Clinical Experience of Thromboembolic Complications of Coil Embolization for Intracranial Aneurysms with Literature Review

Sung-Kyun Hwang · Sung-Hak Kim

Department of Neurosurgery, College of Medicine, Ewha Womans University

Introduction

The endovascular treatment of an intracranial aneurysm by using Guglielmi detachable coils (GDC) was first introduced by Guglielmi et al in 1991. Since then, GDC treatment has undergone rapid change, and recent advanced techniques make it possible to improve anatomic and clinical outcomes. Although this treatment has become an accepted alternative to surgery, thromboembolic events still constitute the main complication.

The development of new materials and advances in vascular catheterization techniques that have made it possible to treat various diseases via an endovascular approach in last decade. With the growing use of endovascular procedures in neurosurgical practice, adequate
Knowledge of basic pathophysiological principals is important.

In this article, we review our thromboembolic complications associated with endovascular procedures, and describe our clinical outcomes with literature review.

**Patients and Method**

Between August 2004 and August 2008, we performed endovascular embolization with GDCs for 160 patients with 176 cerebral aneurysms including ruptured and unruptured aneurysms at our hospital. Sixty-six patients were male and 94 were female. The age of patients ranged from 22 to 82 years (average, 58.8 years). All ruptured aneurysms were treated within 2 weeks. In this study we retrospectively evaluated this group with regard to complications rates and outcome.

The locations of the aneurysms were the internal carotid artery in 23, anterior communicating artery in 8, anterior choroidal artery in 7, posterior communicating artery in 43, middle cerebral artery in 40, basilar tip in 9, vertebral and basilar artery in 10 patients. One hundred sixty-one aneurysms were small (diameter lesser than 10mm), and thirteen were large (diameter 10–25mm), two were giant (diameter more than 25mm). One hundred forty-two aneurysms had small necks (neck diameter 4mm), and thirty four had wide necks (neck diameter 4mm). The method used for such measurements has been published previously\(^\text{[24]}\)\(^\text{[9]}\)\(^\text{[13]}\)\(^\text{[14]}\)\(^\text{[20]}\)\(^\text{[22]}\).

### Table 1. Patient with thromboembolic complications

| No | Age (years)/Sex | Site         | Size (mm)  | Hunt and Hess grade at presentation | Antiplatelet drug response | Comment                              | GOS |
|----|----------------|--------------|------------|-------------------------------------|---------------------------|--------------------------------------|-----|
| 1  | 48/M           | ACOM         | 8×3.5      | 4                                   | Complete                  | Recanalization by intraarterial thrombolysis | 2   |
| 2  | 50/F           | ICA          | 2×3        |                                      | Complete                  | Stent induced thrombosis            | 5   |
| 3  | 45/F           | MCA          | 4×5        | Unruptured                          | Partial                   | Hemorrhagic transformation           | 5   |
| 4  | 44/F           | MCA          | 6.5×7      | Unruptured                          | Partial                   | Hemorrhagic transformation           | 5   |
| 5  | 41/F           | MCA          | 5×7        | Unruptured                          | Complete                  | Transient dysphasia                  | 5   |
| 6  | 65/M           | PCOM         | 2×4        | Unruptured                          | None                      | Infarction                           | 2   |

ACOM: anterior communicating artery aneurysm, ICA: internal carotid artery aneurysm, MCA: middle cerebral artery aneurysm, PCOM: posterior communicating artery aneurysm, GOS: Glasgow outcome scale.

![Fig. 1. No. 1 case. A: Computed tomography shows subarachnoid hemorrhage in basal cistern & interhemispheric cistern with diffuse brain swelling. B: Enhanced computed tomography shows aneurysmal sac at anterior communicating artery.](image)

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Results

Patients were included in this category if they had clinical symptoms of cerebral ischemia, either angiographic or clinical, of thromboembolism. Clots were angiographically noted but, if there was no clinical evidence of cerebral ischemia, such events were not recorded as complications. Of 160 patients, only six patients had the thromboembolic complications during or after procedures (3%, Table 1). Two of them suffered fully recovered transient ischemic attack after procedure, which might be stent and catheter related complication. They did not have a suffered a watershed stroke from hypotension/hypoperfusion during a procedure. Another two patient had a ischemia and delayed hemorrhagic transformation, which were related to use of antiplatelet drug during or after procedures. The other two patients had thromboembolic complications during procedure which were relieved by intraarterial fibrinolytic management. None of patients died of these thromboembolic complications.

