ABSTRACT

BACKGROUND AND PURPOSE: IVT administered in acute ischemic stroke provides low recanalization rates in proximal intracranial occlusions, with consequently poor clinical outcome. The safety and efficacy of an IES by using mechanical thrombectomy after IVT failure were assessed in acute MCA occlusions.

MATERIALS AND METHODS: Patients presenting with acute MCA occlusion within 4.5 hours with an NIHSS score between 8 and 25 and a DWI ASPECTS of >5 were eligible. From September 2009 to September 2010, mechanical thrombectomy by using the Solitaire FR device was systematically performed if no clinical improvement was observed 1 hour after the initiation of IVT (IES group). Results in terms of clinical outcome were compared with those from an IVT series from January 2007 to August 2009 (IVT group).

RESULTS: Alteplase was administered in 123 patients with proximal intracranial occlusion. Fifty-six had a confirmed MCA occlusion: 32 were included in the IVT group; and 24, in the IES group. At 24 hours, the median NIHSS improvement was 8.5 points in the IES group (25%–75% CI, 1.5–13) and 3 points in the IVT group (25%–75% CI, 1–5) (P = .001). At 3 months, 17/22 (77%) patients from the IES group and 15/30 (50%) from the IVT group had an mRS score of ≤2. After adjustment for confounding variables, IES was strongly associated with favorable clinical outcome (77% versus 50%; adjusted odds ratio = 11.9; 95% CI, 1.6–89.1; P < .02). No symptomatic intracranial hemorrhage was observed.

CONCLUSIONS: IES by using systematic mechanical thrombectomy after IVT failure safely improves the clinical outcome at 3 months and could represent an interesting alternative in the management of patients with acute MCA occlusion.

ABBREVIATIONS: ASPECTS = Alberta Stroke Program Early CT Score; CI = confidence interval; IA = intra-arterial; IES = IV-endovascular strategy; IV = IV thrombolysis; mRS = modified Rankin Scale
acute ischemic stroke due to an MCA occlusion within 4.5 hours of onset.

MATERIALS AND METHODS

Patients

From January 2007 to September 2010, all consecutive patients presenting with an acute ischemic stroke at Montpellier University Hospital, France, were included in this study if they fulfilled the following criteria: 1) a symptom onset from IVT infusion of <4.5 hours, 2) a stroke in the MCA territory with a confirmed proximal occlusion (M1 segment) on MRA, and 3) an NIHSS score between 8 and 25. Patients who presented with at least 1 of the following criteria were excluded from the study: 1) a thrombolysis contraindication either in accordance with the National Institute of Neurological Disorders and Stroke guidelines or a derived MRI imaging ASPECTS of <5, indicating an extensive stroke; and 2) an associated occlusion in another intracranial arterial territory or a tandem occlusion in the internal carotid artery.

Study Design

As a routine practice, all patients presenting with an acute stroke underwent MRA a few minutes after their admission in parallel with an initial NIHSS score assessment. Pacemaker and patient agitation were the main MRA contraindications.

After a review of our therapeutic strategy in September 2009, eligible patients were then split into 2 groups: 1) Patients from the first study period (January 2007 to August 2009, 32 months) treated with standard IVT alone were registered in the IVT group, and 2) patients from the second study period (September 2009 to September 2010, 12 months) treated by using the IES were registered in the IES group. The IES group included patients who received either a rescue thrombectomy following IVT failure or a successful IVT. Demographic data, medical history, vascular risk factors, imaging findings, and clinical data for every patient with a confirmed acute ischemic stroke were retrieved from the medical records. The clinical outcome of patients was systematically assessed by a neurologist using the NIHSS during the acute phase (at admission and 24 hours later) and the mRS at 3 months. An mRS score ranging from 0 to 2 was considered a favorable clinical result. Informed consent was obtained from all patients or family members, and the local ethics committee approved the study.

