Stent Placement in Acute Cerebral Artery Occlusion
Use of a Self-Expandable Intracranial Stent for Acute Stroke Treatment

Caspar Brekenfeld, MD; Gerhard Schroth, MD; Heinrich P. Mattle, MD; Do-Dai Do, MD; Luca Remonda, MD; Pasquale Mordasini, MD; Marcel Arnold, MD; Krassen Nedeltchev, MD; Niklaus Meier, MD; Jan Gralla, MD

Background and Purpose—Stent placement has been applied in small case series as a rescue therapy in combination with different thrombolytic agents, percutaneous balloon angioplasty (PTA), and mechanical thromboembolectomy (MT) in acute stroke treatment. These studies report a considerable mortality and a high rate of intracranial hemorrhages when balloon-mounted stents were used. This study was performed to evaluate feasibility, efficacy, and safety of intracranial artery recanalization for acute ischemic stroke using a self-expandable stent.

Methods—All patients treated with an intracranial stent for acute cerebral artery occlusion were included. Treatment comprised interarterial thrombolysis, thromboaspiration, MT, PTA, and stent placement. Recanalization result was assessed by follow-up angiography immediately after stent placement. Complications related to the procedure and outcome at 3 months were assessed.

Results—Twelve patients (median NIHSS 14, mean age 63 years) were treated with intracranial stents for acute ischemic stroke. Occlusions were located in the posterior vertebrobasilar circulation (n=6) and in the anterior circulation (n=6). Stent placement was feasible in all procedures and resulted in partial or complete recanalization (TIMI 2/3) in 92%. No vessel perforations, subarachnoid, or symptomatic intracerebral hemorrhages occurred. One dissection was found after thromboaspiration and PTA. Three patients (25%) had a good outcome (mRS 0 to 2), 3 (25%) a moderate outcome (mRS 3), and 6 (50%) a poor outcome (mRS 4 to 6). Mortality was 33.3%.

Conclusions—Intracranial placement of a self-expandable stent for acute ischemic stroke is feasible and seems to be safe to achieve sufficient recanalization. (Stroke. 2009;40:847-852.)

Key Words: ischemic stroke ■ interventional neuroradiology ■ stent ■ endovascular therapy
lyzed. Data had been collected prospectively and entered into our stroke database. Inclusion criteria for intracranial stent treatment were confirmed vessel occlusion by digital subtraction angiography (DSA), failed IAT, or contraindication to perform intravenous thrombolysis (IVT) or IAT (eg, previous cerebral infarct, surgery, or warfarin therapy). Inclusion criteria for IAT were (1) clinical diagnosis of acute stroke established by a stroke neurologist; (2) baseline NIHSS score ≥ 4, except for isolated aphasia or hemianopia; (3) exclusion of hemorrhage by cranial Computed Tomography (CT) or Magnetic Resonance Imaging (MRI); (4) vessel occlusion correlating to neurological deficit confirmed by 4-vessel angiography; (5) initiation of treatment within 6 hours of symptom onset for hemispheric stroke and within 12 hours for vertebrobasilar stroke; (6) no clinical or laboratory contraindications for IAT; (7) patients ≥ 75 years that their general condition before stroke did not advise against it; (8) informed consent of the patient or next of kin.13–15 Baseline investigations included a neurological and physical examination, assessment of stroke severity using the National Institutes of Health Stroke Scale (NIHSS), routine blood analysis, and 12-lead ECG.

