

Circle of Willis variation and outcome after intra-arterial treatment

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ABSTRACT

Background Intra-arterial treatment (IAT) improves outcomes in acute ischaemic stroke. Presence of collaterals increases likelihood of good outcome. We investigated whether variations in the circle of Willis (CoW) and contributing carotid arteries influence outcome in patients who had a stroke treated with IAT.

Methods CT angiography data on patients who had an acute stroke treated with IAT were retrospectively collected. CoW was regarded complete if the contralateral A1 segment, anterior communicating artery and ipsilateral posterior communicating artery were fully developed, and the P1 segment was visible. Carotid artery contribution was studied with a self-developed carotid artery score ranging from 0 to 2 depending on the number of arteries supplying the occluded side of the CoW. Good clinical outcome was defined as modified Rankin Score ≤ 2 and measured at discharge and 3 months. We calculated risk ratios for the relation between completeness of the CoW, carotid score and good outcome, and performed a trend analysis for good outcome according to the carotid score.

Results 126 patients were included for analysis. Patients with a complete and incomplete CoW had a comparable risk for good outcome at discharge and 3 months. A higher carotid score was associated with a higher likelihood of good clinical outcome (p for trend 0.24 at discharge and 0.05 at 3 months).

Conclusion In patients with acute ischaemic stroke treated with IAT, chances of good clinical outcome tended to improve with number of carotid arteries supplying the cerebral circulation. Completeness of the CoW was not related to clinical outcome.

INTRODUCTION

Stroke is one of the leading causes of death and disability. Several trials showed that intra-arterial treatment (IAT) in acute ischaemic stroke treatment leads to better outcomes.^{1–6} Angiographic success rate is much higher than clinical recovery rate. One of the factors involved in this discrepancy is the presence of collaterals.^{7 8} The presence of adequate collaterals hinders penumbral tissue loss in acute ischaemic stroke⁹ and hence influences outcome after stroke.

The cerebral collateral circulation may be divided into primary collateral circulation through the circle of Willis and secondary collateral circulation through the

WHAT IS ALREADY KNOWN ON THIS TOPIC

⇒ The presence of adequate collaterals aids penumbral tissue preservation in acute ischaemic stroke and hence influences outcome after stroke. Most studies measure collateralisation by visibility of leptomeningeal collaterals in the affected hemisphere as seen on CT angiography.

WHAT THIS STUDY ADDS

⇒ In this study, we investigated whether variations in collaterals, by means of the circle of Willis and number of carotid arteries supplying the cerebral circulation, are associated with clinical outcome in patients with acute ischaemic stroke treated with intra-arterial treatment.

HOW THIS STUDY MIGHT AFFECT RESEARCH, PRACTICE OR POLICY

⇒ This study further defines which patient is more likely to benefit from intra-arterial treatment.

leptomeningeal and ophthalmic arteries. Previous studies on patients with carotid artery occlusion or severe carotid stenosis suggest that an incomplete circle of Willis increases risk of ischaemic stroke.^{10–12} In addition, extent of collateralisation influences outcome in patients who had a stroke treated with IAT.^{3 13–15} However, most of these studies measure collateralisation by visibility of leptomeningeal collaterals in the affected hemisphere as seen on CT angiography (CTA).^{3 13 14 16} In this study, we investigated whether variations in collaterals, by means of the circle of Willis and number of carotid arteries supplying the cerebral circulation, are associated with clinical outcome in patients with acute ischaemic stroke treated with IAT.

METHODS

Patients

All patients with acute anterior ischaemic stroke, due to carotid T-top or middle cerebral artery (MCA) (M1) occlusion, who were treated with IAT in the period of

October 2002–October 2013 (start of MR CLEAN trial) in Haaglanden Medical Center, The Hague and Antonius Hospital Nieuwegein, were included in the study. Demographic and clinical data were recorded at baseline including age, sex, time of symptom onset, National Institutes of Health Stroke Scale (NIHSS), blood pressure and serum glucose on admission. For all patients, type of intra-arterial therapy (intra-arterial thrombolysis or mechanical thrombectomy including type of device used) was registered. Patients with atherosclerotic carotid bifurcation occlusions or carotid dissections were not included in the study. The decision to treat a patient with IAT was made for each patient individually and IAT was performed only after obtaining consent from the patient or his relatives. Secondary preventive treatment was initiated according to European guidelines.¹⁷

Radiological characteristics

For each treated patient, site of intracranial occlusion or stenosis on CTA, presence of extracranial internal carotid artery occlusion or stenosis, Clot Burden Score (CBS),¹⁸ Alberta Stroke Programme Early CT Score (ASPECTS),¹⁹ leptomeningeal collateral flow (Collateral Flow Grading, CFG),²⁰ degree of recanalisation after IAT (Thrombolysis In Cerebral Infarction (TICI) score²⁰) and anatomy of the circle of Willis were assessed. The circle of Willis was regarded complete if there was a fully developed and open anterior (A)1 segment contralateral to the acute occlusion, anterior communicating artery (ACoA), ipsilateral posterior communicating artery (PCoA) and visible posterior (P)1 segment.

