards to mortality and sICH when compared to ACLVO. When adjusting for baseline differences, there was no significant difference between BAO vs ACLVO regarding rates of favorable outcome (aOR 0.986, 95%-CI 0.553 - 1.758). However, BAO were associated with increased rates of FR (aOR 2.146, 95%-CI 1.267 – 3.633). Predictors for FR were age, stroke severity, maneuver count and intracranial stenting. No significant heterogeneity on the relative merits of achieving successful recanalization on several outcome parameters were observed when comparing BAO and ACLVO.

Conclusions In selected patients, similar outcomes can be achieved in BAO and ACLVO patients treated with MT. Randomized controlled trials comparing patient selection and interventional strategies seem warranted to avoid FR.

Trial registration number NCT03496064

INTRODUCTION

About 1% of patients with acute ischemic stroke have a basilar artery occlusion (BAO) that is amendable

to endovascular treatment.¹² Such patients present with particularly severe symptoms and have a poor prognosis compared with those with anterior circulation large vessel occlusion (ACLVO).³⁴

In patients with BAO and other large vessel occlusions (LVOs) of the posterior circulation (PC), indications for mechanical thrombectomy (MT) lack evidence derived from large pivotal thrombectomy trials.⁵ ⁶ Intravenous thrombolysis (IVT) is therefore the standard of care in those patients.⁷ Yet, IVT yields insufficient acute reperfusion rates in patients with BAO.^{8–13} Therefore, many patients subsequently deteriorate clinically and tend to have poor outcomes if reperfusion is not achieved.⁸¹⁴

Recanalization rates of MT were reported to be lower than with AC occlusions and procedure time longer.¹⁵ Furthermore in BAO, poor outcome despite successful recanalization by MT was reported to occur more frequently—a phenomenon termed futile recanalization (FR) (modified Rankin Scale (mRS) score 4-6 despite complete recanalization).¹⁵⁻¹⁸ In this setting, ethics and patient preferences are taken into account when considering the risk of subjecting patients to a treatment that may avoid death but instead creates long-term dependence.^{19 20} Efforts have been made to identify predictors of FR in BAO-namely, stroke severity on admission, amount of radiological lesion on baseline MRI or CT, respiratory instability, and collateral quality.^{16–18 21 22} Despite those efforts, the rates of long-term dependence were reported to be around 30%²³ after MT in BAO.

Although several studies have reported predictors of outcome after endovascular therapy in patients with BAO, few studies have focused solely on MT. Furthermore, reports were limited by small numbers of patients, non-transparent patient selection, heterogeneous endovascular treatments, and inclusion of patients with more distal occlusions.

The aims of this study were hence to compare patients' outcome, the relative merits of achieving recanalization, predictors of FR, and procedural safety between BAO and ACLVO MT, and predictors of FR in BAO.



METHODS

Details of the BEYOND-SWIFT Registry are in the public domain (NCT03496064) and have been published previously.²⁴ Briefly, the registry is a retrospective, multicenter, non-randomized observational study designed to investigate the safety and efficacy of a Medtronic market-released neurothrombectomy device (applied as the initial device used for interventions) in patients with acute ischemic stroke who do not fulfill treatment eligibility requirements according to the current guidelines. An overview of the patients included in the registry, the follow-up data for each center, and ethical approval procedures can be found in online supplementary table I. Ethical approval was obtained in Bern for pooling and analyses of the registry data (KEK Bern, Bern, Switzerland ID 2018–00766).

Most patients included in the registry (n=2046) were treated for large-vessel anterior circulation strokes (n=1832). Of the patients treated for BAO (n=165), 152 (92.1%) had documented 90-day follow-up. Selection criteria for performing MT in BAO were center-specific and are presented in online supplementary table 5.

Variables and image analysis

Local investigators categorized the site of LVO into vertebral artery, basilar artery, or first/second segment of the posterior cerebral artery (P1/P2) occlusions. For the main analysis we compared patients with BAO with those with iICA/carotid-T/ M1-occlusions. Either the operator at each center or an independent research fellow rated the postprocedural modified Thrombolysis in Cerebral Infarction (mTICI) score (see online supplementary table I). mTICI 2b was rated as reperfusion of at least 50% of the initially occluded target vessel territory, according to modifications of the TICI scale.²⁵ For clinical outcome evaluation, we assessed 3-month functional outcomes using the mRS score in routinely scheduled clinical visits or standardized telephone interviews organized at each center. FR was defined as successful recanalization in patients with mRS score 4-6 at 90 days.²⁰ Worsening in the National Institutes of Health Stroke Scale (NIHSS) score of ≥ 4 points between the admission NIHSS score and the score 24 hours later, and not attributable to an intracerebral hemorrhagic event was defined as non-hemorrhagic worsening.²⁶

Statistical analysis

Univariate comparisons between patients with BAO or ACLVO, between patients with favorable outcome (mRS score 0–2) and patients with unfavorable outcome (mRS score 3–6), as well as between patients with futile and non-futile recanalization were made using standard statistical measures (Fisher's exact test for categorical variables, Mann–Whitney U test for non-normally continuous or ordinally scaled variables, and Welsch's t-test for independent normally distributed data).