Discussion

Thromboembolic complications are reported in 5–10% of patients treated endovascularly\(^2\)\(^3\)\(^4\)\(^5\)\(^6\)\(^7\)\(^8\)\(^9\)\(^10\)\(^11\)\(^12\)\(^13\)\(^14\)\(^15\)\(^16\)\(^17\)\(^18\), though permanent deficit is less frequent, with rates of 1–5%\(^19\)\(^20\).

Associations between the development of thrombosis, the size of the catheter/introducer, and the duration of the procedure have been established. Thromboembolic complications of coiling are most commonly recognized as distal emboli\(^14\)\(^15\) on check angiography and less commonly occur with a filling defect at the coil-ball/parent-artery junction\(^21\). However, we had an increased prevalence of local thrombosis. Large aneurysm diameter and coil protrusion are independent risk factors for post-
procedural thromboembolic events. Stroke, usually from thrombus in the parent vessel with downstream embolization, remains a significant problem. The actuarial risk of stroke associated with aneurysm coiling has been found by one group, in a retrospective study that only looked at good-grade patients, to equal 3.8%6). In addition to clinically recognizable clots and strokes, silent infarcts are often detected on postprocedure diffusion-weighted magnetic resonance imaging. There is evidence that these events are quite common12). The outcome of the International Subarachnoid Aneurysm Trial was encouraging to neuroendovascular enthusiasts. There is, unfortunately, no prospectively obtained published data on neuroendovascular technical problems. It is important that endovascular specialists share their poor outcome data, not only to educate their colleagues on the pitfalls of endovascular treatment, but to also improve the level of care.

There are many possible sources for embolic events during GDC treatment of cerebral aneurysms, including friable plaques and iatrogenic dissection in the parent vessels, air bubbles, and thrombus or fresh clots within aneurysms and catheters18). On the other hand, the location of the lesion depends on both the flow patterns in the patient’s brain and other unresolved factors. The brain border zone is a favored destination for microemboli because this area has an inadequate blood supply, and once the blood supply is disturbed by microemboli, this area might fall to infarction19). Thromboembolic complications can be initially silent, becoming symptomatic only when the infarct enlarges.

Treatment of thrombus formation may be local (mechanical clot extraction and/or disruption)523), and abciximab21, or a combination. Treatment of thrombotic complications in ruptured aneurysms requires a fine balance between maintaining intra-aneurysmal occlusion (to which coil packing and thrombosis contribute)3) and lysis of the thrombus in the parent artery. Its action is
prolonged and, though reduced, it may persist as long as a week after administration. In contrast to heparin, abciximab is not easily reversed. Despite these theoretic effects, abciximab has been used for carotid and vertebro-basilar rescue and for the treatment of acute stroke, without a substantial increase in intracerebral hemorrhage. Therefore, abciximab does not appear to have intracranial risk on par with that of thrombolytic agents.

