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# Mechanical Thrombectomy for Isolated M2 Occlusions: A Post Hoc Analysis of the STAR, SWIFT, and SWIFT PRIME Studies

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## ABSTRACT

**BACKGROUND AND PURPOSE:** Mechanical thrombectomy is beneficial for patients with acute ischemic stroke and a proximal anterior occlusion, but it is unclear if these results can be extrapolated to patients with an M2 occlusion. The purpose of this study was to examine the technical aspects, safety, and outcomes of mechanical thrombectomy with a stent retriever in patients with an isolated M2 occlusion who were included in 3 large multicenter prospective studies.

**MATERIALS AND METHODS:** We included patients from the Solitaire Flow Restoration Thrombectomy for Acute Revascularization (STAR), Solitaire With the Intention For Thrombectomy (SWIFT), and Solitaire With the Intention for Thrombectomy as Primary Endovascular Treatment (SWIFT PRIME) studies, 3 large multicenter prospective studies on thrombectomy for ischemic stroke. We compared outcomes and technical details of patients with an M2 with those with an M1 occlusion. All patients were treated with a stent retriever. Imaging data and outcomes were scored by an independent core laboratory. Successful reperfusion was defined as modified Thrombolysis in Cerebral Infarction score of 2b/3.

**RESULTS:** We included 50 patients with an M2 and 249 patients with an M1 occlusion. Patients with an M2 occlusion were older (mean age, 71 versus 67 years;  $P = .04$ ) and had a lower NIHSS score (median, 13 versus 17;  $P < .001$ ) compared with those with an M1 occlusion. Procedural time was nonsignificantly shorter in patients with an M2 occlusion (median, 29 versus 35 minutes;  $P = .41$ ). The average number of passes with a stent retriever was also nonsignificantly lower in patients with an M2 occlusion (mean, 1.4 versus 1.7;  $P = .07$ ). There were no significant differences in successful reperfusion (85% versus 82%,  $P = .82$ ), symptomatic intracerebral hemorrhages (2% versus 2%,  $P = 1.0$ ), device-related serious adverse events (6% versus 4%,  $P = .46$ ), or modified Rankin Scale score 0–2 at follow-up (60% versus 56%,  $P = .64$ ).

**CONCLUSIONS:** Endovascular reperfusion therapy appears to be feasible in selected patients with ischemic stroke and an M2 occlusion.

**ABBREVIATIONS:** ESCAPE = Endovascular Treatment for Small Core and Proximal Occlusion Ischemic Stroke; IMS = Interventional Management of Stroke; MERCI = Mechanical Embolus Removal in Cerebral Ischemia; MT = mechanical thrombectomy; REVASCAT = Endovascular Revascularization With Solitaire Device Versus Best Medical Therapy in Anterior Circulation Stroke Within 8 Hours; STAR = Solitaire FR Thrombectomy for Acute Revascularization; SWIFT = Solitaire FR With the Intention For Thrombectomy; SWIFT PRIME = Solitaire With the Intention for Thrombectomy as Primary Endovascular Treatment

Recent data have shown that mechanical thrombectomy (MT) with a stent retriever is safe and improves functional outcome in patients with acute ischemic stroke and an occlusion of the anterior circulation.<sup>1–5</sup> It is unclear, however, if these results can

be extrapolated to patients with an occlusion of the second segment of the middle cerebral artery (M2 occlusion). Because of its distal location, smaller diameter, and thinner walls, MT of the M2 segment is technically more challenging and may be associated with a higher risk of periprocedural complications. The potential benefit of reperfusion may also be different, in part because M2 occlusions generally respond better to IV thrombolysis.<sup>6</sup>

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**Table 1: Anatomic definition of MCA segments**

MCA Segment	Anatomic Definition
M1	Horizontal segment of the proximal MCA from the bifurcation of the ICA into the anterior and middle cerebral arteries to the genu of the MCA branch or branches at the entrance to the insula
M2	Vertical MCA branches in the Sylvian fissure originating at the genu and extending to the next genu at the level of the operculum; if the anterior temporal artery arises from the horizontal M1 segment, it will not be considered an M2 branch
M3	Branches that continue when the M2 branches change course again to a more lateral/horizontal course in the Sylvian fissure away from the insula and below the operculum
M4	Branch vessels primarily extend vertically over the frontal and parietal lobes and inferiorly over the temporal lobe