The primary endpoint of this analysis was favorable outcome at 90 days (mRS score 0–2) in BAO versus ACLVO according to multivariate analysis. Secondary outcomes consisted of rate of successful recanalization, FR, all-cause mortality at day 90, overall complication rate, and symptomatic intracerebral hemorrhage (sICH), which was assessed at each center by applying the ECASS II criteria, although they were originally established for AC strokes. The association between BAO *versus* ACLVO with all of the previously listed outcome parameters was hence assessed using multivariable logistic regression with adjustments for the following confounders: age (continuous), prestroke independence (categorical), NIHSS score on admission (ordinal, adjusted odds ratio (aOR) per point increase), systolic blood pressure (metric, aOR per mm Hg increase), known onset (categorical), risk factor hypertension (categorical), risk factor smoking (categorical), risk factor previous stroke (categorical), risk factor diabetes (categorical), type of admission imaging (CT vs MRI, categorical), IVT (categorical), and time from symptom onset to admission (metric, aOR per minute increase). The adjustments were made according to significant baseline differences in univariate analysis, except for admission glucose levels and history of coronary heart disease since too many data items were missing.

A potential difference in the relative merits of achieving successful recanalization between BAO and ACLVO was evaluated with interaction terms. For this comparison, we assessed the relative effect of successful recanalization on various dichotomizations of functional outcome and safety parameters using multivariable logistic regression with adjustments for the following prespecified confounders: age (continuous), sex (categorical), NIHSS score on admission (ordinal, aOR per point increase) and IVT (categorical). The adjustments for this analysis were made according to a clinical assumption of the main confounders as the restricted number of patients without successful recanalization limited the number of variables that could be included.

RESULTS

Baseline

For baseline differences, the 165 patients with BAO were younger, more often male, more often transferred from other hospitals, had more severe symptoms on admission, higher glucose on admission, less often anticoagulation pretreatment, less often hypertension, less often dyslipidemia, other TOAST etiology, and MRI was less often performed as the initial imaging modality (table 1) as compared with ACLVO. Furthermore, time from symptom onset to groin puncture was longer in BAO than in ACLVO (median 300 min vs 225 min, p<0.001).

Technical efficacy

Comparison of MT in patients with BAO and patients with ACLVO showed that the procedure was more often effective in BAO, with achievement of a mTICI 3 recanalization in 61.8% and a mTICI 2b/3 in 90.3% of patients requiring fewer maneuvers (p=0.002). Groin puncture to recanalization intervals in patients with BAO were similar to those in ACLVO (median $45 \min vs 47 \min, p=0.824$). However, more intracranial stents were used in patients with BAO (17.0% vs 2.3%, p<0.001) and longer times from symptom onset to groin-puncture (300 min vs 225 min, p<0.001) were noted, probably because interventions usually required general anesthesia (88.2% vs 54.9%, p<0.001) (table 2). Also on multivariate analysis with adjustments for baseline differences, MT in BAO was associated with higher odds of achieving TICI 3 (aOR=2.004, 95% CI 1.227 to 3.274, p=0.005) and successful recanalization \geq TICI 2b (aOR=2.740, CI 1.145 to 6.554).

Safety

Safety endpoints, including sICH, systemic bleeding, craniectomy, and complication rates, were less frequent in patients with BAO than in patients with ACLVO (table 2), but the difference did not reach statistical significance. Also on multivariate analysis, MT in BAO as compared with ACLVO was not associated with either sICH (aOR=0.773, 95% CI 0.228 to 2.618) or mortality (aOR=1.612, 95% CI 0.891 to 2.918). However, patients with MT in BAO more often had postinterventional

