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Blood Pressure in the First 6 Hours Following Endovascular Treatment for Ischemic Stroke Is Associated With Outcome

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BACKGROUND AND PURPOSE: Optimal blood pressure (BP) management in the acute phase of ischemic stroke remains an unresolved issue. It is uncertain whether guidelines for BP management during and after intravenous alteplase can be extrapolated to endovascular treatment (EVT) for stroke due to large artery occlusion in the anterior circulation. We evaluated the associations between systolic BP (SBP) in the first 6 hours following EVT and functional outcome as well as symptomatic intracranial hemorrhage.

METHODS: Patients of 8 MR CLEAN (Multicenter Randomized Controlled Trial of Endovascular Treatment for Acute Ischemic Stroke in the Netherlands) Registry centers, with available data on SBP in the 6 hours following EVT, were analyzed. We evaluated maximum, minimum, and mean SBP. Study outcomes were functional outcome (modified Rankin Scale) at 90 days and symptomatic intracranial hemorrhage. We used multivariable ordinal and binary regression analysis to adjust for important prognostic factors and studied possible effect modification by successful reperfusion.

RESULTS: Post-EVT SBP data were available for 1161/1796 patients. Higher maximum SBP (per 10 mmHg increments) was associated with worse functional outcome (adjusted common odds ratio, 0.93 [95% CI, 0.88–0.98]) and a higher rate of symptomatic intracranial hemorrhage (adjusted odds ratio, 1.17 [95% CI, 1.02–1.36]). The association between minimum SBP and functional outcome was nonlinear with an inflection point at 124 mmHg. Minimum SBP lower and higher than the inflection point were associated with worse functional outcomes (adjusted common odds ratio, 0.85 per 10 mmHg decrements [95% CI, 0.76–0.95] and adjusted common odds ratio, 0.81 per 10 mmHg increments [95% CI, 0.71–0.92]). No association between mean SBP and functional outcome was observed. Successful reperfusion did not modify the relation of SBP with any of the outcomes.

CONCLUSIONS: Maximum SBP in the first 6 hours following EVT is positively associated with worse functional outcome and an increased risk of symptomatic intracranial hemorrhage. Both lower and higher minimum SBP are associated with worse outcomes. A randomized trial to evaluate whether modifying post-intervention SBP results in better outcomes after EVT for ischemic stroke seems justified.

GRAPHIC ABSTRACT: An online [graphic abstract](#) is available for this article.

Key Words: blood pressure ■ cerebral hemorrhage ■ guideline ■ ischemic stroke ■ reperfusion

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Nonstandard Abbreviations and Acronyms

BP	blood pressure
EVT	endovascular treatment
MR ASAP	Multicentre Randomised trial of Acute Stroke treatment in the Ambulance With Nitroglycerin Patch
MR CLEAN	Multicenter Randomized Controlled Trial of Endovascular Treatment for Acute Ischemic Stroke in the Netherlands
SBP	systolic blood pressure
sICH	symptomatic intracranial hemorrhage

In the first 24 hours after stroke, BP is often increased, even after endovascular treatment (EVT), and it takes a few days to return to baseline levels.^{1,2} It has been demonstrated that admission BP is strongly associated with functional outcome after EVT.^{3–5} Since BP is an important factor affecting cerebral perfusion, it is likely that BP within the first hours following EVT has an impact on infarct size and thereby functional outcome.^{6,7} Two observational studies found an association between systolic BP (SBP) peaks in the 24 hours following stroke and increased risks of symptomatic intracranial hemorrhage (sICH) and functional dependency.^{2,8} However, these studies did not relate timing of BP measurement to the occurrence of sICH, so reverse causality could be present and the target BP level in the first few hours after EVT remains unclear. Yet, if BP is causally related to outcome, modification using medication might be a feasible strategy to improve functional outcomes. We aimed to evaluate the associations of SBP in the first 6 hours following EVT with functional outcome and the occurrence of sICH.

METHODS

Study Protocol and Data Availability

We used data from the MR CLEAN (Multicenter Randomized Controlled Trial of Endovascular Treatment for Acute Ischemic Stroke in the Netherlands) Registry, a prospective, multicenter, observational cohort, including all consecutive patients treated with EVT for acute ischemic stroke in the Netherlands between March 2014 and 2017. Detailed information on the description of variables and the methods of MR CLEAN Registry have been reported previously.⁹ Data cannot be made available, as no patient approval has been obtained for sharing coded data. However, R syntax and output files of the analyses will be made available on request.

Study Population

Patients were eligible for inclusion for this analysis if they had been treated in an MR CLEAN Registry center that was able to provide BP data of the first 6 hours after EVT. Individual

patients were included if they were 18 years or older; had a proximal intracranial occlusion in the anterior circulation (intracranial carotid artery/intracranial carotid artery terminus, middle cerebral artery [M1/M2], and anterior cerebral artery [A1/A2]) confirmed on computed tomography angiography; in whom groin puncture was possible within 6.5 hours after symptom onset; and had at least one available BP value within the first 6 hours following EVT.