To investigate the contribution of the carotid arteries, we developed a score to determine the effect of circle of Willis variation on cerebral perfusion (figures 1–3):

- ▶ **Carotid score 0:** no contribution of either carotid artery to the collateral pathway. An example is a carotid T-top occlusion with a hypoplastic A1 segment contralaterally precluding perfusion of the circle through the contralateral anterior circulation.
- ▶ **Carotid score 1:** the circle of Willis is perfused from one carotid artery only. This happens with:
 - An M1 occlusion with a normal ipsilateral A1 segment and a hypoplastic contralateral A1 segment.
 - An M1 occlusion with bilateral symmetrical A1 segments but without a visible ACoA precluding perfusion from the contralateral carotid artery.
 - A carotid T-top occlusion and a dominant contralateral A1 segment, thus feeding both anterior cerebral arteries through the contralateral internal carotid artery (ICA).
- ▶ **Carotid score 2:** perfusion from two carotid arteries:
 - An M1 occlusion with normal bilateral A1 segments and a visible ACoA.

Recanalisation was regarded successful if the TICI score at the end of the intra-arterial procedure was 2b or 3. Most patients underwent a CT scan 24 hours after treatment or after any clinical deterioration. All radiological

scores were assessed by an experienced neuroradiologist (GLaN) blinded for clinical outcome.

Clinical outcomes

Clinical outcome was retrospectively assessed with the modified Rankin Score (mRS)^{21 22} at discharge and after 3 months. Good outcome was defined as an mRS score of ≤ 2 . In addition, we recorded complications both during the intra-arterial procedure and during admission.

Statistical analysis

Descriptive statistics were used for baseline, radiological and treatment characteristics. We calculated risk ratios (RRs) and 95% CIs for the relation between the completeness of the circle of Willis (complete vs incomplete), the carotid score and good clinical outcome with Poisson regression.²³ Moreover, we performed a trend analysis for good clinical outcome according to the three classes of the carotid score by taking this characteristic as a continuous variable. In addition, we adjusted for the predefined clinical (NIHSS, age and time to IAT) and radiological characteristics (ASPECTS, CFG and CBS). Adjusted RRs were calculated in bivariable analyses.

RESULTS

In the period of October 2002–October 2013, 163 patients were treated with IAT in both centres. We were able to retrieve CTAs in 129 patients. Clinical outcome was missing in one patient and in two other patients, the CTA was of low quality and could therefore not be rated appropriately. Hence, 126 patients were included for analysis. Baseline characteristics were essentially the same for both the complete group of 163 patients treated with IAT and the 126 patients who were subsequently included in our analysis (table 1). Fifty-five per cent (n=69) of all patients were men and median age was 62 years; median NIHSS was 14.

Treatment

Overall, 89 patients (71%) were treated with intravenous thrombolysis and 122 patients (97%) were treated with IAT. In four patients (2%), the intra-arterial procedure failed. Most patients treated with IAT had mechanical thrombectomy (table 1).

Radiological parameters

The majority of patients (84%) had an ASPECTS of 7 or more. Successful recanalisation was achieved in 59 patients (49%). The median CFG was 3 (range 0–3) and the median CBS was 3 (range 1–8).

Clinical outcome and completeness of the circle of Willis

Thirty-four patients (27%) had a good clinical outcome at discharge and 53 patients (42%) had good clinical outcome at 3 months (table 2).

Patients with a complete circle of Willis and patients with incomplete circle of Willis had a comparable risk for good clinical outcome at discharge (RR 0.82; 95% CI 0.38