Characteristics	BAO (n=165)	ACLVO (intracranial carotid artery, carotid-T, M1) (n=1574)	P value
Clinical items			
Age (years)	70 (59–80), N=165	73 (61–82), N=1574	0.019
Transfer from another hospital	80/165 (48.5%)	561/1572 (35.7%)	0.002
Sex (female)	69/165 (41.8%)	810/1574 (51.5%)	0.022
NIHSS on admission	18 (8–30), N=155	17 (12–20), N=1558	0.046
Prestroke independence	149/164 (90.9%)	1446/1556 (92.9%)	0.341
Blood pressure systolic (mm Hg)	148 (SD 31), N=86	150 (SD 28), N=1141	0.413
Blood pressure diastolic (mm Hg)	78 (SD 16), N=86	82 (SD 20), N=1139	0.097
Admission glucose (mmol/L)	7.5 (6.4–9.7), N=93	6.6 (5.8–8.1), N=1181	<0.001
Wake-up stroke	13/161 (8.1%)	123/1451 (8.5%)	1.000
TOAST etiology			0.008
Large artery	37/163 (22.7%)	205/1546 (13.3%)	
Cardioembolic	64/163 (39.3%)	733/1546 (47.4%)	
Other specific etiology	12/163 (7.4%)	101/1546 (6.5%)	
Unknown etiology	50/163 (30.7%)	507/1546 (32.8%)	
Medication			0.058
Antiplatelet			
Monotherapy	36/154 (23.4%)	430/1463 (29.4%)	
Dual therapy	0 (0%)	26/1463 (1.8%)	
Statin	32/142 (22.5%)	395/1300 (30.4%)	0.053
Anticoagulation			0.014
VKA	10/154 (6.5%)	180/1464 (12.3%)	
NOAC	2/154 (1.3%)	64/1464 (4.4%)	
Risk factors			
Diabetes	23/163 (14.1%)	271/1555 (17.4%)	0.326
Arterial hypertension	87/163 (53.4%)	1046/1553 (67.4%)	<0.001
Dyslipidemia	60/161 (37.3%)	788/1547 (50.9%)	0.001
Smoking	47/153 (30.7%)	428/1499 (28.6%)	0.574
Previous stroke	17/159 (10.7%)	209/1565 (13.4%)	0.389
Coronary artery disease	17/83 (20.5%)	235/1131 (20.8%)	1.000
Type of imaging			0.010
MRI	42/165 (25.5%)	994/1544 (64.4%)	
СТ	123/165 (74.5%)	550/1544 (35.6%)	
Treatment			
IVT use	71/165 (43.0%)	779/1574 (49.5%)	0.120
Time from onset of symptoms to IVT needle (min)	165 (113–210), N=39/71	120 (83–165), N=437/779	0.003
Time from onset of symptoms to admission in stroke center (min)	228 (IQR 121-369), N=132	143 (71–245), N=1380	<0.001
Time from onset of symptoms to groin puncture (min)	300 (IQR 211-480), N=133	225 (165–315), N=1346	<0.001

ACLVO, anterior circulation large-vessel occlusion; BAO, basilar artery occlusion; IVT, intravenous thrombolysis; NIHSS, National Institutes of Health Stroke Scale; NOAC, nonvitamin K antagonist oral anticoagulants; TOAST, Trial of ORG 10172 in Acute Stroke Treatment; VKA, vitamin K antagonist. Difference in numbers of observations and total numbers of patients in each group due to missing data items.

non-hemorrhagic worsening (aOR=2.797, CI 1.366 to 5.726) than in ACLVO.

Outcome

Patients with BAO had worse outcomes than patients with ACLVO, as shown by a lower rate of moderate outcome (mRS score 0–3, 46.1% vs 56.7%, p=0.013) and a higher mortality rate (36.2% vs 24.4%, p=0.002), although rates of excellent

functional outcome (mRS score 0–1, 25.0% vs 27.3%, p=0.631) and favorable outcome (mRS score 0–2, 36.2% vs 42.9%, p=0.120) were comparable in an unadjusted analysis (online supplementary table 2).

Baseline differences of patients with favorable outcome (mRS score 0–2) and unfavorable outcome (mRS score 3–6) can be found in online supplement table 3 and favoured neither BAO nor ACLVO. After adjusting for baseline differences, there was

Table 2	Safety and efficacy data comparing patients with basilar artery occlusion and patients with large vessel occlusion in the anterior
circulatio	n

	MT of BAO (n=165)	MT of ACLVO (n=1574)	P value	
Efficacy				
mTICI 3	102/165 (61.8%)	706/1571 (44.9%)	<0.001	
mTICI ≥2b	149/165 (90.3%)	1299/1571 (82.7%)	0.011	
Time from groin puncture to recanalization (min)	45 (30–81), N=153	47 (30–75), N=1471	0.824	
General anesthesia	142/161 (88.2%)	792/1442 (54.9%)	<0.001	
Additional intra-arterial thrombolytic agents	18/160 (11.3%)	112/1443 (7.8%)	0.127	
Maneuver count	1 (IQR 1–2), N=101	2 (1–3), N=1106	0.002	
Intracranial stenting	28/165 (17.0%)	36/1572 (2.3%)	<0.001	
Extracranial stenting	13/165 (7.9%)	182/1572 (11.6%)	0.194	
Safety				
sICH ECASS II definition	8/165 (4.8%)	98/1562 (6.3%)	0.608	
Systemic bleeding	1/39 (2.6%)	21/637 (3.3%)	1.000	
Craniectomy	4/161 (2.5%)	51/1448 (3.5%)	0.649	
Any interventional complication	18/165 (10.9%)	198/1572 (12.6%)	0.620	
Complications	 Two vasospasms Five dissections Three perforations Two emboli to new territory Six other 	 51 vasospasms 37 dissections 21 perforations 59 emboli to new territory 32 other 		

ACLVO, anterior circulation large vessel occlusion; BAO, basilar artery occlusion; MT, mechanical thrombectomy; mTICI, modified Thrombolysis in Cerebral Infarction; sICH, symptomatic intracranial hemorrhage according to the European Cooperative Acute Stroke Study II definition.Difference in numbers of observations and total numbers of patients in each group due to missing data items.

no significant difference in excellent (aOR=0.893, 95% CI 0.478 to 1.669), favorable (primary outcome, aOR=0.986, 95% CI 0.553 to 1.758, figure 1) and moderate functional outcome at 90 days (aOR=0.692, 95% CI 0.396 to 1.121).