BP Measures

We collected SBP values recorded between the end of the EVT procedure (defined as time of reperfusion or last contrast bolus) and 24 hours after EVT or until discharge from the intervention center. To limit the risk of confounding by indication based on missing BP data due to early transfer of patients in good condition, we restricted our primary analysis to the first 6 hours following EVT. The predefined BP measures of interest included (1) maximum SBP (reflecting peak in BP course), (2) minimum SBP (reflecting drops in BP), and (3) mean SBP. If >1 SBP measurement was available, maximum and minimum SBP were calculated based on the average of the 2 highest or lowest SBP values in the 6 hours following EVT, to limit the risk of measurement error. When only one SBP value was available, there was no difference between maximum, minimum and mean SBP. Additionally, we performed a sensitivity analysis to evaluate the association between the predefined BP measures in the first 24 hours following EVT and outcomes. Since the majority of sICH and extracranial hemorrhage occurs within 24 hours following EVT, we did not evaluate the association between BP and these outcomes to avoid reverse causality. Details on BP protocols of the included centers are described in Table I in the [Data Supplement](#).

Outcome Measures

The primary outcome measure was functional outcome according to the modified Rankin Scale, which is a 7-point scale ranging from 0 no symptoms to 6 death, assessed at 90 days after EVT.¹⁰ Secondary outcome measures included functional independence (modified Rankin Scale score ≤ 2), mortality within 90 days after EVT, National Institutes of Health Stroke Scale score indicating neurological deficit at 24 to 48 hours after EVT, extracranial hemorrhage (requiring surgery or blood transfusion), and new ischemic stroke (new neurological deficit confirmed with imaging) within 90 days from stroke onset. Furthermore, any occurrence of sICH (neurological deterioration of ≥ 4 points on the National Institutes of Health Stroke Scale and a compatible hemorrhage on noncontrast computed tomography assessed by an independent core laboratory according to the Heidelberg criteria) was included as a secondary outcome measure.^{11,12}

Statistical Analysis

Baseline characteristics of the study population are tabulated by 3 subgroups according to maximum SBP tertiles. Continuous variables are expressed as means (SD) or medians (interquartile ranges), where applicable. Categorical variables are expressed as numbers of patients and percentages.

We evaluated the linearity of the associations between the postprocedural SBP parameters and outcomes by comparing

model fit of a regression model with a linear SBP term to a regression model with a SBP term with a restricted cubic spline transformation with 3 knots. We performed multivariable ordinal logistic regression, binary logistic regression or linear regression analyses, as appropriate with adjustment for the following potential confounders: age, sex, National Institutes of Health Stroke Scale score on admission, pre-stroke modified Rankin Scale score, medical history of hypertension, stroke, diabetes, atrial fibrillation, myocardial infarction, treatment with intravenous thrombolysis, SBP on admission, location of occlusion, Alberta Stroke Program Early CT Score on noncontrast computed tomography,¹³ collateral score on computed tomography angiography according to a 4-point scale (0=absent collaterals [0% filling of the vascular territory downstream of the occlusion], 1=poor collaterals [$>0\%$ and $\leq 50\%$ filling], 2=moderate collaterals [$>50\%$ and $<100\%$ filling], and 3=excellent collaterals [100% filling]),¹⁴ the use of general anesthesia during EVT, time from stroke onset to reperfusion or last contrast bolus, extended Thrombolysis in Cerebral Infarction score at the end of the EVT procedure,¹⁵ number of BP measurements in the 6 hours following EVT, and intervention center. For the outcome sICH, we aimed to reduce the possibility that results were hampered by reverse causality (ie, BP measurements collected during or after occurrence of sICH) by excluding patients in whom sICH occurred within 6 hours following EVT. The associations of BP parameters with outcomes were presented per 10 mm Hg change in BP.

We assessed whether the relation between postprocedural BP and outcomes was modified by the extent of reperfusion. We fitted a similar multivariable regression model as described above including an interaction term for SBP parameter*successful reperfusion, a dichotomized term for extent of reperfusion (unsuccessful, extended Thrombolysis in Cerebral Infarction score $<2B$ versus successful, extended Thrombolysis in Cerebral Infarction score $\geq 2B$).¹⁵ For all regression analyses, missing data were imputed using multiple imputations by chained equations based on relevant covariates and outcomes.¹⁶ All analyses were performed using R software (Version 3.6.1, R foundation for Statistical Computing, Vienna, Austria) with the packages: *tableone*, *mice*, *Hmisc*, *ggplot2*, and *rms*.

Medical Ethics Committee Statement

The medical ethics committee of the Erasmus University MC, Rotterdam, the Netherlands, evaluated the study protocol of the MR CLEAN Registry and granted permission to perform the study as a registry (MEC-2014-235).

RESULTS

Study Population

Of 1796 patients treated with EVT during the study period in the 8 participating centers, 1161 (65%) were included in the current analysis (Figure 1). The median available number of SBP measurements in the first 6 hours following EVT was 7 (interquartile range, 4–11). For 86/1161 patients only one SBP value in the first 6 hours was available. The mean SBP in the first 6 hours following EVT was 150 mm Hg (SD 25). Baseline characteristics of the study population are shown according to maximum SBP tertiles (Table 1). Patients with a higher maximum SBP in the first 6 hours following EVT were on average older and were more likely to have a history of atrial fibrillation, diabetes, hypertension, distal occlusion, and poorer collateral scores.