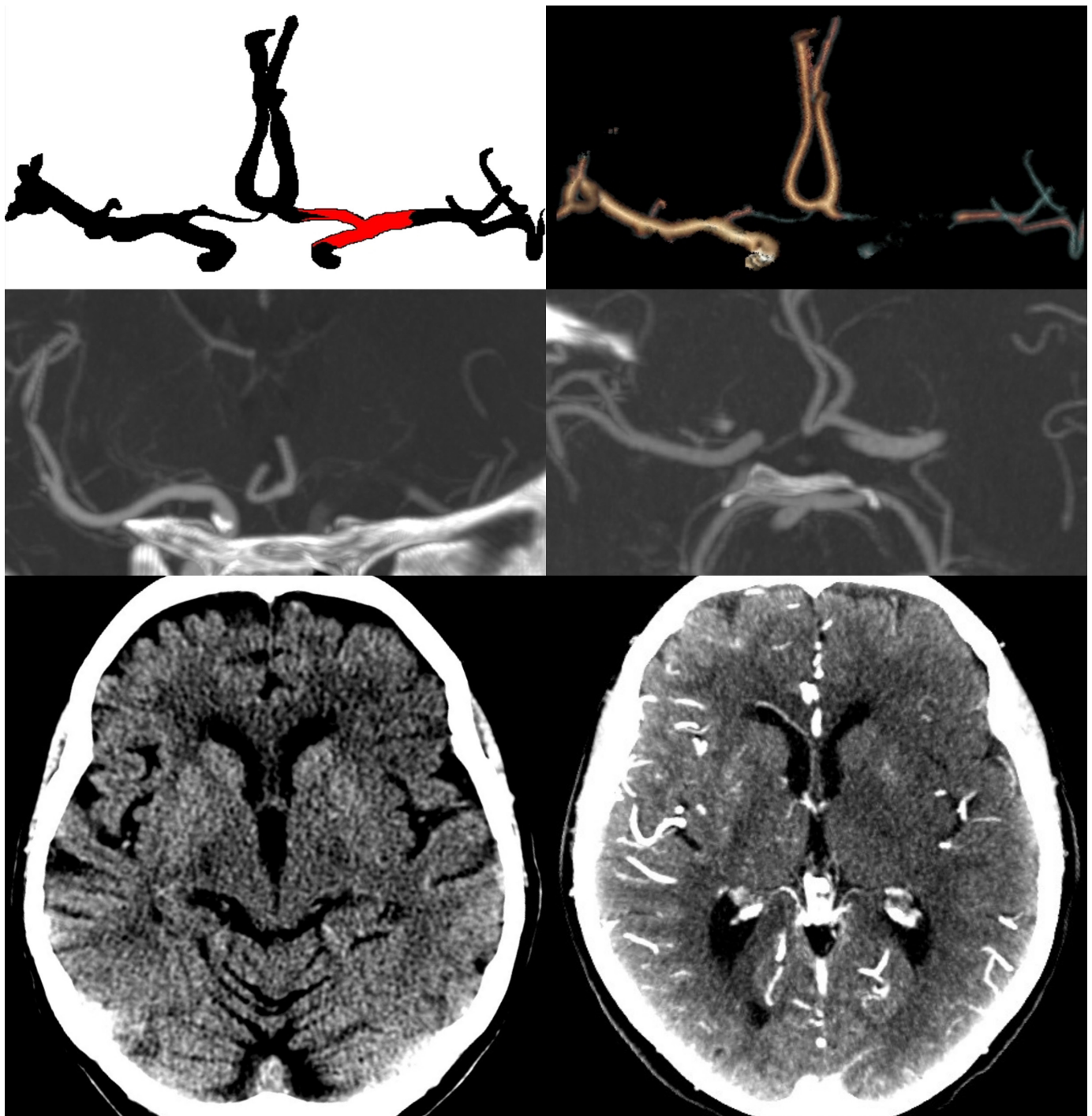


Figure 1 Carotid score 0, no contribution of either carotid artery to the collateral pathway, on non-enhanced CT, CT angiography and maximum intensity projection reconstructions.

to 1.78) and at 3 months (RR 1.01; 95% CI 0.61 to 1.67; table 2).

Clinical outcome and the carotid score

The majority of patients (n=78, 62%) had cerebral perfusion from both carotid arteries (carotid score 2). Of the patients with a complete circle of Willis (n=26, 21%), 22 patients had carotid score 2. Patients with only one carotid artery (carotid score 1) supplying the intracranial circulation tended to have better outcomes

compared with patients with carotid score 0 (RR at discharge 1.29; 95% CI 0.20 to 8.43 and RR 2.00; 95% CI 0.32 to 12.59 at 3 months). Further, patients with a carotid score of 2 tended to have more often a good clinical outcome at discharge (RR 1.85; 95% CI 0.30 to 11.39) and at 3 months (RR 2.92; 95% CI 0.48 to 17.75) compared with those with carotid score 0. In addition, compared with carotid score 1, patients with carotid score 2 also tended to have higher chances of good