The number of patients with an M2 occlusion in the thrombectomy trials was small. The Multicenter Randomized Clinical trial of Endovascular Treatment for Acute Ischemic Stroke in the Netherlands (MR CLEAN) protocol did allow recruitment of these patients, but <8% of included patients had an isolated M2 occlusion.<sup>1</sup> In the Endovascular Revascularization With Solitaire Device Versus Best Medical Therapy in Anterior Circulation Stroke Within 8 Hours (REVASCAT), Endovascular Treatment for Small Core Proximal Occlusion Ischemic Stroke (ESCAPE), and Extending the Time for Thrombolysis in Emergency Neurological Deficits–Intra-Arterial (EXTEND-IA), only 10, 6, and 4 patients with an isolated M2 occlusion were treated with MT, respectively.<sup>2,4,5</sup> The Solitaire With the Intention for Thrombectomy as Primary Endovascular Treatment (SWIFT PRIME) protocol excluded M2 occlusions, but there were some protocol violations.<sup>3</sup> Thus far, none of these studies has specifically examined the subgroup of M2 occlusions. The aim of our study was to examine the technical aspects, safety, and outcomes of MT with a stent retriever in patients with an isolated M2 occlusion who were included in 3 large multicenter prospective studies.

## MATERIALS AND METHODS

### Description of Studies and Patient Selection

We included patients from the Solitaire FR With the Intention for Thrombectomy (SWIFT) trial, Solitaire FR Thrombectomy for Acute Revascularization (STAR) study, and the SWIFT PRIME trial. The design of these studies has been reported previously.<sup>3,7-9</sup> Briefly, the SWIFT trial was a randomized clinical trial that ran from 2010 to 2011, in which patients with acute ischemic stroke and an angiographically confirmed occlusion of a the proximal cerebral artery were allocated to treatment with a stent retriever (Solitaire; Covidien, Irvine, California) or the Merci device (Concentric Medical, Mountain View, California). SWIFT had a roll-in phase during which all patients were treated with a stent retriever, and these patients were included in the current study. STAR was a prospective, multicenter, single-arm study from 2010 to 2012 in which patients with an occlusion of the anterior circulation (intracranial and terminus internal carotid artery and M1 and M2 segments of the middle cerebral artery) were included. All patients in STAR were treated with a stent retriever (Solitaire). In both SWIFT and STAR, patients were eligible only if MT was possible within 8 hours of symptom onset. In SWIFT PRIME, patients with an acute ischemic stroke and confirmed occlusion of the anterior intracranial circulation were randomized between intravenous tPA followed by MT with a stent retriever or intravenous tPA alone. The trial ran from December 2012 until Novem-

ber 2014. In all 3 studies, clinical outcome was determined at 90-day follow-up by using the modified Rankin Scale.

We compared outcomes and technical details of patients with an M2 occlusion with those with an M1 occlusion. We excluded patients from SWIFT treated with the Merci device and patients from the control arm of SWIFT PRIME. Patients with a combined M1 and M2 occlusion were categorized as having M1 occlusion. Those with an occlusion at another location (eg, carotid terminus) were excluded from the analysis. The local ethics committee at every site approved the study protocol, and all patients or their legal representatives gave written informed consent.

Outcome data were adjudicated by an independent CT and MR imaging core laboratory, an angiography core laboratory, and a clinical events committee. The angiography core laboratory assessed the location of the occlusion. Table 1 describes the definition for each segment of the middle cerebral artery used by the core laboratory of the 3 studies. Other variables scored by the core laboratory or Clinical Events Committee were final revascularization grades, hemorrhagic complications, and other adverse events. In SWIFT and SWIFT PRIME, the assessors were blinded to study group assignments.