Association of Maximum SBP With Outcomes

The association between maximum SBP and functional outcome at 90 days (shift towards better modified Rankin Scale score) was linear (Figure 2A, likelihood ratio test $P=0.14$ for maximum SBP). Patients with higher maximum SBP in the 6 hours following EVT were more likely to have worse functional outcomes compared with patients with lower maximum SBP (adjusted common odds ratio [OR], 0.93 per 10 mm Hg [95% CI, 0.88–0.98], Table 2). Higher maximum SBP was associated with a larger neurological deficit (measured with the National Institutes of Health Stroke Scale) at 24 to 48 hours after EVT (β 0.31 [95% CI, 0.14–0.49]), increased risk of sICH (adjusted OR, 1.17 [95% CI, 1.02–1.36]), but not with an increased

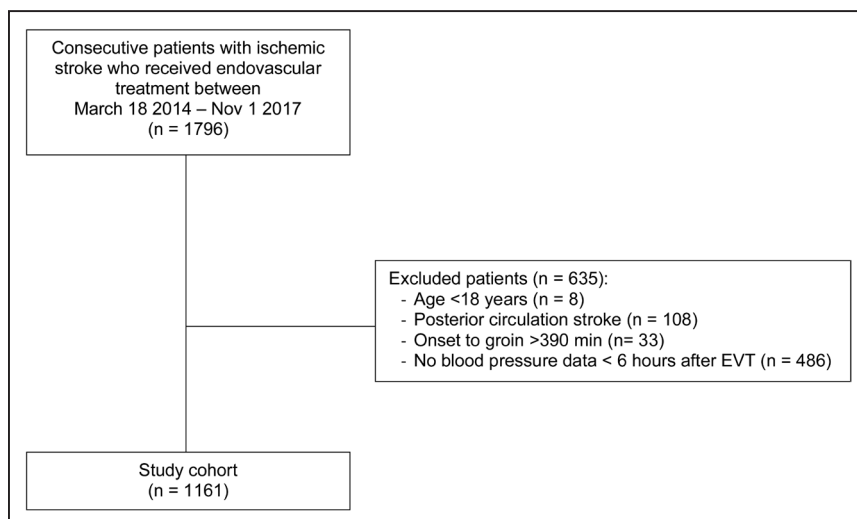


Figure 1. Flowchart of patient inclusion.

EVT indicates endovascular treatment.

Table 1. Baseline Characteristics of All Patients Shown According to Tertiles of Maximum SBP During First 6 Hours Following EVT

	Maximum SBP, <140 mm Hg (n=364)	Maximum SBP, 140–170 mm Hg (n=466)	Maximum SBP, >170 mm Hg (n=331)	Missing
Patient characteristics				
Age, mean (SD)	65 (15)	71 (13)	74 (12)	
Male sex, n (%)	190 (52)	242 (52)	166 (50)	
NIHSS, median [IQR]	16 [11–19]	15 [10–19]	17 [12–20]	2/3/3
Left hemisphere, n (%)	200 (55)	228 (49)	174 (53)	0/0/1
SBP, mean (SD)	134 (20)	150 (22)	162 (24)	10/3/3
DBP, mean (SD)	77 (14)	83 (16)	87 (16)	10/5/5
IVT, n (%)	275 (76)	368 (79)	251 (76)	2/1/0
Medical history, n (%)				
Previous stroke	54 (15)	84 (18)	62 (19)	2/5/5
Atrial fibrillation	87 (24)	118 (26)	91 (28)	3/7/8
Hypertension	151 (42)	237 (53)	199 (61)	7/20/7
Diabetes	44 (12)	79 (17)	66 (20)	2/4/5
Myocardial infarction	50 (14)	67 (15)	52 (16)	3/9/11
Peripheral arterial disease	49 (14)	41 (8.9)	31 (9.7)	4/7/11
Prestroke mRS score				8/8/12
0	261 (73)	332 (73)	220 (69)	
1	37 (10)	51 (11)	43 (14)	
2	18 (5.1)	28 (6.1)	24 (7.5)	
≥3	40 (11)	47 (10)	32 (10)	
Medication, n (%)				
Antihypertensive	170 (48)	249 (55)	199 (61)	6/14/6
Statin	133 (37)	170 (37)	118 (37)	7/12/12
Antiplatelet	111 (31)	133 (29)	114 (35)	4/7/9
DOAC	19 (5.3)	15 (3.3)	8 (2.4)	4/7/4
Coumarin	11 (3.1)	11 (2.4)	4 (1.2)	5/8/7
Imaging, n (%)				
Occluded segment				9/16/15
ICA	14 (3.9)	23 (5.1)	17 (5.4)	
ICA-T	69 (19)	79 (18)	69 (22)	
M1	236 (67)	275 (61)	172 (54)	
M2	32 (9.0)	71 (16)	57 (18)	
Other*	4 (1.1)	2 (0.4)	1 (0.3)	
ASPECTS subgroups				
0–4	16 (4.5)	18 (4.0)	10 (3.2)	
5–7	67 (19)	72 (16)	67 (21)	
8–10	276 (77)	359 (80)	236 (75)	
Collateral score				
Absent	8 (2.3)	21 (4.8)	23 (7.4)	
Filling <50% of occluded area	122 (35)	148 (34)	145 (47)	
Filling ≥50% but less <100%	143 (41)	198 (45)	97 (31)	
Filling 100% of occluded area	76 (22)	74 (17)	45 (15)	
Workflow				
Transfers from primary stroke center, n (%)	208 (57)	308 (66)	213 (64)	1/0/0
Time from stroke onset to groin puncture, min, median [IQR]	190 [150–240]	190 [150–240]	192 [153–250]	