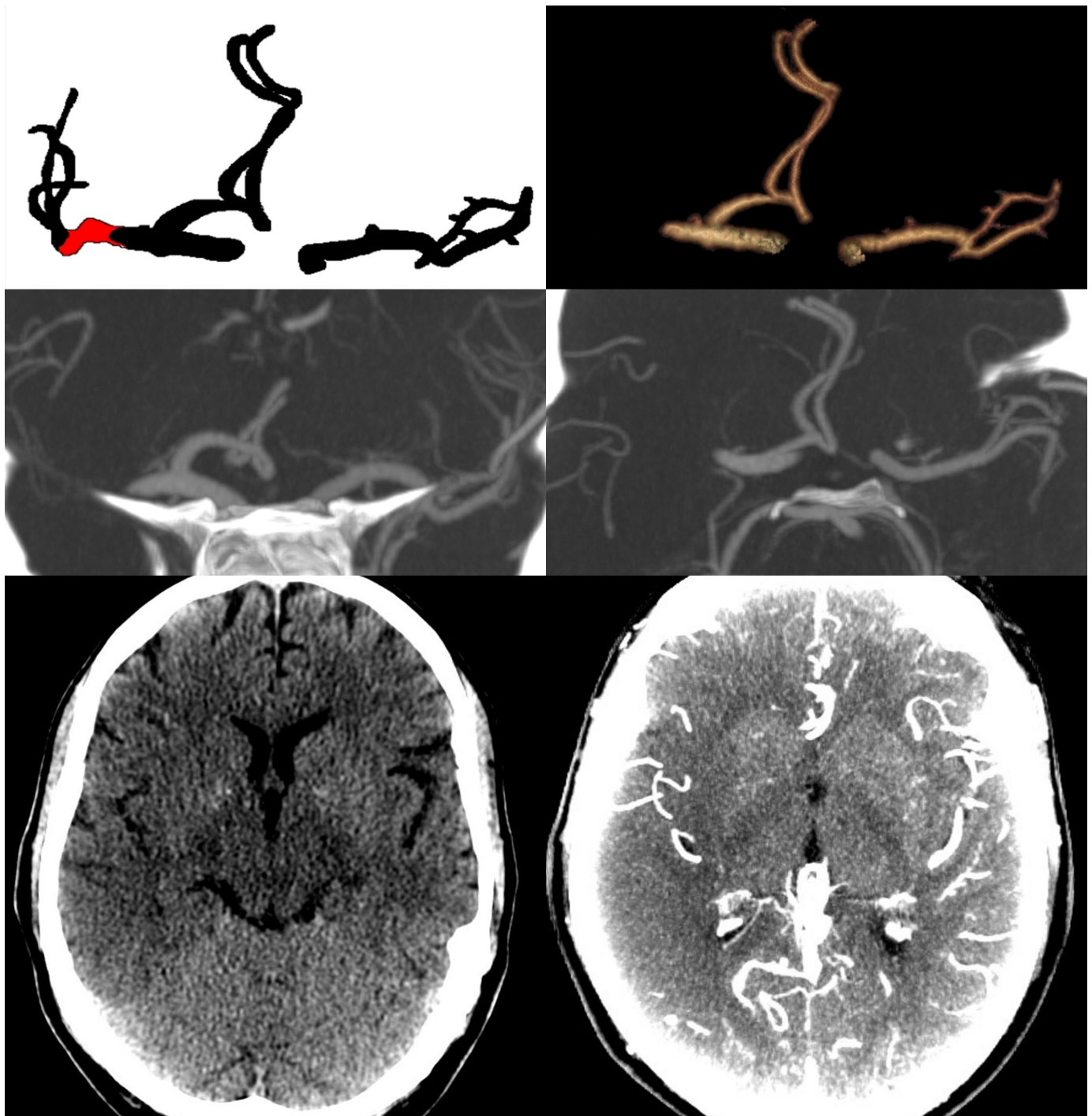


Figure 2 Carotid score 1, the circle of Willis is perfused from one carotid artery only, on non-enhanced CT, CT angiography and maximum intensity projection reconstructions.

clinical outcome (RR at discharge 1.44; 95% CI 0.80 to 2.49 and RR at 3 months 1.46; 95% CI 0.90 to 2.37). A higher carotid score was associated with a higher chance of good clinical outcome at 3 months ($p_{\text{trend}}=0.05$), but not at discharge ($p_{\text{trend}}=0.24$).

After adjustment for age, NIHSS at presentation, time to start of IAT, ASPECTS, CFG and CBS, the RRs remained essentially the same (online supplemental table).

DISCUSSION

The number of carotid arteries supplying the cerebral circulation (carotid score) in the setting of IAT for large (MI or T-top) artery occlusion seems to be an independent predictor of good clinical outcome after 3 months. However, we observed no relation between completeness of the circle of Willis and clinical outcome after IAT.

Incompleteness of the circle of Willis may result in worse cerebral perfusion and hence more damage in case