**Arteriography**

Interventions were performed in the neuroangiography laboratory equipped with a biplane, high-resolution angiography system (CAS 500, Toshiba). Except for one patient who had local anesthesia, all stents were deployed under general anesthesia. After placement of a 7- to 8-Fr catheter sheath in the common femoral artery a 4-vessel DSA including all cerebral arteries was performed to assess the site of vessel occlusion and collateral circulation. Collateral flow was graded regarding to the grading system proposed by the American Society of Interventional and Therapeutic Neuroradiology/Society of Interventional Radiology: Grade 0, no collaterals visible to the ischemic site; grade 1, slow collaterals to the periphery of the ischemic site with persistence of some of the defect; grade 2, rapid collaterals to the periphery of the ischemic site with persistence of some of the defect and to only a portion of the ischemic territory; grade 3, collaterals with slow but complete angiographic blood flow to the ischemic bed occurring by the late venous phase; grade 4, complete and rapid collateral blood flow to the vascular bed of the entire ischemic territory, by retrograde perfusion.16

For endovascular therapy the 5-Fr diagnostic catheter (JB2 Valvanis, Cook) was exchanged for a 7-Fr guide catheter (Guider SoftTip, Boston Scientific), that was placed in the parent major cerebral artery (ie, internal carotid artery [ICA] or vertebral artery [VA]). The guide catheter was continuously flushed with heparinized saline (10 IU/mL).

For IAT a 2.5-Fr microcatheter (Renegade, Boston Scientific) was coaxially introduced through the guide catheter and placed directly in front of or inside the occluding thrombus. Urokinase (Medac) was administered for 60 to 90 minutes up to a dose of 1 000 000 IU. Occasionally the thrombus was gently passed with the microcatheter. Analogous, PTA, and stent placement were performed before successful passage of the occlusion and placement of a long exchange microcatheter (Transend 300 Floppy, Boston Scientific). Occlusion length and vessel diameter were measured before treatment to allow undersizing of the balloon (Gateway PTA Balloon Catheter, Boston Scientific, 1.5 to 3.5 mm diameter, 9 to 20 mm length) to avoid vessel rupture.

In all patients a self-expandable stent ( Wingspan stent system, Boston Scientific) designed for intracranial use was applied. Stents were slightly oversized to allow proper adjustment to the vessel wall (2.5 to 4.5 mm diameter, 15 to 20 mm length). The stent catheter was navigated to the occlusion site using a road map, and the stent was deployed during fluoroscopic control. Aspirin (300 to 500 mg) was administered intravenously immediately after stent placement. PTA was carried out before or after stent placement at the discretion of the operator. Recanalization result was assessed by DSA immediately after stent placement according to the Thrombolysis in Myocardial Infarction (TIMI) trial criteria: Grade 0, no recanalization; grade 1, minimal recanalization; grade 2, partial recanalization; grade 3, complete recanalization.17 Thrombus formation or residual thrombus inside the stent lumen as well as side wall irregularities corresponding to residual atherosclerotic stenosis or fixated thrombotic material between stent and vessel wall were recorded. Dissection and vessel perforation were assessed. Flow in the lenticulostriate arteries (LA) after stent placement in the middle cerebral artery (MCA) was evaluated and perfusion of major branches (eg, M2 segments, cerebellar arteries, posterior cerebral artery) after recanalization of the main vessel was assessed.

**Patient Care and Follow-Up**

After the intervention patients were transferred to the intensive care unit. Brain CT or MRI was performed in the first 24 hours after intervention as well as in case of neurological deterioration to exclude intracranial hemorrhage and to estimate brain edema. sICH was defined as clinical deterioration (4-point or greater increase in the NIHSS score or a 1-point deterioration in the level of consciousness) combined with space-occupying brain hematoma.15,18 After exclusion of hemorrhage, long-term aspirin (100 mg/d) was given and clopidogrel (75 mg/d) was added for the next 30 days. Clinical outcome was assessed at 3 months according to the modified Rankin scale (mRS).19 Outcome was stratified to “good outcome” (mRS 0 to 2), “moderate outcome” (mRS 3), and “poor outcome” (mRS 4 to 6).

Stent patency at follow-up was assessed by transcranial ultrasound (n=8), MR-Angiography (n=6), and CT-Angiography (n=4) in the first 72 hours after stent placement. Two patients died before follow-up examination of vessel patency.