(Continued)

Table 1. Continued

	Maximum SBP, <140 mm Hg (n=364)	Maximum SBP, 140–170 mm Hg (n=466)	Maximum SBP, >170 mm Hg (n=331)	Missing
Procedure				
General anesthesia, n (%)	22 (6.5)	23 (5.3)	23 (7.4)	26/30/20
Duration procedure, min, median [IQR]	60 [38–86]	55 [36–80]	61 [42–80]	49/41/23
Reperfusion grade after intervention (eTICI), n (%)				8/15/13
0	38 (11)	67 (15)	56 (18)	
1	6 (1.7)	17 (3.8)	5 (1.6)	
2A	71 (20)	81 (18)	57 (18)	
2B	92 (26)	98 (22)	66 (21)	
2C	36 (10)	42 (9.3)	35 (11)	
3	113 (32)	146 (32)	99 (31)	

Continuous data are presented as mean (SD) for normal distributed data or as median [IQR] for skewed data. Categorical data are presented as numbers (percentage). SBP tertiles are for data inspection only, analysis is based on the full range of SBP measures. Tertiles of maximum SBP were rounded to tens. ASPECTS indicates Alberta Stroke Program Early CT Score; DBP, diastolic blood pressure; DOAC, direct oral anticoagulant; eTICI, extended Thrombolysis in Cerebral Infarction; EVT, endovascular treatment; ICA-(T), internal carotid artery (terminus); IQR, interquartile range; IVT, intravenous thrombolysis; M(segment), middle cerebral artery; mRS, modified Rankin Scale; n, number; NIHSS, National Institutes of Health Stroke Scale; and SBP, systolic blood pressure.

*A1/A2/M3 occlusion.

risk of death (adjusted OR, 1.02 [95% CI, 0.95–1.08], Table 2). In the sensitivity analysis of SBP measures during the first 24 hours, we observed a similar association between higher maximum SBP and worse functional outcome (adjusted common OR, 0.90 per 10 mm Hg [95% CI, 0.85–0.94], Table II in the [Data Supplement](#)).

Association of Minimum SBP With Outcomes

The association between minimum SBP and functional outcome was nonlinear (Figure 2B) based on multivariable model fit comparing a linear SBP term to a model allowing 3 knots for SBP (likelihood ratio test $P < 0.01$

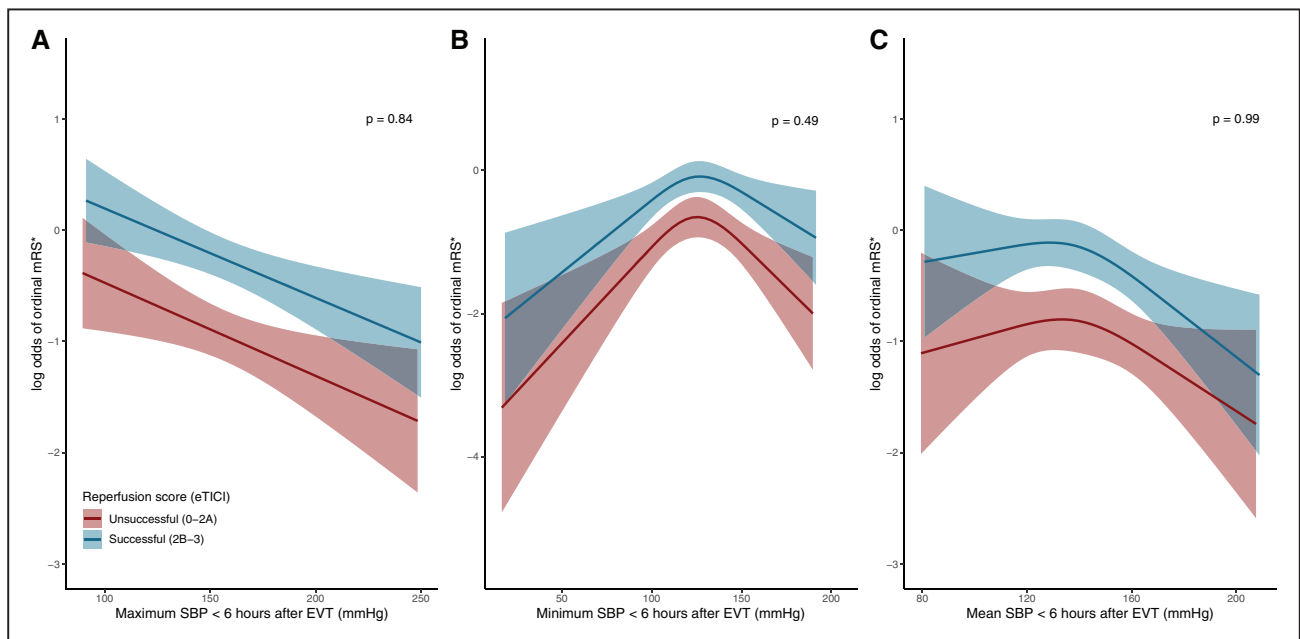


Figure 2. Relationship of systolic blood pressure (SBP) and shift toward better functional outcome.