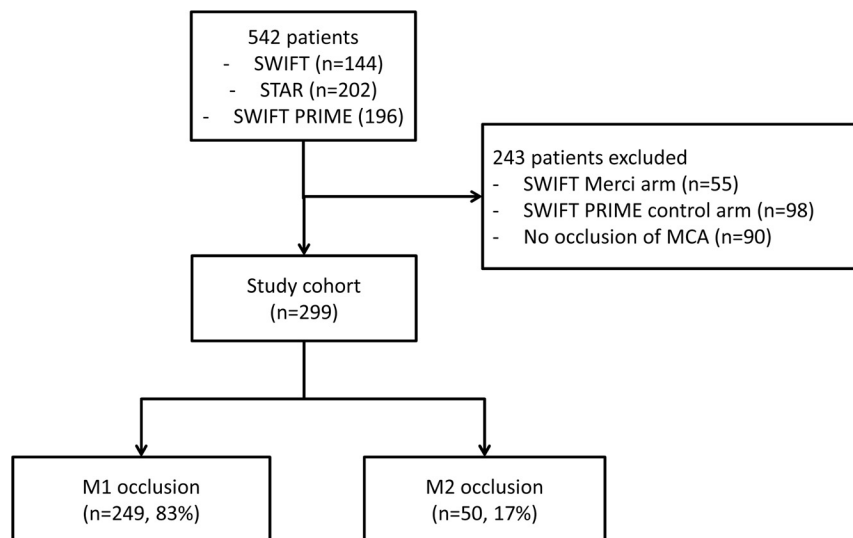
### MT Procedure

The MT procedure used in STAR, SWIFT, and SWIFT PRIME has been described previously.<sup>7-9</sup> The aim was to achieve successful recanalization of the territory of the occluded vessel. The use of a balloon-guide catheter was mandatory in STAR and SWIFT and optional in SWIFT PRIME. Up to 3 passes with the stent retriever were allowed according to the protocols. The decision to perform any additional passes was at the discretion of the interventionalist and was scored as a protocol violation. The procedure could be performed with the patient under local or general anesthesia. Follow-up brain imaging was performed after 24 hours in all patients.

### Data Analysis

We compared the technical details of the procedure, complication rate, and clinical outcome at follow-up between patients with an M1 versus M2 occlusion. Successful reperfusion was defined as a modified Thrombolysis in Cerebral Infarction score of  $\geq 2b$  of the target territory. Both the proportions of symptomatic and any intracranial hemorrhage are provided. Symptomatic intracranial hemorrhage was defined as hemorrhage within 24 hours associated with an increase on the NIHSS of  $\geq 4$  points or that resulted in death. Clinical outcome at 90 days is provided as mRS 0–1, mRS 0–2, and all-cause mortality.

All analyses were by intention-to-treat. Categorical variables



**FIG 1.** Flowchart of patient selection.

**Table 2: Baseline characteristics<sup>a</sup>**

	M2 Occlusion (n = 50)	M1 Occlusion (n = 249)	P Value
Age (yr) (mean)	71 ± 11	67 ± 13	.04
Female	48% (24/50)	57% (143/249)	.28
NIHSS score (median) (IQR)	13 (10–17)	17 (14–20)	<.001
Medical history			
Atrial fibrillation	38% (19/50)	37% (93/249)	1.000
Hypertension	70% (35/50)	61% (152/249)	.27
Diabetes	18% (9/50)	15% (38/249)	.67
Hyperlipidemia	30% (15/50)	37% (93/249)	.42
Peripheral artery disease	0% (0/50)	4% (9/249)	.37
Current smoker	14% (7/49)	21% (53/249)	.33
Prior stroke/TIA	12% (6/50)	17% (42/249)	.53
Systolic BP (mean) (mm Hg)	150 ± 28	147 ± 23	.45
Diastolic BP (mean) (mm Hg)	84 ± 17	80 ± 15	.11
Left-sided occlusion	54% (27/50)	47% (117/249)	.44
Baseline ASPECTS (mean)	9.0 ± 1.3	8.2 ± 1.7	.003
Receipt of IV tPA	77% (36/47)	69% (161/234)	.38
Laboratory data			
Baseline serum glucose level (mean)	127 ± 51	128 ± 55	.89
INR (mean)	1.1 ± 0.5	1.1 ± 0.3	.95
Platelets (mean)	235 ± 75	234 ± 79	.95
aPTT (mean) (sec)	31 ± 20	29 ± 22	.44

**Note:**—IQR indicates interquartile range; BP, blood pressure; INR, international normalized ratio; aPTT, activated partial thromboplastin time.