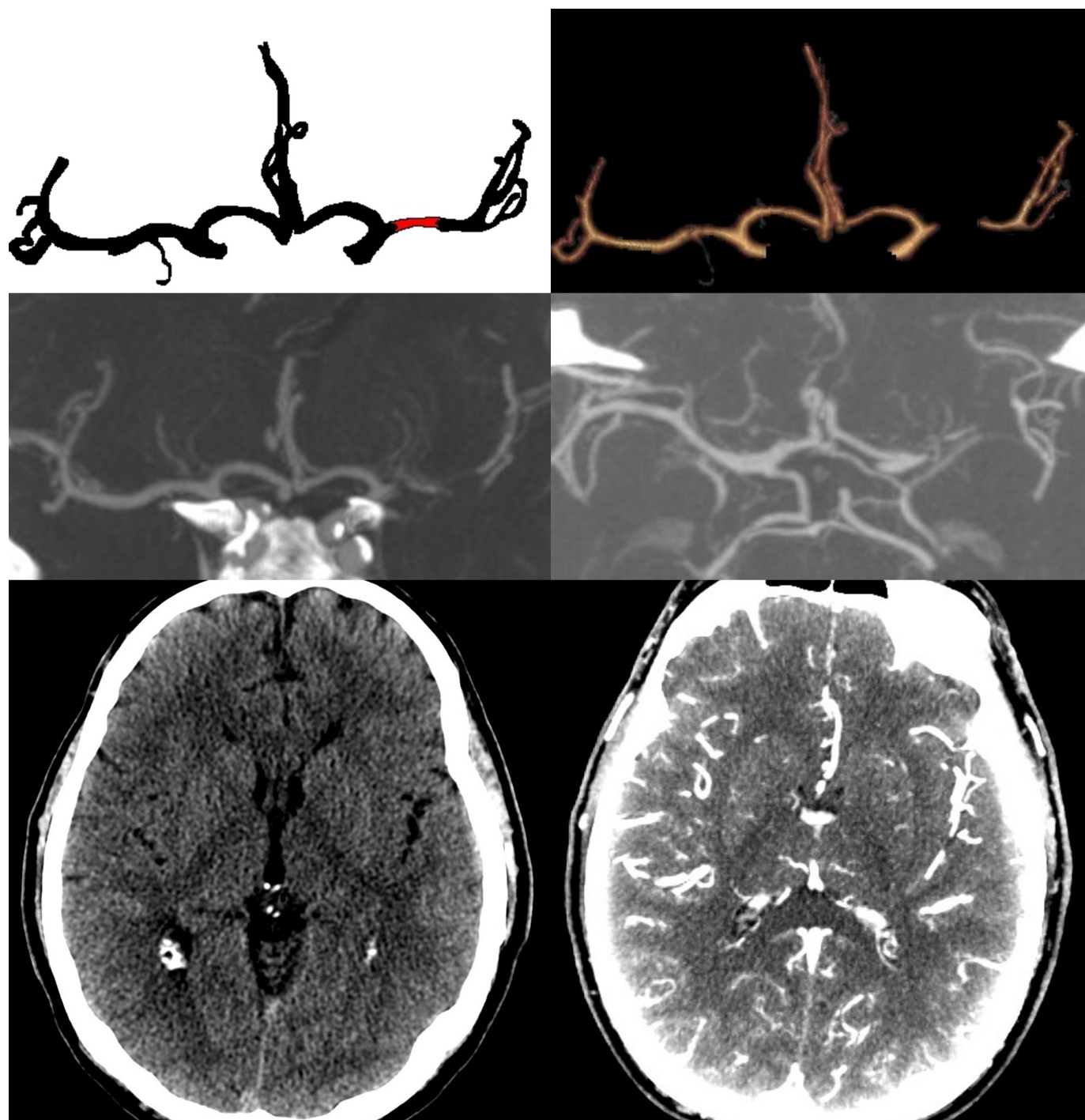


Figure 3 Carotid score 2, perfusion from two carotid arteries, on non-enhanced CT, CT angiography and maximum intensity projection reconstructions.

of acute cerebral ischaemia.^{10–12} Previous studies have shown that completeness of the circle of Willis, particularly the posterior circle of Willis, is a hallmark of deteriorated cerebral perfusion.^{24 25} In addition, patients with ipsilateral carotid occlusion before IAT in whom ACoA or PCoA or both were absent were shown to have less favourable clinical outcomes at 90 days.²⁶ We could not confirm such a detrimental effect in our patients who had a stroke with an incomplete circle of Willis treated with IAT. A possible explanation for this might be that in our

study, only three patients (2%) suffered from an ipsilateral carotid occlusion. We did find a negative influence of impaired carotid contribution by means of our carotid score. Hence, one could hypothesise that the sum of both the carotid occlusion and the incompleteness of the circle of Willis results in less favourable outcome in the aforementioned study. A larger, more recent study combined patients from the DEFUSE2 and CRISP Studies and studied completeness of the circle of Willis in relation to functional outcome after IAT. In line with our results, no

Table 1 Demographics

| | Total (n=163) | With CTA (n=126) |
|--|---------------|------------------|
| Median age (in years, range) | 63 (23–91) | 62 (23–89) |
| Men (%) | 98 (60) | 69 (55) |
| Median NIHSS (range) | 14 (3–38) | 14 (3–38) |
| Site of occlusion (%) | | |
| Carotid | 3 (2) | 3 (2) |
| Carotid T-top | 15 (9) | 11 (9) |
| Middle cerebral artery | 145 (89) | 112 (89) |
| Time onset to ER (median in minutes, range) | 62 (5–359) | 68 (5–359) |
| Time onset to IVT (median in minutes, range) | 105 (27–274) | 110 (27–274) |
| Time onset to IAT (median in minutes, range) | 207 (65–1111) | 207 (65–1111) |
| IVT (%) | 113 (69) | 89 (71) |
| IAT (%) | 159 (98) | 122 (97) |
| Use of thrombolytics (%) | 120 (74) | 97 (77) |
| Mechanical (%) | 133 (82) | 101 (82) |
| Failed procedure (%) | 4 (3) | 4 (3) |
| Radiological parameters | | |
| ASPECTS >7 (%) | 126 (75) | 106 (84) |
| Median Collateral Flow Grading (range) | | 3 (0–3) |
| Median Clot Burden Score (range) | | 3 (1–8) |
| Successful recanalisation (%) | 68 (44) | 59 (49) |
| Complete circle of Willis (%) | | 26 (21) |
| Carotid score | | |
| 0 | | 6 (5%) |
| 1 | | 42 (33%) |
| 2 | | 78 (62%) |

ASPECTS, Alberta Stroke Programme Early CT Score; CTA, CT angiography; ER, emergency room; IAT, intra-arterial treatment; IVT, intravenous thrombolysis; NIHSS, National Institutes of Health Stroke Scale.

association was found between incomplete circle of Willis and functional outcome.²⁷

Contralateral carotid occlusion is well known to increase risk of stroke or death in patients treated with carotid endarterectomy because of symptomatic carotid stenosis.²⁸ In addition, contralateral carotid artery stenosis has been shown to be an independent predictor of poor clinical outcome in patients who had a stroke with acute tandem occlusion treated with IAT.²⁹ These studies are in line with our results, showing that that loss of carotid arteries supplying the cerebral circulation reduces the likelihood of good clinical outcome.