The models are fitted with a linear function for maximum SBP and restricted cubic spline function with 3 knots for minimum SBP and mean SBP parameters. All models include the following variables: age, National Institutes of Health Stroke Scale (NIHSS) at baseline, Alberta Stroke Program Early CT Score (ASPECTS) at baseline, history of hypertension, time between stroke onset to reperfusion, and an interaction term for SBP parameter*reperfusion grade. The graphs depict the log odds for a shift towards better modified Rankin Scale (mRS) score (*ordinal mRS) with 95% CI, for each level of maximum SBP (A), minimum SBP (B), and mean SBP (C) in the first 6 h following endovascular treatment (EVT) for successful and unsuccessful reperfusion, with corresponding P value for interaction. The ranges of the x axes correspond to the lowest and highest SBP value in the data. eTICI indicates extended Thrombolysis in Cerebral Infarction.

Table 2. Associations Between Continuous Maximum SBP Within First 6 Hours Following EVT and Outcomes Shown per 10 mm Hg Increment in SBP

	n=1161	(c)OR/ β coefficient	a(c)OR/a β coefficient*
Primary outcome			
mRS at 90 d, median [IQR]	3 [1-6]	0.85 (0.85–0.92)	0.93 (0.88–0.98)
Secondary outcomes, clinical			
mRS score ≤ 2 at 90 d, n (%)	474 (44)	0.89 (0.85–0.92)	0.92 (0.86–0.98)
NIHSS 24–48 h, median [IQR]	9 [4-16]	0.51 (0.34–0.69)†	0.31 (0.14–0.49)†
Mortality at 90 d, n (%)	278 (26)	1.11 (1.05–1.16)	1.02 (0.95–1.08)
Symptomatic intracranial hemorrhage, n (%)	56 (4.8)	1.19 (1.07–1.33)‡	1.17 (1.02–1.36)‡
Extracranial hemorrhage, n (%)	23 (2.0)	1.05 (0.90–1.21)	1.00 (0.84–1.19)
New ischemic stroke, n (%)	19 (1.6)	0.89 (0.75–1.06)	0.90 (0.74–1.13)

a(c)OR indicates adjusted (common) odds ratio; ASPECTS, Alberta Stroke Program Early CT Score; a β , adjusted β coefficient; eTICI, extended Thrombolysis in Cerebral Infarction; EVT, endovascular treatment; IQR, interquartile range; mRS, modified Rankin Scale; n, number; NIHSS, National Institutes of Health Stroke Scale; SBP, systolic blood pressure; and sICH, symptomatic intracranial hemorrhage.

*Variables in the model: maximum SBP, age, sex, history of stroke, diabetes, hypertension, atrial fibrillation, myocardial infarction, prestroke mRS, intravenous thrombolysis, SBP on hospital admission, NIHSS at baseline, collateral score, ASPECTS at baseline, occlusion location, general anesthesia, eTICI after EVT, time from stroke onset to reperfusion, number of blood pressure measurements, and intervention center.

†Reported effect measure is β coefficient.

‡Patients with sICH ≤ 6 h following EVT were excluded (n=17).

for minimum SBP). Due to the nonlinearity of this association, we obtained effect estimates for lower minimum and higher minimum SBP separately (inflection point at around 124 mmHg). Minimum SBP below 124 mmHg and minimum SBP above 124 mmHg were both associated with worse functional outcome (adjusted common OR per 10 mmHg decrement 0.85 [95% CI, 0.76–0.95] for minimum SBP <124 mmHg and 0.81 per 10 mmHg increment [95% CI, 0.71–0.92] for minimum SBP ≥ 124 mmHg). Also, minimum SBP lower than 124 mmHg and minimum SBP higher than 124 mmHg were associated with higher mortality rates and a more frequent occurrence of extracranial hemorrhage. Minimum SBP higher than 124 mmHg was associated with more neurological deficit at 24 to 48 hours, which was not observed for lower minimum SBP (Table III in the [Data Supplement](#)).

Association of Mean SBP With Outcomes

The associations between mean SBP and functional outcome were also nonlinear (Figure 2C) based on multivariable model fit comparing a linear SBP term to a model allowing 3 knots for SBP (likelihood ratio test $P < 0.01$ for mean SBP). Therefore, we obtained effect estimates for lower mean SBP and higher mean SBP separately (inflection point at around 138 mmHg). Mean SBP below 138 mmHg was associated with higher likelihood of extracranial hemorrhage (adjusted OR, 1.66 per 10 mmHg decrement [95% CI, 1.07–2.51]). We did not observe an association between mean SBP higher than 138 mmHg and any of the outcomes (Table IV in the [Data Supplement](#)). The distribution of outcomes according to maximum, minimum and mean SBP tertiles is shown in Figure I and Table V in the [Data Supplement](#).

We did not find an interaction between extend of reperfusion and the relation of SBP with functional outcome (P values for interaction: maximum SBP=0.84; minimum SBP=0.49 and mean SBP=0.99, Figure 2) or any of the secondary outcomes (Figure II in the [Data Supplement](#)). We observed a decline in maximum SBP from baseline during the 6 hours following EVT for both reperfusion categories, with higher maximum SBPs among patients with unsuccessful reperfusion at the end of EVT procedure compared with patients with successful reperfusion (Figure III in the [Data Supplement](#)).