**Results**

From May 2006 until July 2007, 77 patients were treated by endovascular means for acute ischemic stroke at our department. Twelve patients (7 men and 5 women, mean age 63±13 years) suffering acute cerebral artery occlusion were treated by intracranial stent placement and are included in this study. Indication for stent placement was failed IAT or MT (n=7), previous surgery, warfarin therapy, progressive stroke with preexisting brain infarct, vessel occlusion during aneurysm coiling after acute SAH, and recanalization of the intradural VA to allow IAT for thrombosis of the basilar artery (BA; n=1 each). Median NIHSS score at admission was 14 (range 5 to 38). Either CT (n=4) or MRI (n=8) was performed before treatment was initiated.

In 11 patients 1 stent was deployed, and in 1 patient 3 stents were delivered. Stent placement was feasible in all 14 stent procedures. Occlusion sites are given in the Table. All patients had collateral flow grade 2. We made the decision to place a stent in 4 patients in whom PTA had been unsuccessful. PTA was performed in a further 4 patients before and in 3 patients after stent placement. Partial or complete recanalization (TIMI 2 and 3) was achieved in 11/12 patients (91.6%). Median time from symptom onset to recanalization was 393 minutes (range 20 to 510 minutes). One dissection of a VA was visible after aspiration and PTA. No vessel perforations or dissections related to the stent placement were noted, and no SAH or sICH was detected by follow-up CT or MRI. Vessel wall irregularities were found immediately after stent placement in all patients (Figure 1).
Preservation of LA was possible in all stent placements in the middle cerebral artery (MCA). Occlusion of a major vessel branch at the site of stent placement persisted in 6 patients. Assessment of the dependent vessel branch territory by follow-up MRI or CT showed infarction in 3 of 6 patients (50%), whereas no infarction of this particular territory was noted in the remaining 3 patients. In 8 patients major branches at the site of the stent showed sufficient perfusion (Figure 1).

Thrombus formation inside the stent was found in 1 patient. It started during coil embolization of an anterior communicating artery aneurysm and continued after a stent had been deployed to recanalize the acutely occluded A1 segment. Subsequent intraarterial application of a glycoprotein IIb/IIIa receptor inhibitor (ReoPro 20 mg, Lilly) resulted in regression of luminal clot. In a further case repeated passages with the aspiration catheter and microcatheter to recanalize the BA caused a dislocation of the proximal stent markers without effect on revascularization result (Figure 2).

At 3 months follow-up 3 patients (25%) had a good outcome (mRS 0 or 1), 3 (25%) had a moderate outcome (mRS 3), and 6 (50%) had a poor outcome (mRS 4 to 6). Mortality was 33.3%.

Discussion

A recent meta-analysis of several stroke studies revealed a strong association of recanalization and good outcome after acute ischemic stroke.1 IVT has been shown to improve patient outcome and is approved by the FDA and EMEA.20 However, only a minority of patients admitted for acute stroke receive IVT.21 IAT is also effective for vessel recanalization and is supposed to achieve higher recanalization rates than IVT.13,18 On the other hand, application of thrombolytic drugs increases the risk of sICH.6,15,20 These