<sup>a</sup> Categorical variables are given as percentages with *n*/*N* in parentheses, where *n* is the number of patients in whom the variable was present and *N* the total number of patients for whom that particular variable was reported.

were compared between groups by using a Fisher exact test. Continuous variables were compared by using a *t* test, except in the cases in which medians and interquartile ranges are reported; then, a Wilcoxon rank sum test was used. Statistical analyses were conducted in SAS, Version 9.3 (SAS, Cary, North Carolina).

### Role of the Funding Source

In all 3 studies, an academic steering committee supervised the trial design and operations. The sponsor of the study (Covidien) was responsible for site management, data management, and safety reporting. The study data were independently monitored. The statistical analyses were prepared by independent external statisticians (J. Schafer; Namsa, Minneapolis, Minnesota; S Brown, Altair Biostatistics, St. Louis Park, Minnesota).

The sponsor had no role in the preparation of the article. The corresponding author had full access to all the data in the study and had final responsibility for the decision to submit for publication. The studies were registered at ClinicalTrials.gov, numbers NCT01054560 (SWIFT), NCT01327989 (STAR), and NCT01657461 (SWIFT-PRIME).

### RESULTS

Of the 542 patients enrolled in SWIFT, STAR, and SWIFT PRIME, 243 patients were excluded because they were allocated to the Merci arm of SWIFT (*n* = 55) or the control arm of SWIFT PRIME (*n* = 98) or because they did not have an occlusion of the middle cerebral artery (*n* = 90, Fig 1). Of the 299 included patients, 50 had an M2 occlusion; and 249, an M1 occlusion. Patients with an M2 occlusion were older (71 versus 67 years, *P* = .04) and had a lower median NIHSS score (13 versus 17, *P* < .001) compared with patients with an M1 occlusion (Table 2). There were no important differences in medical histories between the groups. Baseline ASPECTS was higher in those with an M2 occlusion (9.0 versus 8.2, *P* = .003). The proportion of patients who received intravenous tPA before MT did not differ significantly (77% versus 69%, *P* = .38).

Time from groin puncture to recanalization was nonsignificantly shorter in patients with an M2 occlusion (median, 29 versus 35 minutes; *P* = .41, Table 3). The average number of passes with the stent retriever was also nonsignificantly lower in patients with an M2 occlusion (mean, 1.4 versus 1.7; *P* = .07). There was no difference in the proportion of patients in whom modified TICI ≥2b was achieved (85% versus 82%,

*P* = .82) or in the incidence of symptomatic intracranial hemorrhages (both 2%). An mRS of 0–1 at 90 days was achieved in 50% of patients with an M2 occlusion, compared with 41% in those with an M1 occlusion (*P* = .27).

### DISCUSSION

We examined the feasibility and safety of MT with a stent retriever in patients with an isolated M2 occlusion. We found no significant differences in the reperfusion rate or risk of complications between patients with an M1 or M2 occlusion. If anything, there was a trend toward a lower number of passes required with stent retrievers in patients with an M2 occlusion. Clinical outcomes were nonsignificantly better in patients with an M2 occlusion,