DISCUSSION

Increased maximum SBP in the first 6 hours following EVT was associated with worse functional outcome, a greater risk of sICH and more severe early neurological deficits. Minimum SBP lower and higher than the inflection point of 124 mmHg were associated with worse functional outcome. A mean SBP lower than 138 mmHg was associated with an increased risk of extracranial hemorrhage. None of the associations between BP and outcomes were modified by successful reperfusion at the end of the EVT procedure.

Our results are in line with previous studies reporting that higher maximum SBPs in the 24 hours following EVT are associated with worse clinical outcomes.^{28,17–19} The explanation for the worse outcome observed in patients with higher maximum SBP is likely to be multifactorial, including disruption of the blood-brain barrier, hemorrhagic transformation, elevated serum catecholamine levels, and larger infarcts.²⁰ The association between higher BP and worse outcomes following EVT has been observed up to 3 days after treatment, stressing the

importance of patient monitoring and support following EVT.²¹ In contrast with our findings, no association between maximum SBP after EVT and risk of sICH was observed in a subgroup analysis of a recent meta-analysis including 791 patients.¹⁹

We observed a nonlinear association between minimum SBP and functional outcome, with an inflection point at 124 mmHg during the first 6 hours following EVT. Previous studies evaluating minimum SBP did not find an association with functional outcome. However, these studies were small, no test for nonlinearity was performed, and functional outcome was assessed dichotomously.^{18,22} Only one other study reported that an increase in minimum SBP was associated with an increased likelihood of functional independence.¹⁷ Low SBP in the (sub)acute phase of ischemic stroke might be associated with impaired cerebral perfusion, infarct expansion, or complications like impending sepsis.^{22,23}

We observed a small decrease of maximum SBP following EVT in patients with successful compared to unsuccessful reperfusion, similar to previous findings.¹ It has been hypothesized that optimal BP regime varies with the reperfusion status (ie, successful or unsuccessful). For example, higher SBP might be associated with hemorrhagic transformation given complete reperfusion.^{24,25} However, maintaining hypertension might be of benefit in patients with unsuccessful reperfusion to optimize collateral blood flow and maintain cerebral perfusion pressure.^{7,17,26} Several studies reported modification of the effect of BP on outcome by reperfusion status.^{18,22} However, in our large study cohort, we did not observe different associations between SBP and functional outcome for patients with successful and unsuccessful reperfusion, which was also observed by another cohort study.² This might partially be explained by the fact that high SBP is a marker of tissue damage rather than reperfusion success. Therefore, successful reperfusion should probably be regarded as a confounder of the association between BP and outcome and not only as an effect modifier.

Given the clear association between BP and outcome after EVT, the lack of evidence on optimal BP management, the variation in hemodynamic management among EVT centers, and the possibility of a modifiable effect of BP on outcome, a clinical trial seems justified.²⁷ Currently, the BEST-II trial (Blood Pressure After Endovascular Stroke Therapy-II; URL: <https://www.clinicaltrials.gov>; Unique identifier: NCT04116112) aims to evaluate the safety of lower SBP in patients treated with EVT in whom successful reperfusion is achieved. In this trial, patients will be randomly assigned to one of the following SBP targets: ≤ 180 , < 160 , and < 140 mmHg. Intravenous antihypertensive treatment will be started after reperfusion to maintain SBP below the assigned target for 24 hours.²⁸

Furthermore, the BP-TARGET trial (Blood Pressure Target in Acute Stroke to Reduce Hemorrhage After Endovascular Therapy; URL: <https://www.clinicaltrials.gov>; Unique

identifier: NCT03160677) aims to determine whether strict SBP control (intervention arm: SBP between 110 and 129 mmHg) versus standard SBP control (control arm: SBP between 130 and 185 mmHg) during 24 hours following EVT in patients with successful reperfusion will reduce the risk of any intracranial hemorrhage.^{29,30} Besides, the ongoing MR ASAP trial (Multicentre Randomised Trial of Acute Stroke Treatment in the Ambulance With Nitroglycerin Patch) aims to assess the effect of transdermal glyceryl trinitrate started within 3 hours of symptom onset in the prehospital setting on functional outcome in patients with ischemic stroke or intracerebral hemorrhage. This intervention is suggested to improve outcome after stroke by an increase in the intracranial collateral flow and a reduction of the BP.³¹ Although these further studies on hemodynamic management in stroke patients are warranted, one of the major challenges of hemodynamic management remains to extrapolate population-based data to determine the target BP for an individual stroke patient.

Limitations

Our study has several limitations. First, due to the retrospective observational design, results could have been confounded by variables not adjusted for in the analyses, so residual confounding might be present. Second, our observed associations do not prove causality between SBP and outcome measures. SBP could have been measured during the asymptomatic phase preceding sICH. Hence, definitive inferences on effects of SBP treatment are not possible. Furthermore, as we did not have data on individual SBP targets or information on administration of either a vasopressor or an antihypertensive agent after EVT, we do not know how well SBP was managed. Besides, as data on follow-up infarct volumes were not available systematically, we could not evaluate if patients with higher SBP were more likely to have larger infarcts.

Conclusions

Patients with higher maximum SBP in the 6 hours following EVT are more likely to have worse functional outcome or sICH compared with patients with lower maximum SBP. Lower as well as higher minimum SBP are associated with worse functional outcome. Randomized trials are needed to evaluate whether modifying SBP post-EVT improves outcome.