<table>
<thead>
<tr>
<th>Patient</th>
<th>Age (years)</th>
<th>NIHSS</th>
<th>Occlusion Site</th>
<th>Stent</th>
<th>IAT</th>
<th>MT</th>
<th>PTA Before Stent*</th>
<th>Postdilatation*</th>
<th>T to treat.†</th>
<th>T to Recan.†</th>
<th>TIMI</th>
<th>Small Perforators</th>
<th>Major Branch</th>
<th>mRS‡</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>49</td>
<td>NA</td>
<td>ACA</td>
<td>ACA</td>
<td>ReoPro</td>
<td>No</td>
<td>No</td>
<td>No</td>
<td>5</td>
<td>20</td>
<td>2</td>
<td>NA</td>
<td>Perfused</td>
<td>1</td>
</tr>
<tr>
<td>2</td>
<td>64</td>
<td>14</td>
<td>M1</td>
<td>M1</td>
<td>No</td>
<td>Yes</td>
<td>Yes</td>
<td>300</td>
<td>350</td>
<td>2</td>
<td>Perfused</td>
<td>Perfused</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>81</td>
<td>18</td>
<td>M1</td>
<td>M1</td>
<td>UK</td>
<td>No</td>
<td>No</td>
<td>285</td>
<td>375</td>
<td>2</td>
<td>Perfused</td>
<td>Perfused</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>67</td>
<td>14</td>
<td>M1</td>
<td>M1</td>
<td>UK</td>
<td>No</td>
<td>No</td>
<td>250</td>
<td>410</td>
<td>3</td>
<td>Perfused</td>
<td>Occluded</td>
<td>4</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>61</td>
<td>17</td>
<td>M1</td>
<td>M1</td>
<td>UK</td>
<td>Yes</td>
<td>No</td>
<td>165</td>
<td>340</td>
<td>2</td>
<td>Perfused</td>
<td>Occluded</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>6</td>
<td>55</td>
<td>23</td>
<td>CTO</td>
<td>M1/2</td>
<td>UK</td>
<td>Aspiration</td>
<td>Yes</td>
<td>240</td>
<td>410</td>
<td>1</td>
<td>Perfused</td>
<td>Occluded/Perfused</td>
<td>6</td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>76</td>
<td>14</td>
<td>Distal BA</td>
<td>BA/PCA</td>
<td>No</td>
<td>No</td>
<td>Yes</td>
<td>300</td>
<td>325</td>
<td>2</td>
<td>Perfused</td>
<td>Perfused</td>
<td>6</td>
<td></td>
</tr>
<tr>
<td>8</td>
<td>83</td>
<td>5</td>
<td>Distal BA</td>
<td>BA/PCA</td>
<td>No</td>
<td>No</td>
<td>Yes</td>
<td>400</td>
<td>475</td>
<td>2</td>
<td>NA</td>
<td>Occluded/Perfused</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>9</td>
<td>44</td>
<td>10</td>
<td>Middle BA</td>
<td>BA</td>
<td>UK</td>
<td>No</td>
<td>Yes</td>
<td>255</td>
<td>355</td>
<td>3</td>
<td>NA</td>
<td>Occluded/Perfused</td>
<td>5</td>
<td></td>
</tr>
<tr>
<td>10</td>
<td>61</td>
<td>38</td>
<td>VA/BA</td>
<td>VA</td>
<td>UK</td>
<td>Aspiration</td>
<td>Yes</td>
<td>350</td>
<td>510</td>
<td>2</td>
<td>NA</td>
<td>NA</td>
<td>6</td>
<td></td>
</tr>
<tr>
<td>11</td>
<td>49</td>
<td>7</td>
<td>VA/BA</td>
<td>VA/BA</td>
<td>UK</td>
<td>Aspiration, catch</td>
<td>No</td>
<td>270</td>
<td>480</td>
<td>3</td>
<td>NA</td>
<td>Occluded</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>12</td>
<td>64</td>
<td>8</td>
<td>VA/BA</td>
<td>BA</td>
<td>UK</td>
<td>Aspiration, catch</td>
<td>Yes</td>
<td>375</td>
<td>500</td>
<td>2</td>
<td>NA</td>
<td>Perfused</td>
<td>6</td>
<td></td>
</tr>
</tbody>
</table>

NA indicates not assessable; ACA, anterior cerebral artery; M1, middle cerebral artery main stem; BA, basilar artery; CTO, carotid-T-occlusion; PCA, posterior cerebral artery; VA, vertebral artery; IAT, intraarterial thrombolysis; UK, urokinase; MT, mechanical thromboembolectomy; PTA, percutaneous transluminal balloon angioplasty; *Before and after stent placement; †time from stroke onset to treatment, resp. recanalization in minutes; ‡modified Rankin scale score at 3 months follow-up.