**Table 3: Details of thrombectomy procedure and clinical and safety outcomes<sup>a</sup>**

	M2 Occlusion (N = 50)	M1 Occlusion (N = 249)	P Value
Time from groin puncture to recanalization (min) (median) (IQR)	29 (22–45)	35 (25–52)	.41
No. of passes with stent retriever (mean)	1.4 ± 0.8	1.7 ± 1.0	.07
≥3 Passes with stent retriever	13% (5/38)	23% (52/227)	.21
mTICI 2b or 3 reperfusion	85% (34/40)	82% (193/235)	.82
Rescue therapy	6% (3/50)	8% (19/249)	1.000
Complications			
Device-related serious adverse events	6% (3/50)	4% (10/249)	.46
Symptomatic ICH	2% (1/50)	2% (5/249)	1.000
Outcome at 90-day follow-up			
mRS 0–1	50% (25/50)	41% (100/243)	.27
mRS 0–2	60% (30/50)	56% (136/243)	.64
Mortality	12% (6/50)	10% (25/249)	.62

**Note:**—mTICI indicates modified Thrombolysis in Cerebral Infarction; ICH, intracerebral hemorrhage; IQR, interquartile range.

<sup>a</sup> Categorical variables are given as percentages with *n*/*N* in parentheses, where *n* is the number of patients in whom the variable was present, and *N* is the total number of patients for whom that particular variable was reported.

though this result most likely is because these patients had milder strokes, as shown by the lower baseline NIHSS scores.

Few studies have previously examined endovascular treatment of patients with an M2 occlusion, and in most of these, intra-arterial thrombolysis or earlier generation thrombectomy devices were used. A post hoc analysis of the Prolyse in Acute Cerebral Thromboembolism (PROACT) II, Interventional Management of Stroke (IMS), and IMS II studies found successful reperfusion in approximately half of all patients with an M2 occlusion.<sup>10</sup> Shi et al<sup>11</sup> studied 28 patients with an M2 occlusion who were enrolled in the Mechanical Embolus Removal in Cerebral Ischemia (MERCİ) and multi-MERCİ trials. They found that compared with patients with an M1 occlusion, the number of required passes with the thrombectomy device was lower in those with an M2 occlusion, similar to our data. They also found higher recanalization rates in patients with an M2 occlusion. Sheth et al<sup>12</sup> collected data on M2 occlusions from their single-center cohort on MT. They found no difference in recanalization rates between patients with an M2 or M1 occlusion, but the risk of intracranial hemorrhagic complications was lower in patients with an M2 occlusion. However, this observation could be related to different endovascular techniques that were used in each group. Most patients with an M2 occlusion were treated with intra-arterial tPA, while the Merci device was predominantly used in patients with an M1 occlusion. Various randomized trials have shown that the methods of endovascular reperfusion that were used in the above-mentioned studies do not improve clinical outcome in patients.<sup>13–15</sup> With the introduction of stent retrievers, these older techniques are now rarely applied; thus, the external validity of these studies is now limited.

Only 2 recent retrospective studies have examined the use of stent retrievers for patients with an M2 occlusion. Dorn et al<sup>16</sup> reported outcomes of 15 patients with an M2 occlusion from a single-center retrospective registry. Similar to our results, their data do not suggest that patients with an M2 occlusion have a higher risk of procedural complications. The other study used data from a regional stroke registry.<sup>17</sup> Most of the patients in this series were treated with a stent retriever, though other techniques were used in 14% of patients. Their rate of symptomatic intracerebral

hemorrhage was 9%, which is somewhat higher than that in our study. However, they did not assess patients with an M1 occlusion for comparison; this feature makes interpretation more difficult. Their patients were also in a more severe clinical condition, with a median NIHSS score of 16, which is comparable with the NIHSS scores of the patients with an M1 occlusion in our study. A drawback of both studies is that they did not use an independent core laboratory to adjudicate the angiographic data and no information is provided on the definition that was used for an M2 occlusion.

Our study had no control group of medically treated patients available for comparison. As a result, we cannot de-

termine whether MT with a stent retriever is superior to best medical management in patients with an M2 occlusion. Individually, neither of the recent thrombectomy trials has a sufficient number of patients with an M2 occlusion to answer this question. One issue that is important in any study that examines MT for M2 occlusion is a clear description of the definition of an M2 branch. The original classification by Fischer in 1938<sup>18</sup> designated the M1 segment as terminating at the genu where the MCA takes a 90° upward turn around the limen insulae; this definition was used in the STAR, SWIFT, and SWIFT PRIME studies. The M1 segment is then further divided into prebifurcation and postbifurcation segments with up to 82% of MCAs reported to bifurcate proximal to the genu.<sup>19,20</sup> This terminology creates some degree of confusion because postbifurcation branches proximal to the genu are commonly designated as M2 segments.