In conclusion, patients with BAO had worse outcomes on univariate analysis, but after adjustment for baseline differences,

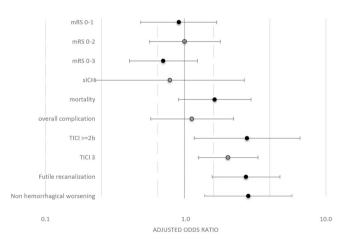


Figure 1 Comparison of outcomes of mechanical thrombectomy (MT) in posterior versus anterior circulation. Adjusted OR of MT in basilar artery occlusion versus anterior circulation large vessel occlusion for outcome parameters in the binary multivariable logistic regression analysis, adjusting for age, National Institutes of Health Stroke Scale score on admission, systolic blood pressure, prestroke independence, known onset, diabetes, arterial hypertension, smoking, previous stroke, imaging type, intravenous thrombolysis, and time from onset to admission. mRS, modified Rankin scale; sICH, symptomatic intracranial hemorrhage; TICI, Thrombolysis in Cerebral Infarction.

we could observe no significant differences of outcome of MT in BAO vs ACLVO, except an increased occurrence of FR.

FR was found more often in BAO than in ACLVO (aOR=2.146, CI 1.267 to 3.633). Factors associated with FR in univariate analysis were older age, higher stroke severity, coronary artery disease, higher maneuver count, and intracranial stenting (table 3). Prestroke dependence, lower posterior circulation Alberta Stroke Program Early CT Score (pcASPECTS), longer time from groin puncture to recanalization and administration of additional intra-arterial thrombolytic agents showed a trend towards association with FR.

On unadjusted analysis, better outcomes were observed in patients with successful recanalization of BAO (online supplement table 4) as compared with patients without successful recanalization. The relative benefit of achieving successful recanalization for various dichotomizations of the mRS scale, mortality, and sICH were comparable between patients presenting with recanalization of BAO and ACLVO according to multivariable binary logistic regression analysis, adjusting for prespecified confounders outlined in the methods section (figure 2). However, significantly higher rates of independence at 3 months were found only in the cohort of patients presenting with ACLVO.

In successfully recanalized BAO patients (TICI $\ge 2b$, n=149), excellent recanalization (TICI 3, n=102) approached significance to increase the odds of a favorable outcome (mRS 0–2) at day 90 (aOR=2.177, 95% CI 0.923 to 5.135) and decreased the odds of all-cause mortality at day 90 (aOR=0.408, 95% CI 0.177 to 0.937).

DISCUSSION

This registry-based retrospective analysis of patients with BAO treated with stent-retriever MT has the following main findings:

	Futile recanalization (n=70)	Non-futile recanalization (n=79)	P value
Clinical items			
Age (years)	74 (63–81), N=70	67 (57–79), N=79	0.025
Transfer from another hospital	34/70 (48.6%)	37/79 (46.8%)	0.870
Sex (female)	34/70 (48.6%)	30/79 (38.0%)	0.246
NIHSS score on admission	21 (12–36), N=65	12 (6–23), N=75	0.002
Prestroke independence	60/70 (85.7%)	75/79 (94.9%)	0.089
Blood pressure systolic (mm Hg)	154 (SD 32), N=41	144 (SD 28), N=39	0.170
Blood pressure diastolic (mm Hg)	78 (SD 17), N=41	78 (SD 15), N=39	0.840
Admission glucose (mmol/L)	7.6 (6.3–9.3), N=45	7.4 (6.4–9.8), N=41	0.959
Noticed symptom onset	50/70 (71.4%)	62/79 (78.5%)	0.347
Wake up	7/70 (10.0%)	5/77 (6.5%)	0.551
In-hospital stroke	2/70 (2.9%)	0/79	0.219
Vedication			0.417
Antiplatelet			
Monotherapy	17/67 (25.4%)	14/75 (18.7%)	
Dual therapy	0/67	0/75	
Statin	15/63 (23.8%)	12/69 (17.4%)	0.394
Anticoagulation			0.401
None	63/67 (94.0%)	69/75 (92.0%)	
VKA	4/67 (6.0%)	4/75 (5.3%)	
NOAC	0/67	2/75 (2.7%)	
Risk factors			
Diabetes	10/69 (14.5%)	9/79 (11.4%)	0.628
Arterial hypertension	36/69 (52.2%)	40/79 (50.6%)	0.870
Dyslipidemia	26/67 (38.8%)	26/79 (32.9%)	0.491
Smoking	16/65 (24.6%)	25/74 (33.8%)	0.267
Previous stroke	8/67 (11.9%)	8/79 (10.1%)	0.794
Coronary artery disease	11/35 (31.4%)	4/40 (10.0%)	0.040
TOAST etiology			0.286
long,			0.200
Large artery	13/69 (18.8%)	21/79 (26.6%)	
Cardioembolic	27/69 (39.1%)	33/79 (41.8%)	
Other specific etiology	4/69 (5.8%)	7/79 (8.9%)	
Unknown etiology	25/69 (36.2%)	18/79 (22.8%)	
Type of imaging			1.000
MRI	16/70 (22.9%)	19/79 (24.1%)	
СТ	54/70 (77.1%)	60/79 (75.9%)	
pcASPECTS	7 (5–9), N=25	8 (7–9), N=24	0.076
Treatment			
IVT use	29/70 (41.4%)	33/79 (41.8%)	1.000
Time from onset of symptoms to IVT needle (min)	165 (117–225), N=21/29	178 (109–228), N=16/33	0.914
Time from onset of symptoms to admission (min)	240 (122–446), N=55	222 (112–335), N=67	0.231
Time from onset of symptoms to groin puncture (min)	305 (211–539), N=55	300 (200–450), N=67	0.365
Time groin to recanalization (min)	47 (34–86), N=67	41 (28–68), N=76	0.096
General anesthesia	61/70 (87.1%)	68/77 (88.3%)	1.000
Additional intra-arterial thrombolytic agents	11/70 (15.7%)	4/76 (5.3%)	0.055
Balloon-guiding catheter	2/70 (2.9%)	7/76 (9.2%)	0.169
Intracranial stent	17/70 (24.3%)	7/79 (8.9%)	0.014
Extracranial stent	3/70 (4.3%)	8/79 (10.1%)	0.219