Procedures

In our institution, all patients presenting with a suspected acute ischemic stroke had systematic MRA on admission. DWI, T2, FLAIR, 3D time-of-flight sequences, and neck vessels were acquired and analyzed to confirm the ischemic lesion and the MCA occlusion. As described, the ASPECTS12 was applied to the DWI sequence to evaluate the parenchymal extension of the ischemic lesion.14 Once the decision to treat was made, IVT was immediately administered. In the first study period, alteplase alone (0.9 mg/Kg) was infused with systematic clinical examinations (NIHSS score) at 1 hour and 24 hours. In the second study period, additional mechanical thrombectomy was discussed for each patient after the 1-hour NIHSS score. If this second NIHSS score was <8 or if it improved by at least 4 points in comparison with the baseline score, IVT was considered successful without additional therapy.

In cases of failed IVT, a mechanical thrombectomy was undertaken as soon as possible, by using a stent retriever (Solitaire FR, 4 × 20 mm; ev3, Irvine, California), with the patient under general anesthesia. An 8F Merci balloon-guiding catheter (Concentric Medical, Mountain View, California) was introduced through a femoral sheath into the appropriate carotid artery. A 0.021-inch-internal-diameter microcatheter, Prowler Select Plus (Cordis, Miami Lakes, Florida), was navigated distal to the point of occlusion over an 0.014-inch steerable microwire. A microcatheter angiographic run was then carried out to confirm the microcatheter position distal to the thrombus. The Solitaire FR was then introduced through the microcatheter, and the device was deployed across the occluding thrombus. The microcatheter distal marker was positioned at the exact level of the proximal marker of the stent. An intravenous bolus of 1000 IU of standard heparin was administered at this time. On Solitaire FR deployment, an angiographic control was performed to evaluate the correct placement and expansion of the device. After the angiographic run, the Solitaire FR was maintained in place for a few minutes (3–7 minutes) to allow device expansion. After this time, the fully deployed Solitaire FR and the delivery microcatheter were gently pulled back together and recovered through the guiding catheter. The Solitaire FR was at no point completely or partially reshathed. During the retrieval, the balloon-guide catheter was inflated to suspend antegrade flow.

Manual aspiration with a 50-mL syringe was performed through the hemostatic valve during the retrieval to reverse the flow and to aspirate clot debris eventually present in the lumen of the guide catheter. A maximum of 5 passes was performed by using the device, if necessary. The integrity of the stent was consistently checked after every pass. No additional devices were used with the Solitaire FR, and it was not used in combination with intra-arterial rtPA in any case. The final MCA recanalization rate was assessed on digital subtraction angiography by using the Thrombolysis In Cerebral Infarction scoring system. A noncontrast CT was performed at day 1 postprocedure in all patients to exclude intracranial hemorrhage. Intracranial hemorrhage was considered symptomatic if a related increase of the NIHSS score of >4 points was observed. The Solitaire FR device is certified by the European Union.

Statistical Analysis

The main characteristics of patients were described at baseline to identify any significant clinical difference between the 2 groups. The Fisher exact and t tests were used to compare qualitative and quantitative variables between the 2 groups, respectively. The NIHSS score was also compared at 24 hours. At 3 months, an mRS score of ≤2 was used to ascertain a good clinical outcome. Association between IES and favorable outcome at 3 months was assessed by using an exact logistic regression analysis, adjusting for the initial NIHSS score, the most important confounding factor in our study. Due to the relatively small number of events in our sample, a regular logistic regression model would not be appropriate because the underlying asymptotic theory would not apply.
All statistical analyses were carried out by using STATA, Version 11 software (StataCorp, College Station, Texas).