ARTICLE INFORMATION

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Supplemental Materials

Online Tables I–V

Online Figures I–III

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REFERENCES

- John S, Hazaa W, Uchino K, Hussain MS. Timeline of blood pressure changes after intra-arterial therapy for acute ischemic stroke based on recanalization status. *J Neurointerv Surg*. 2017;9:455–458. doi: 10.1136/neurintsurg-2016-012369
- Matusevicius M, Cooray C, Bottai M, Mazya M, Tsvigoulis G, Nunes AP, Moreira T, Ollikainen J, Tassi R, Strbian D, et al. Blood pressure after endovascular thrombectomy: modeling for outcomes based on recanalization status. *Stroke*. 2020;51:519–525. doi: 10.1161/STROKEAHA.119.026914
- Venema E, Mulder MJHL, Roozenbeek B, Broderick JP, Yeatts SD, Khatri P, Berkhemer OA, Emmer BJ, Roos YBWEM, Majoie CBLM, 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. doi: 10.1136/bmj.j1710
- Mulder M, Ergezen S, Lingsma HF, Berkhemer OA, Franssen PSS, Beumer D, van den Berg LA, Lycklama ANG, Emmer BJ, van der Worp HB, et al. Baseline blood pressure effect on the benefit and safety of intra-arterial treatment in mr clean (multicenter randomized clinical trial of endovascular treatment of acute ischemic stroke in the Netherlands). *Stroke*. 2017;48:1869–1876. doi: 10.1161/STROKEAHA.116.016225
- van den Berg SA, Uniken Venema SM, Mulder MJHL, Treurniet KM, Samuels N, Lingsma HF, Goldhoorn RB, Jansen IGH, Coutinho JM, Roozenbeek B, et al; MR CLEAN Registry Investigators. Admission blood pressure in relation to clinical outcomes and successful reperfusion after endovascular stroke treatment. *Stroke*. 2020;51:3205–3214. doi: 10.1161/STROKEAHA.120.029907
- Ding X, Xu C, Zhong W, Gong X, Zhou Y, Chen Z, Lou M. Association of maximal systolic blood pressure with poor outcome in patients with hyperattenuated lesions on immediate NCCT after mechanical thrombectomy. *J Neurointerv Surg*. 2020;12:127–131. doi: 10.1136/neurintsurg-2019-014846
- Oliveira-Filho J, Silva SC, Trabuco CC, Pedreira BB, Sousa EU, Bacellar A. Detrimental effect of blood pressure reduction in the first 24 hours of acute stroke onset. *Neurology*. 2003;61:1047–1051. doi: 10.1212/01.wnl.0000092498.75010.57
- Mistry EA, Mistry AM, Nakawah MO, Khattar NK, Fortuny EM, Cruz AS, Froehler MT, Chitale RV, James RF, Fusco MR, et al. Systolic blood pressure within 24 hours after thrombectomy for acute ischemic stroke correlates with outcome. *J Am Heart Assoc*. 2017;6:e006167. doi: 10.1161/JAHA.117.006167
- Jansen IGH, Mulder MJHL, Goldhoorn RB; MR CLEAN Registry investigators. Endovascular treatment for acute ischaemic stroke in routine clinical practice: prospective, observational cohort study (MR CLEAN Registry). *BMJ*. 2018;360:k949. doi: 10.1136/bmj.k949
- van Swieten JC, Koudstaal PJ, Visser MC, Schouten HJ, van Gijn J. Interobserver agreement for the assessment of handicap in stroke patients. *Stroke*. 1988;19:604–607. doi: 10.1161/01.str.19.5.604
- Brott T, Adams HP Jr, Olinger CP, Marler JR, Barsan WG, Biller J, Spilker J, Holleran R, Eberle R, Hertzberg V. Measurements of acute cerebral infarction: a clinical examination scale. *Stroke*. 1989;20:864–870. doi: 10.1161/01.str.20.7.864
- von Kummer R, Broderick JP, Campbell BC, Demchuk A, Goyal M, Hill MD, Treurniet KM, Majoie CB, Marquering HA, Mazya MV, et al. The Heidelberg bleeding classification: classification of bleeding events after ischemic stroke and reperfusion therapy. *Stroke*. 2015;46:2981–2986. doi: 10.1161/STROKEAHA.115.010049
- Barber PA, Demchuk AM, Zhang J, Buchan AM. Validity and reliability of a quantitative computed tomography score in predicting outcome of hyperacute stroke before thrombolytic therapy. ASPECTS Study Group. Alberta stroke programme early CT score. *Lancet*. 2000;355:1670–1674. doi: 10.1016/s0140-6736(00)02237-6
- Tan IY, Demchuk AM, Hopyan J, Zhang L, Gladstone D, Wong K, Martin M, Symons SP, Fox AJ, Aviv RI. 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. doi: 10.3174/ajnr.A1408
- Goyal M, Fargen KM, Turk AS, Mocco J, Liebeskind DS, Frei D, Demchuk AM. 2C or not 2C: defining an improved revascularization grading scale and the need for standardization of angiography outcomes in stroke trials. *J Neurointerv Surg*. 2014;6:83–86. doi: 10.1136/neurintsurg-2013-010665
- Donders AR, van der Heijden GJ, Stijnen T, Moons KG. Review: a gentle introduction to imputation of missing values. *J Clin Epidemiol*. 2006;59:1087–1091. doi: 10.1016/j.jclinepi.2006.01.014
- Goyal N, Tsvigoulis G, Pandhi A, Dillard K, Alsbrook D, Chang JJ, Krishnaiah B, Nickle C, Hoit D, Alsherbini K, et al. Blood pressure levels post mechanical thrombectomy and outcomes in non-recanalized large vessel occlusion patients. *J Neurointerv Surg*. 2018;10:925–931. doi: 10.1136/neurintsurg-2017-013581
- Goyal N, Tsvigoulis G, Pandhi A, Chang JJ, Dillard K, Ishfaq MF, Nearing K, Choudhri AF, Hoit D, Alexandrov AW, et al. Blood pressure levels post