Table 2 Clinical outcome at discharge and at 3 months

| | Circle of Willis | | | Carotid score* | | | P for trend |
|--|---------------------|---------------------|--------------|----------------|----------------------|----------------------|-------------|
| | Complete N=26 | Incomplete N=100 | All N=126 | 0 N=6 | 1 N=42 | 2 N=78 | |
| Good clinical outcome at discharge (%) | 34 (27) | 28 (28) | 62 (34) | 1 (17) | 9 (21) | 24 (31) | |
| Risk ratio (95% CI) | 0.82 (0.38 to 1.78) | Ref | | Ref | 1.29 (0.20 to 8.43) | 1.85 (0.30 to 11.39) | 0.24 |
| Good clinical outcome at 3 months (%) | 53 (42) | 42 (42) | 95 (53) | 1 (17) | 14 (33) | 38 (49) | |
| Risk ratio (95% CI) | 1.01 (0.61 to 1.67) | Ref | | Ref | 2.00 (0.32 to 12.59) | 2.92 (0.48 to 17.75) | 0.05 |

*Carotid score 0 — no carotid artery supplying cerebral circulation; carotid score 1 — one carotid artery supplying cerebral circulation; carotid score 2 — two carotid arteries supplying cerebral circulation.

Conversely, another study showed that there was no strong effect of a coexisting ICA stenosis in acute MCA stroke on tissue status or perfusion parameters on MRI.³⁰ Moreover, cerebral blood volume was found to be elevated in patients who had a stroke with coexisting ICA stenosis, possibly reflecting improved peripheral collateral circulation. However, patients with carotid occlusions were excluded in this study, only ipsilateral carotid stenosis was studied.

Our study has several limitations. First, our study focused on the collateral circulation by the circle of Willis and carotid arteries. Previous studies have shown that the leptomeningeal collaterals play a key role in chances of successful recovery after IAT.^{3 13–15 31 32} We, therefore, included a score for leptomeningeal collaterals (CFG)¹⁹ and corrected the RR for good clinical outcome for this score (see online supplemental table). After this adjustment, however, the RRs remained essentially the same. Second, we did not measure the flow in the ophthalmic artery. When flow is reversed in the ophthalmic artery, it also functions as collateral for the cerebral circulation. Nevertheless, this reversed flow is merely considered an indicator of diminished cerebral perfusion²⁴ and as such not a rescue pathway for cerebral circulation in case of an acute cerebral arterial occlusion. Third, extracranial carotid disease and basilar artery disease were not included in the study. One could hypothesise that any atherosclerotic disease on these locations also influences the cerebral collateral circulation. In addition, we did not register the presence of fetal-type posterior cerebral artery. The presence of such artery might substitute loss of blood flow in case of an occluded ipsilateral carotid artery and hence influence extent of infarction and functional outcome. However, given the rather low number of patients included in our study, we assume that these effects are limited. Fourth, patients included in our cohort were treated with IAT in the era before IAT was standard of care. We realise that our series is relatively old; however, we think we still retrieve valuable information from it about the pathophysiology of reperfusion and outcome. Although the applied IAT techniques were probably less sophisticated as they are nowadays, we do not expect that this had a substantial impact on our results because our study aimed at finding causal relation between completeness of the circle of Willis and clinical outcome after IAT and hence studied cause of disease.

A possible explanation for the lack of a positive relation between completeness of the circle of Willis and clinical outcome might be that we studied the primary circulation in the *acute* setting. Previous studies showed that collateralisation may take months to develop and animal models show that restoration of blood flow through collateral vessels after MCA occlusion takes a month.^{33 34} Most of these studies focused on the role of the collateral circulation by the circle of Willis in patients with atherosclerotic carotid artery disease. We did not select our cohort based on the presence of carotid artery disease. However, we did study the effect of the number of carotid arteries

supplying cerebral circulation and found a relation between this carotid score and clinical outcome.

In patients with acute ischaemic stroke treated with IAT, chances of good clinical outcome improve with the number of carotid arteries supplying the cerebral circulation. Completeness of the circle of Willis does not seem to relate to clinical outcome after IAT. Further studies are needed to confirm our findings in a larger cohort.

Contributors Conception—AR, GLaN, JB, MW, AA. Design—AR, AA, GLaN. Acquisition and analysis—AR, HH, WS, J-AV. Interpretation of data—AR, JB, MW, GLaN, AA. Draft of the work or substantive revision—AR, JB, J-AV, WS, MW, GLaN, AA. Guarantor - AR.

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Patient consent for publication Not required.