RESULTS

Patient and Procedure Characteristics

From January 2007 to September 2010, a total of 256 patients received IVT for an acute ischemic stroke within 4.5 hours of symptom onset. Among them, 123 (48%) presented with a confirmed proximal artery occlusion, including 56 (22%) within the M1 segment. Thirty-two of 56 (57%) were in the IVT group, and 24/56 (43%) were in the IES group (Fig 1). Mean age, sex ratio, medical history, clinical score, and time from the onset of symptoms to IVT are summarized in Table 1. Overall, the patient characteristics were similar between the 2 groups. However, patients included in the IVT group tended to have a better initial NIHSS score than those in the IES group (median NIHSS = 15 [25%–75% CI, 7–19] and 16 [25%–75% CI, 12–19], respectively) and a higher mean time from symptom onset (195 ± 46.4 versus 169 ± 50.7 minutes, respectively). These were not shown to be significant (Table 1), and an adjustment for confounding variables between the 2 groups was performed for the interpretation of our results. The mean door-to-needle time was 62 ± 13 minutes in the IES group and 69 ± 12 minutes in the IVT group (Table 1). The IES group included 7/24 (29%) cases of successful IVT and 17/24 (71%) cases requiring additional mechanical thrombectomy secondary to IVT failure (Fig 1).

All 56 included patients underwent MRA with a sufficient image quality to make the decision to treat. Four (7%) were lost to follow-up (2 patients in each group) and did not have clinical assessment at 3 months. Two of 56 (3.6%) patients died from malignant ischemic stroke (1 patient in the IVT group and 1 patient in the IES group) and were both considered as having unfavorable results at 3 months (mRS = 6).

Recanalization

A noncontrast CT was performed at 24 hours in all patients to exclude an intracranial hemorrhage. However, recanalization of the MCA was not systematically assessed after alteplase infusion. In the IES group, all patients undergoing thrombectomy had a persistent M1 occlusion confirmed on initial digital subtraction angiography. Arteries of 16 of 17 patients (94%) recanalized after thrombectomy with 15/17 (88%) achieving complete recanalization (Thrombolysis In Cerebral Infarction 3) and 1/17 (6%) achieving partial recanalization (Thrombolysis In Cerebral Infarction 2a). In 1 case, the MCA did not reconstitute following mechanical thrombectomy (6%) (Thrombolysis In Cerebral Infarction 0). In the 16 patients who benefited from a MCA recanalization, the mean time from symptom onset to recanalization was 297 ± 61.6 minutes and the mean time from femoral artery puncture to recanalization was 38 ± 20.9 minutes. The mean time from femoral puncture to deployment of the Solitaire device was 26.5 ± 7.6 minutes.

Clinical Outcome

The median NIHSS score improvement between admission and 24 hours was 3 (25%–75% CI, 1–5) points in the IVT group and 8.5 (25%–75% CI, 1.5–13) points in the IES group (difference, 5.5 points; P < .001) (Table 2). For 6 of 56 participants (11%), it was
DISCUSSION

The results of this observational study suggested that our therapeutic strategy (IES) based on rescue thrombectomy after initial IVT failure provided better clinical benefits for the patients in comparison with standard IVT alone. As expected, these favorable results were certainly due to higher recanalization rates (94%). Furthermore, this strategy did not induce additional adverse events.

Previous results of IVT studies reported that MCA recanalization could be achieved in one-third of patients (between 32% and 35%), with subsequent favorable clinical outcome at 3 months. Recently, it has been demonstrated that this recanalization rate can be significantly improved by using a combined IV-IA approach and different mechanical devices in cases of thrombectomy. However, results in terms of clinical benefit at 3 months are mixed. On the basis of these observations and our preliminary protocol evaluation, the management of acute ischemic stroke in our institution was reviewed by our multidisciplinary stroke team in September 2009 and a compromise between IVT alone and combined IV-endovascular therapy was chosen. An IES by using mechanical thrombectomy (Solitaire FR) as a rescue therapy only in cases of IVT failure was systematically introduced for MCA occlusions.

