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Resolution of Mass Effect and Compression Symptoms following Endoluminal Flow Diversion for the Treatment of Intracranial Aneurysms

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ABSTRACT

BACKGROUND AND PURPOSE: Alleviation of aneurysm induced mass effect has been difficult with both conventional endovascular and surgical techniques. Our aim was to study the efficacy of endovascular flow modification on aneurysm-induced mass effect and compression syndrome, as demonstrated by cross-sectional imaging studies and clinical follow-up.

MATERIALS AND METHODS: Thirty aneurysms larger than 10 mm were treated by flow diversion alone and previously had undergone pre- and posttreatment cross-sectional imaging. Pretreatment MR imaging or contrast CT, follow-up angiography at 6 months, and follow-up MR imaging studies between 6 and 18 months were retrospectively analyzed. The neurologic and neuro-ophthalmologic statuses of all patients were recorded before treatment and at the time of follow-up cross-sectional imaging.

RESULTS: At 6 months, 28 aneurysms were completely occluded, 1 had a neck remnant, and 1 had residual filling on angiography. Between 6 and 18 months, 3 aneurysms decreased in size and 27 completely collapsed as demonstrated on MR imaging. Before treatment, 6 patients had vision loss, 10 had double vision due to a third or sixth nerve palsy or both, and 1 had hemiparesis due to brain stem compression. On MR imaging follow-up, vision loss had either improved or resolved in all except 1 patient, double vision had resolved completely (7/10) or partially (3/10), and the patient with brain stem compression became asymptomatic. There was no bleeding observed in this series. One parent artery thrombosis resulted in a major infarct.

CONCLUSIONS: Endovascular flow diversion is a highly effective technique for resolving radiologic mass effect and clinical compression syndromes.

ABBREVIATIONS: FD = flow diverter; GPIIb/IIIa = glycoprotein IIb/IIIa; PED = Pipeline Embolization Device; PUFS = Pipeline for Uncoilable or Failed Aneurysms

Large and giant intracranial aneurysms pose an increased health risk due to a high incidence of rupture1-2 and resultant neurologic deficits caused by compression of neighboring neural structures.3 Successful treatment of aneurysms presenting with mass effect is limited by high surgical morbidity4,5 and low endovascular efficacy, due to a high rate of incomplete occlusion and recurrence.5,6 Parent artery occlusion is effective but restricted to aneurysms with sufficient collateral circulation.5,7 Surgical bypass can potentially compensate for the lack of collaterals but adds substantial morbidity.8,9 Modification of flow within the parent artery is a method that attempts to reduce intra-aneurysmal circulation, leading to aneurysm thrombosis. Reports published to date demonstrate a very high rate of complete and stable occlusion of aneurysms that were considered difficult or impossible to treat by conventional endovascular techniques.12-17 Collapse of large and giant aneurysms has also been observed.13 The purpose of this work was to analyze the effect of flow modification on the radiologic mass effect and clinical compression syndromes induced by large and giant aneurysms.

MATERIALS AND METHODS

Ninety-eight aneurysms in 81 patients were treated at a single center with flow diversion by using either the PED (Covidien/ev3, Irvine, California) (97 aneurysms) or the Silk stent (Balt Extrusion, Montmorency, France) (1 aneurysm) from 2006 to 2011. All subjects were treated under institutional review board–approved protocols as part of the Pipeline Embolization Device for the Intracranial Treatment of Aneurysms Trial study,15 the PUFS study,18 or the Reconstructive Aneurysm Treatment study (un-
published); informed consent was obtained from all subjects. To properly evaluate the effect of flow diversion alone, we excluded from the current analysis aneurysms that had coils in their sacs because they were either previously treated or packed during the FD treatment. For the purpose of this study, the aneurysms that were selected had a diameter larger than 10 mm, were treated by flow diversion alone (ie, without coils), and previously had the following imaging studies: cross-sectional imaging either by MR imaging or contrast CT before treatment; follow-up angiography at 6 months; and T1WI, T2WI, and time-of-flight MRA between 2 and 18 months. Twelve aneurysms were studied by MR imaging at 3 months, and all, between 12 and 18 months after treatment. All these patients had a neurologic and/or neuro-ophthalmologic evaluation before treatment and at the time of follow-up cross-sectional imaging. Clinical and imaging records were evaluated retrospectively.