mechanical thrombectomy and outcomes in large vessel occlusion strokes. *Neurology*. 2017;89:540–547. doi: 10.1212/WNL.0000000000004184

19. Malhotra K, Goyal N, Katsanos AH, Filippatou A, Mistry EA, Khatri P, Anadani M, Spiotta AM, Sandset EC, Sarraj A, et al. Association of blood pressure with outcomes in acute stroke thrombectomy. *Hypertension*. 2020;75:730–739. doi: 10.1161/HYPERTENSIONAHA.119.14230
20. Warach S, Latour LL. Evidence of reperfusion injury, exacerbated by thrombolytic therapy, in human focal brain ischemia using a novel imaging marker of early blood-brain barrier disruption. *Stroke*. 2004;35:2659–2661. doi: 10.1161/01.STR.0000144051.32131.09
21. McCarthy DJ, Ayodele M, Luther E, Sheinberg D, Bryant JP, Elwardany O, Kimball J, Starke RM. Prolonged heightened blood pressure following mechanical thrombectomy for acute stroke is associated with worse outcomes. *Neurocrit Care*. 2020;32:198–205. doi: 10.1007/s12028-019-00803-7
22. Martins AI, Sargento-Freitas J, Silva F, Jesus-Ribeiro J, Correia I, Gomes JP, Aguiar-Gonçalves M, Cardoso L, Machado C, Rodrigues B, et al. Recanalization modulates association between blood pressure and functional outcome in acute ischemic stroke. *Stroke*. 2016;47:1571–1576. doi: 10.1161/STROKEAHA.115.012544
23. Verschoof MA, Groot AE, Vermeij JD, Westendorp WF, van den Berg SA, Nederkoorn PJ, van de Beek D, Coutinho JM. Association between low blood pressure and clinical outcomes in patients with acute ischemic stroke. *Stroke*. 2020;51:338–341. doi: 10.1161/STROKEAHA.119.027336
24. Nour M, Scalzo F, Liebeskind DS. Ischemia-reperfusion injury in stroke. *Interv Neurol*. 2013;1:185–199. doi: 10.1159/000353125
25. Kim TJ, Park HK, Kim JM, Lee JS, Park SH, Jeong HB, Park KY, Rha JH, Yoon BW, Ko SB. Blood pressure variability and hemorrhagic transformation in patients with successful recanalization after endovascular recanalization therapy: a retrospective observational study. *Ann Neurol*. 2019;85:574–581. doi: 10.1002/ana.25434
26. Caplan LR, Hennerici M. Impaired clearance of emboli (washout) is an important link between hypoperfusion, embolism, and ischemic stroke. *Arch Neurol*. 1998;55:1475–1482. doi: 10.1001/archneur.55.11.1475
27. Mistry EA, Mayer SA, Khatri P. Blood pressure management after mechanical thrombectomy for acute ischemic stroke: a survey of the StrokeNet sites. *J Stroke Cerebrovasc Dis*. 2018;27:2474–2478. doi: 10.1016/j.jstrokecerebrovasdis.2018.05.003
28. Mistry AM. Blood Pressure After Endovascular Stroke Therapy-II (Best-II). 2019. <https://clinicaltrials.gov/ct2/show/NCT04116112>.
29. Mazighi M, Labreuche J, Richard S, Gory B, Lapergue B, Sibon I, Berge J, Olivot JM, Reiner P, Houdart E, et al. Blood pressure target in acute stroke to reduce Hemorrhage after endovascular therapy: The randomized BP TARGET study protocol. *Front Neurol*. 2020;11:480. doi: 10.3389/fneur.2020.00480
30. Mazighi M, Richard S, Lapergue B, Sibon I, Gory B, Berge J, Consoli A, Labreuche J, Olivot JM, Broderick J, et al; BP-TARGET investigators. Safety and efficacy of intensive blood pressure lowering after successful endovascular therapy in acute ischaemic stroke (BP-TARGET): a multicentre, open-label, randomised controlled trial. *Lancet Neurol*. 2021;20:265–274. doi: 10.1016/S1474-4422(20)30483-X
31. van den Berg SA, Dippel DWJ, Hofmeijer J, Franssen PSS, Caminada K, Siegers A, Kruijff ND, Kerkhoff H, de Leeuw FE, Nederkoorn PJ, et al; MR ASAP Investigators. Multicentre randomised trial of acute stroke treatment in the ambulance with a nitroglycerin patch (MR ASAP): study protocol for a randomised controlled trial. *Trials*. 2019;20:383. doi: 10.1186/s13063-019-3419-z