Figure 1. 81-year-old patient presenting with acute right-sided hemiplegia and aphasia (NIHSS 18). Diffusion-weighted MR image at the level of the basal ganglia (a) depicted acute infarction in the territory of the lenticulostriate arteries (LA). Left carotid angiogram confirmed occlusion (arrow, b) of the left middle cerebral artery (MCA, M1 segment). Unsubtracted image of anterior-posterior (AP) projection documented the position of the tip of the microcatheter (c). The thrombus had been passed at the side of the LA (arrow, c). After local application of Urokinase minimal recanalization was achieved (d), considered as hemodynamically insufficient. Therefore the thrombus was passed with a long exchange microwire placed in the superior branch of left MCA. Consequently a stent was deployed at the occlusion site, and flow was immediately restored (e). Flow into LA through the stent mesh (white arrow) was achieved. Residual thrombus remained at the origin of the inferior branch (black arrow head). Follow-up MRI performed 2 days later revealed a small embolic infarction (arrow, f) in the left MCA territory but no significant increase of infarcted tissue.
Mortality (33%) was similar to former studies (32% to 40%), and a good outcome after 3 months was observed only in a quarter of the patients. However, stenting was performed as a rescue therapy in patients with major artery occlusions after failure of other techniques. This resulted in a median time from symptom onset to recanalization of 393 minutes and might explain the poor clinical outcome. Additionally, half of our patients suffered of BA occlusion, which is known to be associated with a high rate of poor clinical outcome. Three of 4 deaths in our study are in that group and were caused by aspiration pneumonia. The fourth deceased patient had a carotid-T-occlusion, which is also known to respond poorly to recanalization and often has a poor prognosis. The latter case represents the only insufficient recanalization in our study: stent deployment had been uneventful in this patient, but blood flow to the dependent brain parenchyma was obviously reduced resulting in early reocclusion. The patient died 2 days later because of increasing brain edema.

In accordance with the results of a previous study no dissection or vessel perforation was recorded attributable to stent placement itself. One dissection of a VA was noticed before stent placement related to repeated thrombectomy and PTA. Remarkably, occurrence of sICH and SAH in our patients was significantly lower than reported (none versus 14% to 50%). One explanation might be the use of a self-expandable stent compared to balloon-mounted stents investigated in prior studies. Balloon-mounted stents developed for cardiology are less flexible than self-expandable stents. Therefore, their application causes more mechanical stress to the vessel wall during the navigation into the relatively small intracranial arteries. Additionally, the necessity for stiffer microwires that have to be placed in the periphery of the cerebral arteries increases the risk of perforation, eg, all 4 patients treated with a balloon-mounted stent in the study of Sauvageau et al suffered SAH. Moreover, self-expandable stents apply a lower radial force than the inflation of balloon-mounted stents, reducing the risk of vessel rupture in case of hard thrombus or atherosclerotic stenosis. After deployment the self-expanding force persists and might improve recanalization result with time as thrombus gets resorbed. Besides the mechanical aspects of the applied stent, concomitant antiplatelet therapy in combination with thrombolysis might have increased bleeding rates in previous studies: whereas Levy et al applied glycoprotein IIb/IIIa receptor inhibitors in addition to aspirin, clopidogrel, heparin, and thrombolitics, a glycoprotein IIb/IIIa receptor inhibitor was given only in one of our patients who had not received thrombolysis. Additionally, heparin was not used in our patients aside from catheter flushing. The combination of UK, aspirin, and clopidogrel did not cause sICH in our small patient group.