There are also differences in the number and dominance of the divisions of the MCA, meaning that not all M2 divisions are the same with regard to the volume or importance of the territory that they supply. The variation in size of the divisions also means that technically not all M2 branches are accessible for mechanical thrombectomy. M2 divisions are smaller; therefore, navigating a microcatheter and deploying a stent retriever can be more challenging and associated with greater risk. Thus, when reviewing the patients included in this study, one must consider that the M2 occlusions were likely selected because they had features that made them favorable for mechanical thrombectomy, either because of their size or because they were critical to recanalize due to their supply to and area of eloquence. Patients also were required to have an NIHSS score of at least an 8, which decreases the probability of smaller M2 branches being included. This selection of M2 occlusions is an important limitation of this study, and one that probably applies to all studies that examined MT for M2 occlusions.

Only high-volume stroke centers with extensive experience with MT participated in the SWIFT, STAR and SWIFT PRIME studies. In SWIFT, only centers that had participated in the MERCİ or multi-MERCİ trials or who had an annual MT volume of at least 30 patients were eligible. In STAR, the steering committee selected centers with 24-hour availability of MT that were

experienced with the use of stent retrievers. As a result, caution is required when generalizing these results to stroke centers in general. On the other hand, given the positive results of the recent trials, MT volumes and thus experience of neurointerventionalists are likely to increase substantially in stroke centers within the next few years.

## CONCLUSIONS

In this study, we examined the technical aspects, safety, and outcomes of mechanical thrombectomy in patients with an isolated M2 occlusion who were included in 3 large multicenter prospective studies. Our data suggest mechanical thrombectomy with a stent retriever can be considered in selected patients with acute ischemic stroke and an isolated M2 occlusion.

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UNRELATED: Consultancy: Covidien; Expert Testimony: Tanoury, Nauts, McKinney & Garbarino, PLLC\*; Grants/Grants Pending: Stryker, Hackney, Grover, Hoover & Bean, PLC,\* Covidien,\* Genentech,\* MicroVention,\* Comments: research grants; OTHER: Dr Zaidat serves as a consultant to Covidien. Adnan H. Siddiqui—RELATED: Consulting Fee or Honorarium: Covidien,\* Comments: SWIFT PRIME Steering Committee member, Advisory Board; UNRELATED: Consultancy: Codman & Shurtleff, Guidepoint Global Consulting, Penumbra (Penumbra 3D Separator Trial Steering Committee member; received honoraria), Stryker, Pulsar Vascular, MicroVention Flow-Redirection Endoluminal Device trial Steering Committee member, Lazarus Effect, Blockade Medical, Reverse Medical, W.L. Gore & Associates; Payment for Lectures (including service on Speakers Bureaus): Codman & Shurtleff; Other: Financial Interests: Lazarus Effect, Blockade Medical, Hotspur, Intratech Medical, StimSox, Valor Medical, Medina Medical; Advisory Board: Intersocietal Commission for the Accreditation of Vascular Laboratories, Medina Medical; Honoraria: Abbott Vascular, Toshiba American Medical Systems; OTHER: Dr Siddiqui serves as a consultant to Covidien. Mayank Goyal—RELATED: Consulting Fee or Honorarium: Covidien, Comments: SWIFT PRIME Executive Committee member as the Global Imaging and Workflow Principal Investigator; also received honoraria for speaking and teaching engagements; UNRELATED: Grants/Grants Pending: Covidien,\* Comments: Dr Goyal is one of the Principal Investigators for the ESCAPE Trial, which was partially funded by Covidien through an unrestricted grant to the University of Calgary; Patents (planned, pending or issued): GE Healthcare, Comments: systems and methods for diagnosing strokes, multiphase CTA technology: licensed for further development and marketing of technology; OTHER: Dr Goyal serves as a consultant to Covidien. \*Money paid to the institution.

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