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Ischemic Stroke

Table 3 Continued					
	Futile recanalization (n=70)	Non-futile recanalization (n=79)	P value		
Any interventional complication	7/70 (10.0%)	6/79 (7.6%)	0.773		
Maneuver	1 (1–2), N=45	1 (1–2), N=50	0.016*		
sICH ECASS II	2/70 (2.9%)	2/79 (2.5%)	1.000		

*Maneuver count higher in futile recanalization.

IVT, intravenous thrombolysis; NIHSS, National Institutes of Health Stroke Scale; NOAC, non-vitamin K antagonist oral anticoagulants; pcASPECTS, posterior circulation Alberta Stroke Program Early CT Score; sICH, symptomatic intracranial hemorrhage according to the European Cooperative Acute Stroke Study II definition; TOAST, Trial of ORG 10172 in Acute Stroke Treatment; VKA, vitamin K antagonist. Difference in numbers of observations and total numbers of patients in each group due to missing data items.

(1) In comparison with patients presenting with ACLVO, interventional procedures in patients with BAO were more often technically effective and equally safe as regards mortality and sICH. (2) FR occurred more frequently in BAO, and predictors of FR included age, stroke severity, maneuver count, and intracranial stenting. (3) Patients with MT in BAO had worse functional outcome than in ACLVO on univariate analysis, but after multivariable adjustment, this difference was non-significant. (4) Successful recanalization in patients with BAO had similar associations with lower mortality, lower rates of sICH, and functional outcome as seen in ACLVO.

The recanalization rate of 90.3% in BAO from our study is in line with recent MT observational studies,²⁷ although it is slightly higher than in other systematic reviews, which included earlier studies^{10 17} and recent registries.^{15 28} Advances in MT technique, operator adjudication,²⁹ use of TICI score in PCLVO³⁰ in our registry, and different patient selection probably best explain the differences observed.

The rate of complications in patients with BAO was 10.9%, which was similar to rates previously reported^{31 32} and matched the frequency of complications seen in patients presenting with ACLVO.³¹ Serious complications observed were three artery perforations with one associated death. Patients with unsuccessful recanalization had a higher complication rate (31.3% vs 8.7%, p=0.018).

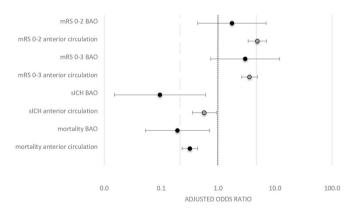


Figure 2 Effect of successful reperfusion on outcomes stratified for posterior versus anterior circulation. Adjusted OR of achieving successful recanalization (modified Thrombolysis in Cerebral Infarction $\geq 2b/3$) for outcome parameters stratified for patients with basilar artery occlusion versus anterior circulation. Notice the broad CIs in the posterior circulation due to the small number of patients in whom recanalization could not be achieved. Numbers indicate the p value for interaction of IV thrombolysis and successful recanalization in the binary multivariable logistic regression analysis, including age, sex, National Institutes of Health Stroke Scale score on admission, and IV thrombolysis. BAO, basilar artery occlusion; mRS, modified Rankin scale; sICH, symptomatic intracranial hemorrhage.

Irrespective of treatment modalities, patients with BAO more often have a deleterious course than their counterparts presenting with ACLVO.^{2 33} The mortality rate of 36.2% is in line with the published data.¹⁷ In accordance with previous data, we found a higher chance for FR in BAO than in ACLVO (47% vs 34%, p=0.002).¹⁵⁻¹⁸ Regarding factors associated with FR, our data confirm previously reported preinterventional factors, including age, baseline NIHSS score and lower pcASPECTS. Contrary to the literature, cardioembolic stroke subtype was no predictor of FR in our study. Importantly, we could identify interventional predictors of FR, mainly higher maneuver count and intracranial stenting. After adjusting for confounders, only age, baseline NIHSS score, and intracranial stenting remained significant prognostic factors for FR in our dataset. Together with a lower mortality rate in successfully recanalized patients, the observation of increased FR suggests that by MT, death was avoided but long-term dependence created. Those finding should be included in the discussion to treat or withhold treatment according to patient preferences.