However, local platelet activation attributable to thrombus, vessel wall injury, and the stent itself remains a major concern. Immediate application of aspirin after stent placement followed by a combination therapy of aspirin and clopidogrel for the next 4 weeks after exclusion of hemorrhage seems efficient to avert stent occlusion. Keeping in mind the higher bleeding rates reported in studies using a combination of aspirin, clopidogrel, glycoprotein IIb/IIIa receptor inhibitor, and thrombolitics as well as the
unfavorable results of the Abciximab in Emergency Treatment of Stroke Trial (AbESTT),26 we advocate the use of glycoprotein IIb/IIIa receptor inhibitors only in case of acute thrombus formation after stent placement. Using this regimen, we have observed no recurrences or embolic events caused by the stent during follow-up.

From our point of view important side branches like the LA can be preserved if the thrombus is passed on the ipsilateral side by the microwire and the stent. Stent expansion will fixate the thrombus at the contralateral wall. This technique was successfully performed in all 5 of our patients suffering MCA occlusion and can be translated to the BA and P1 segments with their perforating arteries as well. However, occlusion of major vessel branches (eg, M2 segment, superior cerebellar artery) at the site of stent placement persisted in 6 patients. Remarkably, no infarction occurred in the dependent vessel territory in half of those patients, pointing to a sufficient collateral circulation. In 3 patients follow-up MRI or CT revealed infarcts in the dependent vessel territory. These infarcts were apparent to some extent at MRI before the interventional treatment, and it remains uncertain whether they are related to the primary occlusive disease or to the stent placement.

In our experience care has to be taken if the deployed stent has to be passed repeatedly with other devices. As long as the stent is not covered by neointima, devices might get caught in the stent struts with subsequent complications (ie, stent dislocation or deformation [Figure 2], device rupture, or failure to remove the device). Hence stent placement was performed to reestablish sufficient cerebral blood flow with as little mechanical manipulation as possible, additional postdilatation was performed in only 25% of our patients.

Before the introduction of stents for the treatment of atherosclerotic stenosis of intracranial vessels, PTA performed alone for acute ischemic stroke yielded some success.27–29 However, when treating thrombus rather than atherosclerotic stenosis, recollection attributable to thrombaggretation and thrombus expansion might occur.7 In our study all PTAs performed for vessel recanalization failed to establish sufficient flow.

Limitations
Because of the retrospective nature of the study, the inhomogeneous and small patient collective, and the different therapies used in addition to the placement of an intracranial stent, results have to be interpreted with caution.

Summary
This study illustrates the feasibility of intracranial stent placement for acute stroke treatment. Recanalization rate was high, and the rate of major complications was low. However, clinical outcome, at least partially influenced by patient selection, was poor. Further controlled randomized trials to prove its safety and efficacy in a larger number of patients are needed.

Disclosures
None.

References


Stent Placement in Acute Cerebral Artery Occlusion: Use of a Self-Expandable Intracranial Stent for Acute Stroke Treatment
Caspar Brekenfeld, Gerhard Schroth, Heinrich P. Mattle, Do-Dai Do, Luca Remonda, Pasquale Mordasini, Marcel Arnold, Krassen Nedeltchev, Niklaus Meier and Jan Gralla

Stroke. 2009;40:847-852; originally published online January 29, 2009; doi: 10.1161/STROKEAHA.108.533810
Stroke is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
Copyright © 2009 American Heart Association, Inc. All rights reserved.
Print ISSN: 0039-2499. Online ISSN: 1524-4628

The online version of this article, along with updated information and services, is located on the World Wide Web at:
http://stroke.ahajournals.org/content/40/3/847

Permissions: Requests for permissions to reproduce figures, tables, or portions of articles originally published in Stroke can be obtained via RightsLink, a service of the Copyright Clearance Center, not the Editorial Office. Once the online version of the published article for which permission is being requested is located, click Request Permissions in the middle column of the Web page under Services. Further information about this process is available in the Permissions and Rights Question and Answer document.

Reprints: Information about reprints can be found online at:
http://www.lww.com/reprints

Subscriptions: Information about subscribing to Stroke is online at:
http://stroke.ahajournals.org/subscriptions/