RESULTS
Thirty aneurysms were identified in 27 patients.

Indications. One patient was treated for an incidentally discovered aneurysm. Twelve patients presented with headaches, and 17, with neurologic symptoms related to their aneurysms. Among those, 1 patient had hemiparesis due to brain stem compression, 6 had vision loss of different degrees related to optic nerve or chiasmal compression, and 10 had diplopia related to either third (n = 3) or sixth (n = 2) cranial nerve palsy or both (n = 5).

Aneurysm Characteristics
Twenty-nine aneurysms were located on the ICA, and 1, on the basilar trunk. ICA aneurysms either involved the cavernous (n = 10) or the paraophthalmic (n = 16) segment of the ICA or arose at the origin of either the superior hypophyseal artery (n = 1) or the posterior communicating artery (n = 1). Sixteen aneurysms had a diameter between 10 and 20 mm, 12 measured 20–30 mm, and 2 were ≥30 mm in diameter. The mean aneurysm size was 18.9 mm, and the mean neck size was 9.7 mm. Three of the 30 aneurysms were fusiform. Two aneurysms of the ICA demonstrated calcifications in their wall on CT.

Procedural Technique
All patients were pretreated with oral aspirin (either 100 or 300 mg/day) and clopidogrel (75 mg/day) for a minimum of 4 days before the procedure. All subjects were intravenously heparinized during the endovascular procedure to reach an activated clotting time of twice their baseline level. No additional heparin was used following the procedure, except in cases of intraprocedural thrombotic complications.

For flow diversion, the PED (Covidien/ev3) was used in 29 aneurysms, and a combination of a LEO and a Silk stent (Balt Extrusion), in 1 case. A single-layer FD was used in 1 case (the aneurysm was located at the posterior communicating artery origin), and the aneurysm neck was covered with 2 layers in 18 cases. Three coaxial layers were used in 8 cases, and 4 layers, in 1 case. A long FD construct was built to cover the lengths of fusiform aneurysms by using 5 pieces in one case and 15 pieces in another case. All patients were maintained on double antiplatelet treatment for 1.5–3 months. Steroids were not routinely administered perioperatively, but subjects with aggravated postprocedure symptoms received methylprednisolone, 2 × 250 mg/day for 2–3 days, with a gradually tapering dose for an additional 10–14 days.

Complications
As a technical complication, incomplete vessel wall apposition was observed in 1 case and was corrected with balloon dilation. In-stent thrombosis occurred during the procedure in 2 cases and was resolved by thrombolysis with application of both tissue plasminogen activator and GP IIb/IIIa inhibitor in one case and GP IIb/IIIa inhibitor only in another case. Subacute occlusion of the parent artery occurred in 1 case 2 days postprocedure, resulting in transient hemiparesis. This was not treated due to good collateral circulation, and it later recanalized spontaneously. Delayed occlusion resulting in hemiparesis was found in 1 case 3 months after treatment. Overall, transient neurologic deficit occurred in 2 patients, and a permanent deficit occurred in 1 patient.