Corroborating our observations about the beneficial effect of successful recanalization, the meta-analyses by Kumar et al also reported that mortality was bisected in patients in whom successful recanalization (mTICI 2b or mTICI 3) was achieved.¹⁰ Although partially present on unadjusted analysis, the higher rates of moderate, favorable, and excellent functional outcome were not statistically significant after adjustment for confounders when recanalization was achieved in BAO, although this could be shown by others.^{14 28 34} The small number of patients in whom recanalization could not be achieved was associated with broad confidence intervals, which probably explains the lack of statistical significance. More importantly, the effect of achieving successful recanalization with respect to mortality rates and sICH was independent of IVT administration. Sensitivity analysis disclosed no interaction between IVT and recanalization success.

It is noteworthy that excellent recanalization (TICI 3) was more beneficial for all-cause mortality at day 90 compared with successful recanalization (TICI \geq 2b), which could be also shown in ACLVO.^{35–37} Therefore, also in BAO, efforts should be made to improve the quality of recanalization by preventing or treating distal emboli.³⁸

Kumar *et al* concluded that the significant treatment effect of recanalization, confirmed by observational data and meta-analyses in this severe disease, would make a randomized controlled trial (RCT) to compare MT and the best medication-based treatment unethical.¹⁰ However, at least two such RCTs are currently being conducted.^{39 40} In our opinion, the most relevant question is not whether or not to treat patients with BAO with MT, but rather in which patients recanalization would be futile or cause long-term dependence. Therefore, RCTs comparing patient selection strategies with advanced imaging techniques and concomitant medical therapy are urgently warranted. Possible predictors

for good outcome including stroke risk factors, stroke severity (NIHSS score), age, respiratory stability, thrombus length, atherosclerotic stenosis versus embolic occlusion, and collateral status have been published.^{14 41-52} They should be combined in a study protocol of patient selection in BAO to minimize the incidence of futile recanalization.

Limitations

This is a single-arm multicenter retrospective registry, which has associated limitations. No comparison with patients treated with medical management only was performed, and patient selection for MT was center-specific. Several factors, including final mTICI score and initial pcASPECTS, were not adjudicated by a core laboratory and the exact location of BAO (proximal, middle, distal portion) was unknown. Data quality was incomplete for important predictors of outcome, such as collateral quality and pcASPECTS. Subgroup analyses were generally confined to small cohorts, which introduces a large uncertainty of the presented effects, as indicated by relatively wide confidence intervals.

CONCLUSION

In selected patients treated with MT, similar outcomes can be achieved in BAO and ACLVO. Patients with BAO in whom recanalization could be achieved, had lower mortality and rate of sICH than those with unsuccessful recanalization, without effect heterogeneity in comparison with ACLVO. The aim of MT should be complete recanalization. RCTs comparing patient selection and interventional strategies to avoid futile interventions seem warranted.

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Competing interests UF is a consultant for Medtronic and Stryker and co-principal investigator of the SWIFT DIRECT trial (Medtronic). JG is a global principal investigator of STAR (Solitaire FR Thrombectomy for Acute

Revascularisation), clinical event commitee member of the PROMISE study (European Registry on the ACE Reperfusion Catheters and the Penumbra System in the Treatment of Acute Ischemic Stroke; Penumbra), and a principal investigator and consultant for the SWIFT DIRECT study (Medtronic) and receives Swiss National Science Foundation (SNSF) grants for magnetic resonance imaging in stroke. LPi serves as a consultant for Balt, Microvention, and Penumbra. JK has received travel grants from Pfizer and Stryker. MRi serves as a consultant for Medtronic, Stryker, Anaconda, Apta Targets, and Perflow Medical and as a speaker for Neuravi. PMi has received funding for speaker honoraria from Boehringer. He has served on scientific advisory boards also for Boehringer. He has received research grants from Bristol-Myers Squibb, Boehringer, and the Swiss Heart Foundation. VMP is a consultant for Stryker (SC for DAWN trial), Penumbra (SC for PROMISE study), BALT (proctorship of products unrelated to ischemic stroke), Phenox, Rapid Medical, Neurovasc and receives research a grant from Philips. All the other authors have nothing to disclose.