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REFERENCES

- 1 Berkhemer OA, Fransen PSS, Beumer D, *et al*. A randomized trial of intraarterial treatment for acute ischemic stroke. *N Engl J Med* 2015;372:11–20.
- 2 Campbell BCV, Mitchell PJ, Kleinig TJ, *et al*. Endovascular therapy for ischemic stroke with perfusion-imaging selection. *N Engl J Med* 2015;372:1009–18.
- 3 Goyal M, Demchuk AM, Menon BK, *et al*. Randomized assessment of rapid endovascular treatment of ischemic stroke. *N Engl J Med* 2015;372:1019–30.
- 4 Saver JL, Goyal M, Bonafe A, *et al*. Stent-retriever thrombectomy after intravenous t-PA vs. t-PA alone in stroke. *N Engl J Med* 2015;372:2285–95.
- 5 Jovin TG, Chamorro A, Cobo E, *et al*. Thrombectomy within 8 hours after symptom onset in ischemic stroke. *N Engl J Med* 2015;372:2296–306.
- 6 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.
- 7 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.
- 8 Angermaier A, Langner S, Kirsch M, *et al*. CT-angiographic collateralization predicts final infarct volume after intra-arterial thrombolysis for acute anterior circulation ischemic stroke. *Cerebrovasc Dis* 2011;31:177–84.
- 9 Jung S, Gilgen M, Slotboom J, *et al*. Factors that determine penumbral tissue loss in acute ischaemic stroke. *Brain* 2013;136:3554–60.

- 10 Henderson RD, Eliasziw M, Fox AJ, *et al.* Angiographically defined collateral circulation and risk of stroke in patients with severe carotid artery stenosis. *Stroke* 2000;31:128–32.
- 11 Hendrikse J, Hartkamp MJ, Hillen B, *et al.* Collateral ability of the circle of Willis in patients with unilateral internal carotid artery occlusion: border zone infarcts and clinical symptoms. *Stroke* 2001;32:2768–73.
- 12 Bisschops RHC, Klijn CJM, Kappelle LJ, *et al.* Collateral flow and ischemic brain lesions in patients with unilateral carotid artery occlusion. *Neurology* 2003;60:1435–41.
- 13 Nambiar V, Sohn SI, Almekhlafi MA, *et al.* Cta collateral status and response to recanalization in patients with acute ischemic stroke. *AJNR Am J Neuroradiol* 2014;35:884–90.
- 14 Sung SM, Lee TH, Cho HJ, *et al.* Functional outcome after recanalization for acute pure M1 occlusion of the middle cerebral artery as assessed by collateral cta flow. *Clin Neurol Neurosurg* 2015;131:72–6.
- 15 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.
- 16 Seeta Ramaiah S, Churilov L, Mitchell P, *et al.* The impact of arterial collateralization on outcome after intra-arterial therapy for acute ischemic stroke. *AJNR Am J Neuroradiol* 2014;35:667–72.
- 17 European Stroke Organisation (ESO) Guideline for management of acute stroke, 2009. Available: <http://www.eso-stroke.org> [Accessed 8 Sep 2018].
- 18 Puetz V, Dzialowski I, Hill MD, *et al.* Intracranial thrombus extent predicts clinical outcome, final infarct size and hemorrhagic transformation in ischemic stroke: the clot burden score. *Int J Stroke* 2008;3:230–6.
- 19 Barber PA, Demchuk AM, Zhang J, *et al.* Validity and reliability of a quantitative computed tomography score in predicting outcome of hyperacute stroke before thrombolytic therapy. *The Lancet* 2000;355:1670–4.
- 20 Higashida RT, Furlan AJ, Roberts H. Technology assessment Committee of the American Society of Interventional and therapeutic Neuroradiology: technology assessment Committee of the Society of interventional Radiology. Trial design and reporting standards for intra-arterial cerebral thrombolysis for acute ischemic stroke. *Stroke* 2003;34:e109–37.
- 21 Rankin J. Cerebral vascular accidents in patients over the age of 60. II. prognosis. *Scott Med J* 1957;2:200–15.
- 22 van Swieten JC, Koudstaal PJ, Visser MC, *et al.* Interobserver agreement for the assessment of handicap in stroke patients. *Stroke* 1988;19:604–7.
- 23 Knol MJ, Le Cessie S, Algra A, *et al.* Overestimation of risk ratios by odds ratios in trials and cohort studies: alternatives to logistic regression. *CMAJ* 2012;184:895–9.
- 24 Kluytmans M, van der Grond J, van Everdingen KJ, *et al.* Cerebral hemodynamics in relation to patterns of collateral flow. *Stroke* 1999;30:1432–9.
- 25 Rutgers DR, Klijn CJM, Kappelle LJ, *et al.* Recurrent stroke in patients with symptomatic carotid artery occlusion is associated with high-volume flow to the brain and increased collateral circulation. *Stroke* 2004;35:1345–9.
- 26 Zhao H, Wang B, Xu G, *et al.* Collateral grade of the Willis' circle predicts outcomes of acute intracranial internal carotid artery occlusion before thrombectomy. *Brain Behav* 2019;9:e01452.
- 27 Seifert-Held T, Eberhard K, Christensen S, *et al.* Circle of Willis variants are not associated with thrombectomy outcomes. *Stroke Vasc Neurol* 2021;6:e000491:310–3.
- 28 Touzé E, Trinquart L, Felgueiras R, *et al.* A clinical rule (sex, contralateral occlusion, age, and restenosis) to select patients for stenting versus carotid endarterectomy. *Stroke* 2013;44:3394–400.
- 29 Maus V, Behme D, Borggrefe J, *et al.* Carotid artery stenosis contralateral to acute tandem occlusion: an independent predictor of poor clinical outcome after mechanical thrombectomy with concomitant carotid artery stenting. *Cerebrovasc Dis* 2018;45:10–17.
- 30 Kaesemann P, Thomalla G, Cheng B, *et al.* Impact of severe extracranial ICA stenosis on MRI perfusion and diffusion parameters in acute ischemic stroke. *Front Neurol* 2014;5:1–7.
- 31 Jadhav AP, Diener H-C, Bonafe A. On behalf of the swift prime Investigators. Correlation between clinical outcomes and baseline CT and CT angiographic findings in the swift prime trial. *AJNR Am J Neuroradiol* 2017;38:2270–6.
- 32 van den Wijngaard IR, Boiten J, Holswilder G, *et al.* Impact of collateral status evaluated by dynamic computed tomographic angiography on clinical outcome in patients with ischemic stroke. *Stroke* 2015;46:3398–404.
- 33 Widder B, Kleiser B, Krapf H. Course of cerebrovascular reactivity in patients with carotid artery occlusions. *Stroke* 1994;25:1963–7.
- 34 Coyle P, Heistad DD. Blood flow through cerebral collateral vessels one month after middle cerebral artery occlusion. *Stroke* 1987;18:407–11.