This was observed at the time of diagnostic angiography immediately before emergency coiling. Abciximab immediately cleared the soft basilar thrombus, but the focal thrombus persisted at the origin of the superior cerebellar artery. This clot did not respond to attempted mechanical disruption by using the guidewire, leading us to conclude that this was hard, atheroembolic material dislodged from the vertebral artery dissection. We suggest that an infusion may potentially confer an additional and unnecessary risk of hemorrhage. Postprocedural anticoagulation can be effectively maintained with heparin followed by aspirin and/or clopidogrel. Technical complications associated with the use of GDCs include aneurysmal perforation and rupture, parent artery occlusion, cerebral embolism, coil migration, vasospasm, and hemodynamic ischemia with inadequate collateral reserve. The largest contributor to complications of GDC treatments is iatrogenic brain ischemia caused by thromboembolic events. This may be caused by thrombus form athrombosis in the aneurysm sac, or from herniation of the coils into the parent vessel. Although clinical thromboembolic complications have been reported to occur in a range of 1.0% to 28%, the complications in these studies were defined as new focal deficits, a change in mental status, or abnormalities at postprocedural computed tomography or conventional magnetic resonance imaging. Rordorf et al suggested that thromboembolic events related to GDC techniques may be more common than what has been reported: in fact, silent embolism occurred in 61% of the 14 cases in their study. GDC treatment is effective with an acceptable morbidity rate for small aneurysms with a small neck owing to the high initial success rate and low mortality and morbidity rates. In wide-neck and large aneurysms, however, the combined use of the balloon-assisted technique enables a denser packing of aneurysms and markedly improves the anatomic results after the treatment. Some authors report that the occurrence of thromboembolic events with the use of the balloon-assisted technique is not different from that of the conventional GDC technique. However, this technique requires sophisticated handling, frequent use of guidewires and microcatheters, the need for temporary occlusion of the parent vessel, and repeated inflation and deflation of the balloon. Use of the balloon-assisted technique is theoretically a risk factor for thromboembolic complications, because the risk of significant emboli will likely increase with increasing procedural complexity, potential sites for the generation of microemboli increase, and longer procedure time. On the other hand, the process of intra-aneurysmal clot organization in large aneurysms is delayed and incomplete, and it may cause delayed postprocedure ischemic events.

Bendzus et al reported that a silent embolism related to diagnostic angiography occurred in 26% of their cases. Their embolic lesions were located in the distal vascular territory of small cortical, subcortical, or perforating vessels. Whenever possible, clot disruption by fragmentation is carried out before and in combination with chemical thrombolysis: clot fragmentation alone is never used as the sole solution. The purpose of fragmentation is twofold: to establish flow beyond the thrombus, permitting passage of the fibrinolytic drug, the heparin, and the intrinsic fibrinolytic factors into the occluded area, and to increase the surface of the clot accessible to the drug. The results of combination of clot fragmentation and fibrinolysis suggest that fragmentation and superselective drug infusion may improve the rate of recanalization. The rate of recanalization indicates that the early administration of a fibrinolytic drug increases the chances of favorable clot dissolution. Spontaneous thrombolysis
has been estimated to be 20% during the first 24 hours and 80% within the first week after ictus.

We believe that fibrin or thrombus formation was induced at the tip of the catheter or around the coil. However, on some occasions, the occlusive material may have been an embolus from a preexisting thrombosis within the aneurysmal sac that was dislodged during coil maneuvering. These emboli originate from thrombogenic material a long time before anticoagulant therapy is given, resulting in clots with different characteristics. Another factor that must be considered is the different vulnerability to ischemia found in different regions of the brain. In our opinion, the risks of clotting do not correlate with the size of the aneurysm but rather with its location and with the extent of experience of the doctor in charge.

Most thromboemboli in our series occurred in the middle cerebral artery territory, which was also the most frequent location of the aneurysms. Thrombosis is more likely in cases in which coils protrude from the aneurysm into the lumen of the main artery. Complications related to fibrinolytic therapy were seen in three patients. Two suffered aneurysmal rebleeding, in both instances the patients underwent embolization during the acute phase of an subarachnoid hemorrhage.

**Conclusion**

In endovascular procedures, arterial injury, and the use of catheter, contrast agents, and implanted devices with thrombogenic potential place patients at risk for thrombosis and embolization. Extensive research has been performed to elucidate the pathophysiological features underlying thrombosis associated with endovascular procedures.

Although permanent clinical sequelae are rare, the high rate of thromboembolic events suggests that alterations in the technique, such as the addition of antplatelet agents and development of new embolic materials that could be packed by means of simple methods, should be considered. Experience, attention to procedural details, proper patient selection, and measures to reduce the rate of symptomatic and silent embolisms may permit safer aneurysm treatment in the future.

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