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REFERENCES

- Smith WS, Lev MH, English JD, et al. Significance of large vessel intracranial occlusion causing acute ischemic stroke and TIA. Stroke 2009;40:3834–40.
- 2 Mattle HP, Arnold M, Lindsberg PJ, *et al*. Basilar artery occlusion. *Lancet Neurol* 2011;10:1002–14.
- 3 Merwick Á, Werring D. Posterior circulation ischaemic stroke. *BMJ* 2014;348:g3175.
- 4 Schulz UG, Fischer U. Posterior circulation cerebrovascular syndromes: diagnosis and management. J Neurol Neurosurg Psychiatry 2017;88:45–53.
- 5 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;387:1723–31.
- 6 Powers WJ, Rabinstein AA, Ackerson T, et al. 2018 Guidelines for the early management of patients with acute ischemic stroke: a guideline for healthcare professionals from the American Heart Association/American Stroke Association. Stroke 2018;49:e46–110.
- 7 Sarikaya H, Arnold M, Engelter ST, *et al*. Outcomes of intravenous thrombolysis in posterior versus anterior circulation stroke. *Stroke* 2011;42:2498–502.
- 8 Schonewille WJ, Wijman CA, Michel P, et al. Treatment and outcomes of acute basilar artery occlusion in the Basilar Artery International Cooperation Study (BASICS): a prospective registry study. *Lancet Neurol* 2009;8:724–30.
- 9 Pagola J, Ribo M, Álvarez-Sabin J, et al. Thrombolysis in anterior versus posterior circulation strokes: timing of recanalization, ischemic tolerance, and other differences. J Neuroimaging 2011;21:108–12.
- Kumar G, Shahripour RB, Alexandrov AV. Recanalization of acute basilar artery occlusion improves outcomes: a meta-analysis. J Neurointerv Surg 2015;7:868–74.
- 11 Lindsberg PJ, Mattle HP. Therapy of basilar artery occlusion: a systematic analysis comparing intra-arterial and intravenous thrombolysis. *Stroke* 2006;37:922–8.
- 12 Lindsberg PJ, Soinne L, Tatlisumak T, *et al.* Long-term outcome after intravenous thrombolysis of basilar artery occlusion. *JAMA* 2004;292:1862.
- 13 Ritvonen J, Strbian D, Silvennoinen H, et al. Thrombolysis and adjunct anticoagulation in patients with acute basilar artery occlusion. Eur J Neurol 2019;26:128–35.
- 14 Bouslama M, Haussen DC, Aghaebrahim A, et al. Predictors of good outcome after endovascular therapy for vertebrobasilar occlusion stroke. Stroke 2017;48:3252–7.
- 15 Alonso de Leciñana M, Kawiorski MM, Ximénez-Carrillo Á, et al. Mechanical thrombectomy for basilar artery thrombosis: a comparison of outcomes with anterior circulation occlusions. J Neurointerv Surg 2017;9:1173–8.
- 16 Lindsberg PJ, Sairanen T, Nagel S, et al. Recanalization treatments in basilar artery occlusion—systematic analysis. Eur Stroke J 2016;1:41–50.
- 17 Gory B, Eldesouky I, Sivan-Hoffmann R, et al. Outcomes of stent retriever thrombectomy in basilar artery occlusion: an observational study and systematic review. J Neurol Neurosurg Psychiatry 2016;87:520–5.
- 18 Singer OC, Berkefeld J, NoÎte CH, et al. Mechanical recanalization in basilar artery occlusion: the ENDOSTROKE study. Ann Neurol 2015;77:415–24.

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- 19 Thiel A, Schmidt H, Prange H, *et al.* [Treatment of patients with thromboses of the basilar artery and locked-in syndrome. An ethical dilemma]. *Nervenarzt* 1997;68:653–8.
- 20 Strbian D, Sairanen T, Silvennoinen H, et al. Thrombolysis of basilar artery occlusion: impact of baseline ischemia and time. Ann Neurol 2013;73:688–94.
- 21 Nagel S, Herweh C, Köhrmann M, *et al.* MRI in patients with acute basilar artery occlusion DWI lesion scoring is an independent predictor of outcome. *Int J Stroke* 2012;7:282–8.
- 22 Mokin M, Sonig A, Sivakanthan S, *et al*. Clinical and procedural predictors of outcomes from the endovascular treatment of posterior circulation strokes. *Stroke* 2016;47:782–8.
- 23 Phan K, Phan S, Huo YR, et al. Outcomes of endovascular treatment of basilar artery occlusion in the stent retriever era: a systematic review and meta-analysis. J Neurointerv Surg 2016;8:1107–15.
- 24 Kaesmacher J, Chaloulos-lakovidis P, Panos L, et al. Clinical effect of successful reperfusion in patients presenting with NIHSS < 8: data from the BEYOND-SWIFT registry. J Neurol 2019;266:598–608.
- 25 Higashida RT, Furlan AJ, Roberts H, et al. Trial design and reporting standards for intraarterial cerebral thrombolysis for acute ischemic stroke. J Vasc Interv Radiol 2003;14:E1–31.
- 26 Mazya M V, Cooray C, Lees KR, *et al.* Minor stroke due to large artery occlusion. When is intravenous thrombolysis not enough? Results from the SITS International Stroke Thrombolysis Register. *Eur Stroke J* 2017;3:239698731774600.
- 27 Kang DH, Jung C, Yoon W, et al. Endovascular thrombectomy for acute basilar artery occlusion: a multicenter retrospective observational study. J Am Heart Assoc 2018;7:e009419.
- 28 Weber R, Minnerup J, Nordmeyer H, et al. Thrombectomy in posterior circulation stroke: differences in procedures and outcome compared to anterior circulation stroke in the prospective multicentre REVASK registry. *Eur J Neurol* 2019;26:299–305.
- 29 Zhang G, Treurniet KM, Jansen IGH, et al. Operator versus core lab adjudication of reperfusion after endovascular treatment of acute ischemic stroke. Stroke 2018;49:2376–82.
- 30 Jung C, Yoon W, Ahn SJ, et al. The revascularization scales dilemma: is it right to apply the treatment in cerebral ischemia scale in posterior circulation stroke? AJNR Am J Neuroradiol 2016;37:285–9.
- 31 Behme D, Gondecki L, Fiethen S, et al. Complications of mechanical thrombectomy for acute ischemic stroke-a retrospective single-center study of 176 consecutive cases. *Neuroradiology* 2014;56:467–76.
- 32 LII Y, Holmberg A, Mpotsaris A, et al. Posterior circulation occlusions may be associated with distal emboli during thrombectomy: factors for distal embolization and a review of the literature. *Clin Neuroradiol* 2018.
- 33 Sharma S, Mazya MV, Wahlgren N, et al. IV thrombolysis in very severe and severe ischemic stroke: results from the SITS-ISTR Registry. *Neurology* 2016;86:2115.
- 34 Haussen DC, Dharmadhikari SS, Snelling B, et al. Posterior communicating and vertebral artery configuration and outcome in endovascular treatment of acute basilar artery occlusion. J Neurointerv Surg 2015;7:864–7.
- 35 Kleine JF, Wunderlich S, Zimmer C, *et al.* Time to redefine success? TICI 3 versus TICI 2b recanalization in middle cerebral artery occlusion treated with thrombectomy. *J Neurointerv Surg* 2017;9:117–21.