Web table Adjusted Risk ratios for good clinical outcome at discharge and 3 months

| | | Complete vs incomplete | | Carotid score | |
|--------------|----------------------------------|------------------------|---------------------|---------------------|--------------------|
| | | Circle of Willis | 1 vs 0 | 2 vs 0 | 1 vs 2 |
| At discharge | Risk ratio (RR, 95% CI) | 0.82 (0.38 - 1.78) | 1.29 (0.20 - 8.43) | 1.85 (0.30 - 11.39) | 1.44 (0.74 - 2.80) |
| | Adjusted RR Age | 0.79 (0.38 - 1.67) | 1.13 (0.18 - 7.10) | 1.48 (0.24 - 9.11) | 1.32 (0.68 - 2.60) |
| | Adjusted RR NIHSS [†] | 0.84 (0.42 - 1.65) | 1.22 (0.20 - 7.46) | 1.48 (0.25 - 8.58) | 1.22 (0.63 - 2.33) |
| | Adjusted RR time to IAT | 0.86 (0.33 - 2.23) | 1.05 (0.16 - 6.90) | 1.49 (0.25 - 8.91) | 1.41 (0.61 - 3.25) |
| | Adjusted RR ASPECTS [‡] | 0.90 (0.44 - 1.85) | 1.76 (0.28 - 11.11) | 2.41 (0.40 - 14.41) | 1.37 (0.72 - 2.62) |
| | Adjusted RR CFG [§] | 0.96 (0.46 - 2.01) | 1.00 (0.19 - 5.12) | 1.26 (0.26 - 6.12) | 1.28 (0.67 - 2.44) |
| | Adjusted RR CBS [¶] | 0.82 (0.38 - 1.75) | 1.30 (0.20 - 8.66) | 1.83 (0.29 - 11.34) | 1.39 (0.71 - 2.72) |
| At 3 months | Risk ratio (RR, 95 CI) | 1.01 (0.61 - 1.67) | 2.00 (0.32 - 12.59) | 2.92 (0.48 - 17.75) | 1.46 (0.90 - 2.37) |
| | Adjusted RR Age | 0.98 (0.61 - 1.57) | 1.83 (0.29 - 11.56) | 2.53 (0.41 - 15.52) | 1.38 (0.84 - 2.24) |
| | Adjusted RR NIHSS [†] | 1.02 (0.62 - 1.67) | 1.90 (0.32 - 11.41) | 2.68 (0.46 - 15.72) | 1.42 (0.87 - 2.31) |
| | Adjusted RR time to IAT | 1.10 (0.61 - 2.00) | 1.82 (0.34 - 9.62) | 2.68 (0.54 - 13.30) | 1.47 (0.82 - 2.63) |
| | Adjusted RR ASPECTS [‡] | 1.08 (0.66 - 1.76) | 2.30 (0.37 - 14.21) | 3.30 (0.56 - 19.58) | 1.43 (0.89 - 2.32) |
| | Adjusted RR CFG [§] | 1.05 (0.62 - 1.76) | 1.56 (0.29 - 8.37) | 2.25 (0.43 - 11.66) | 1.45 (0.88 - 2.39) |
| | Adjusted RR CBS [¶] | 1.00 (0.60 - 1.65) | 1.99 (0.32 - 12.53) | 2.98 (0.48 - 18.32) | 1.51 (0.92 - 2.48) |

[†]NIHSS: National Institutes Health Stroke Scale. [‡]ASPECTS: Alberta Stroke Program Early CT score²¹. [§]CFG: Collateral Flow Grading²². [¶]CBS: Clot Burden Score²⁰.