In our center, the careful assessment of the acute ischemic lesions represents an important point in the decision to treat. Although parenchymal extension of the ischemic lesion by using the CT-ASPECTS was described as an alternative approach to the rule of one-third of the cerebral hemisphere, the DWI sequence is more objective in detecting ischemic lesions with a clearer contrast than CT. Thus, the DWI-ASPECTS applied was systematically assessed before the IVT decision, and patients having a predictive score of death (DWI-ASPECTS ≤ 4) were not treated and were excluded from the study. In addition, a DWI-ASPECTS of > 7 has been shown to be a good prognostic factor for patients. This was also observed in our setting, though it did not reach statistical significance, possibly due to a suboptimal sample size. Our findings also confirmed that the initial clinical severity on presentation is correlated to the clinical prognosis. As shown before, the initial NIHSS score was identified as an independent predictive factor for favorable clinical outcome at 3 months.

Previous studies of acute MCA occlusions reported a favorable clinical outcome at 3 months in 41% and 63% of patients by using a combined IV-endovascular strategy, in 56% by using combined IV-IA thrombolysis, and in 40% and 53% by using IA thrombolysis alone. Comparatively, patients included in our IES group presented with high rates of favorable outcome at 3 months (77%). This was also the case in patients who underwent only mechanical thrombectomy (67%). The implementation of thrombectomy after initial IVT failure markedly improved the favorable outcome rate at 3 months, most particularly after adjustment for the initial clinical severity (NIHSS) between the 2 groups, with statistical significance. Stroke severity at admission was identified as the most important predictive factor of clinical outcome. Given that patients included in the IES group had higher initial NIHSS scores, a causal relationship between IES and 3-month favorable outcome is very likely, though our study was not randomized. Beyond the beneficial effect of the IES, our IVT control group also presented with high rates of favorable outcome at 3 months (50%) in comparison with previous reports, which ranged from 35% to 49% in the MCA location.

All the series presented above suggested that an early arterial recanalization is a significant end point in acute stroke. However, it should not be considered alone because the favorable outcome in our setting could not be explained by the time from symptom onset to IVT, which was longer than that in other studies. We believe this is secondary to a combination of the geographic dif-
ficulty presented by our rural catchment area and the relatively long door-to-needle time in this study.5,6,9,11,16,20 The strict patient-selection criteria based on the location-weighted imaging system (ASPECTS) systematically used in our center for all patients might be another additional explanation. It may be that the parenchymal extension of the ischemic lesion could have been underestimated on CT in most of the previous studies, increasing worse outcome and hemorrhagic complication rates. Another potential factor influencing the favorable clinical outcome in our setting could be related to the use of the Solitaire FR stent device, which achieved an MCA recanalization rate of 94%. The stent was easily navigated to the occlusion point, and its deployment across the thrombus was obtained at first attempt in all cases.22 No symptomatic complications (ie, intracranial hemorrhage) were observed in the IES group, suggesting a good safety profile of this stent retriever in the MCA territory. Recent reports have also demonstrated equally encouraging results in terms of the efficacy (ie, recanalization) and safety with this device.22–24

The major limitation of this study was the absence of patient randomization between IVT and IES groups, which may have biased our results. However, these 2 groups consisted of consecutive cases comparable in regard to classic prognostic factors, and the results were based on a multivariable analysis with an adjustment on confounding factors. Given the high significant association between IES and favorable clinical outcome at 3 months, it is unlikely that any residual confounding factors would alter our findings dramatically. The IVT procedure and the inclusion/exclusion criteria did not change during the study period. Our stroke center optimized with time the rapid referral of patients with stroke, which could explain the lower time from symptom onset to treatment in the second study period. However, this difference was accounted for in the analysis and was not associated with the study outcome.

**CONCLUSIONS**

Our results highlight the 24-hour and 3-month clinical benefit of an IV–endovascular strategy by using thrombectomy as a second-line therapy in patients presenting with an MCA occlusion within 4.5 hours. This strategy did not provide additional symptomatic complications. Arterial recanalization is still the most relevant goal, particularly when methodic selection of patients by using the DWI ASPECTS is performed. While the results of large multicenter trials that are evaluating the efficacy and safety of thrombectomy as a first-line therapy are pending, our strategy could represent an interesting alternative in the management of acute MCA occlusions.

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