- 36 Kaesmacher J, Dobrocky T, Heldner MR, et al. Systematic review and meta-analysis on outcome differences among patients with TICI2b versus TICI3 reperfusions: success revisited. J Neurol Neurosurg Psychiatry 2018;89:910–7.
- 37 Rizvi A, Seyedsaadat SM, Murad MH, et al. Redefining 'success': a systematic review and meta-analysis comparing outcomes between incomplete and complete revascularization. J Neurointerv Surg 2019;11:9–13.
- 38 Boeckh-Behrens T, Pree D, Lummel N, et al. Vertebral artery patency and thrombectomy in basilar artery occlusions. Stroke 2019;50:389–95.
- 39 Liu X, Xu G, Liu Y, et al. Acute basilar artery occlusion: endovascular interventions versus standard medical treatment (BEST) trial-design and protocol for a randomized, controlled, multicenter study. *Int J Stroke* 2017;12:779–85.
- 40 van der Hoeven EJ, Schonewille WJ, Vos JA, *et al.* The Basilar Artery International Cooperation Study (BASICS): study protocol for a randomised controlled trial. *Trials* 2013;14:200.
- 41 Alqadri S, Adil MM, Watanabe M, et al. Patterns of collateral formation in basilar artery steno-occlusive diseases. J Vasc Interv Neurol 2013;6:9–13.
- 42 Werner M, Lopez-Rueda A, Zarco F, *et al*. Mechanical thrombectomy in acute basilar artery occlusion: a safety and efficacy single centre study. *Interv Neuroradiol* 2016;22:310–7.
- 43 Yoon W, Kim SK, Heo TW, *et al*. Predictors of good outcome after stent-retriever thrombectomy in acute basilar artery occlusion. *Stroke* 2015;46:2972–5.
- 44 Vergouwen MD, Compter A, Tanne D, et al. Outcomes of basilar artery occlusion in patients aged 75 years or older in the Basilar Artery International Cooperation Study. J Neurol 2012;259:2341–6.
- 45 Brandt T, von Kummer R, Müller-Küppers M, et al. Thrombolytic therapy of acute basilar artery occlusion. Variables affecting recanalization and outcome. Stroke 1996;27:875–81.
- 46 Cross DT, Moran CJ, Akins PT, et al. Collateral circulation and outcome after basilar artery thrombolysis. AJNR Am J Neuroradiol 1998;19:1557-63.
- 47 Gilberti N, Gamba M, Premi E, *et al*. Endovascular mechanical thrombectomy in basilar artery occlusion: variables affecting recanalization and outcome. *J Neurol* 2016;263:707–13.
- 48 Goyal N, Tsivgoulis G, Nickele C, et al. Posterior circulation CT angiography collaterals predict outcome of endovascular acute ischemic stroke therapy for basilar artery occlusion. J Neurointerv Surg 2016;8:783–6.
- 49 Jung S, Mono ML, Fischer U, et al. Three-month and long-term outcomes and their predictors in acute basilar artery occlusion treated with intra-arterial thrombolysis. *Stroke* 2011;42:1946–51.
- 50 Kim YW, Hong JM, Park DG, et al. Effect of intracranial atherosclerotic disease on endovascular treatment for patients with acute vertebrobasilar occlusion. AJNR Am J Neuroradiol 2016;37:2072–8.
- 51 Mundiyanapurath S, Möhlenbruch M, Ringleb PA, et al. Posterior circulation acute stroke prognosis early computed tomography score using hypointense vessels on susceptibility weighted imaging independently predicts outcome in patients with basilar artery occlusion. PLoS One 2015;10:e0132587.
- 52 Raymond S, Rost NS, Schaefer PW, *et al*. Patient selection for mechanical thrombectomy in posterior circulation emergent large-vessel occlusion. *Interv Neuroradiol* 2018;24:309–16.