Angiographic Follow-Up
All aneurysms were followed by angiography at 6 months. In addition, 13 aneurysms underwent follow-up angiographies at 1

FIG 1. Complete angiographic occlusion of a paraophthalmic aneurysm and aneurysm collapse following FD treatment. A, Large paraophthalmic aneurysm (arrow) causing optic nerve neuropathy on DSA, lateral view. The patient’s visual acuity was 0.4. B, Complete angiographic occlusion of the aneurysm and reconstruction of the parent artery (arrow) 6 months after treatment with a PED. The patient’s visual acuity was 0.5. C, The same aneurysm demonstrated by T2WI with mass effect (arrow) and mixed signal intensity. D, Lack of signal void within the aneurysm on T2WI in the same patient 2 months later indicates aneurysm thrombosis. The size of the aneurysm and related mass effect remain unchanged (arrow). Optic nerve neuropathy progressed; the patient’s visual acuity was 0.1. E, T2WI in the same patient demonstrates aneurysm collapse and elimination of mass effect at 18 months after treatment. The patient’s visual acuity at this point was 0.9.
The patient's visual acuity was 0.2. D related to aneurysm wall calcification, and the aneurysm size remained unchanged. The patient's visual acuity deteriorated slightly (0.1). (Fig 2).

was found on the T2-weighted images within the aneurysm sac was detected by the time-of-flight MRA, but high signal intensity study, in aneurysms that did not yet fully collapse, no flow signal was detected by the time-of-flight MRA, but high signal intensity was seen on MR imaging, the patient's visual acuity slightly deteriorated. Seven were completely cured, and 3 had vision improved from as low as finger counting only to quantifiable vision of 0.6 and from 0.3 to 0.8. In 1 case in which the aneurysm worsened from as low as finger counting only to quantifiable vision of 0.6 and from 0.3 to 0.8. In 1 case in which the aneurysm was incompletely occluded on angiography and did not collapse as seen on MR imaging, the patient’s visual acuity slightly deteriorated from 0.2 to 0.1 (Fig 2). All 10 subjects who had presented with diplopia improved: Seven were completely cured, and 3 had minor diplopia at the time of the last MR imaging follow-up. Two of the minor diplopia cases were related to a residual partial third nerve palsy, and 1 was caused by residual dysfunction of both the third and sixth nerves. Hemiparesis in 1 patient presenting with brain stem compression was relieved completely by 6 months after treatment.

DISCUSSION

Large and giant aneurysms represent a challenge for both surgical and endovascular approaches. Such aneurysms often present with mass effect and corresponding compression syndrome on the neighboring neural tissue. Therefore, the goal of treatment is not only to prevent rupture but also to reduce the pulsating mass and eliminate the associated mass effect. Unfortunately, this goal is difficult to achieve by conventional endovascular techniques. Packing of the aneurysm with soft coils can reduce the pulsation on the surrounding tissue only if the neck of the aneurysm is completely sealed. Because most of these aneurysms typically have a broad neck, such complete sealing is very difficult if not impossible to achieve, resulting in the low observed efficacy of coil packing in relieving compression symptoms and the high rate of recanalization. Embolization by using liquid embolics may provide better long-term stability; however, such embolization turns a soft pulsating mass into a hard volume that, even if completely occluding the aneurysm, transfers the pulsation of the parent artery to the surrounding tissue. Surgical treatment is considered superior in cases of compression syndromes because it is capable of immediate decompression of the neural tissue. However, surgery is made difficult by the large mass of the aneurysm, often requiring temporary clipping or proximal balloon occlusion and necessitating induced hypothermia or even cardiac arrest during surgery. Despite these difficulties, the efficacy of direct surgery in eliminating cranial nerve symptoms remains limited. Many of these aneurysms are repaired by bypass surgery rather than by clipping. While this technique can be performed without interrupting antegrade blood flow, the rate of morbidity-mortality associated with the method is considerable. The limited success of conventional techniques justifies the search for safer and more effective treatment options.
Flow diversion by using high-mesh-attenuation intraluminal endoprostheses, the newest advance in endovascular aneurysm therapy, has been recently introduced.13-16,29 The primary goal of diverting flow away from the aneurysm cavity is to allow the aneurysm to thrombose and the intima of the parent artery to cover the neck of the aneurysm once flow through the neck has ceased. Early reports suggest that in addition to better primary occlusion rates and improved long-term stability, this technique is capable of effectively eliminating mass effect and related compression symptoms.13 More recently, Piano et al30 reported shrinkage of the aneurysm following flow diversion in 61% of cases that were available for assessment.

In contrast to these favorable results, temporary aggravation of compression-related cranioptathies may occur.14 In addition, delayed rupture of FD-treated large and giant aneurysms has been reported.31,32 In a voluntary self-audited survey conducted by the European Society of Minimally Invasive Neurologic Therapy, delayed rupture was found in 0.96% of 1421 aneurysms but in 2.1% of large and giant lesions.33 Flow diversion is further complicated by the necessity of aggressive antithrombotic treatment and potential thrombosis of the parent artery in the case of insufficient antiaggregation.

The present report focuses on the radiologic and clinical effects of endoluminal flow alteration on large and giant aneurysms treated solely by the FD technique. Cases were retrospectively selected from a consecutive series of 98 aneurysms that were treated with FD for various indications in our institution. Only aneurysms that were larger than 10 mm and treated without coils were included. The 6-month and 1-year angiographic occlusion rates were similar to those reported in other flow diversion studies,13-15 and clinical outcomes closely matched radiologic findings.

On posttreatment MR imaging follow-up, 27 of 30 aneurysms (90%) collapsed completely. In another 3 cases (10%), the aneurysm shrank but did not fully disappear. In 2 of those cases, image analysis disclosed a heavily calcified aneurysm wall (Fig 2C, -D). The rigid, calcified wall of the aneurysm may prevent or delay collapse even if there is no flow inside the sac. In 1 case, some shrinkage of the sac was first detected 8 months after treatment even though the aneurysm was completely occluded on angiography at 6 months. Despite every effort (including aggressive steroid treatment), the patient’s visual acuity, which deteriorated 8 weeks after treatment, only started to improve at this time. The visual acuity finally improved from the pretreatment status of 0.4–0.25 to 0.8–0.8. In the other case, the role of wall calcification could not be evaluated because complete aneurysm occlusion was not achieved and the aneurysm had residual flow at 6-month angiography. In this case, no aneurysm shrinkage was detected and the sac did not lose its mixed high T1WI-T2WI signal intensity. This patient’s visual acuity slightly deteriorated from 0.2 to 0.1 (Fig 2).

In the third case, the ICA was occluded following bridging of a giant fusiform carotid cavernous aneurysm by using a long construct of PEDs. In this case, the size of the aneurysm decreased but it did not fully disappear, though the associated double vision had resolved.

In 27 cases, complete angiographic occlusion was achieved with the exception of a very small neck remnant. The aneurysms completely collapsed and became isointense. In all of these cases, the relevant cranial nerve compression symptoms either improved or resolved completely. All subjects in this series were treated in a single session, and none have been retreated.

FDs are thought to be effective by reducing blood flow through the aneurysm orifice. Similar to other stents, the implanted FD induces neointimal growth over its surface. Once blood flow entering and exiting the aneurysm has been substantially reduced, the neointimal layer may completely cover the orifice and seal the aneurysm from the parent artery. Reduced flow within the sac initiates thrombosis. During the acute stage of thrombosis, the aneurysm mass may increase in size, further increasing the associated mass effect34,35 and possibly leading to temporary aggravation of related clinical symptoms, as observed in 4 of 27 patients in our series, thereby necessitating the application of high-dose corticosteroids. The reduction of mass effect is expected by cessation of aneurysm pulsation and collapse of the mass of the aneurysm, which later requires thrombus organization that may take weeks to months following complete cessation of intra-aneurysmal flow. In our study, aneurysm shrinkage was seen as early as 3 months and as late as 8 months after treatment. Lack of aneurysm filling on angiography was paralleled by a reduction of aneurysm size on MR imaging in most (but not all) cases. Clinical symptoms, however, did not improve until the aneurysm started shrinking and losing its mixed signal intensity on MR imaging. Angiographic occlusion after FD treatment indicates redirection of flow into the parent artery but does not demonstrate completion of intra-aneurysmal thrombosis.

There were no delayed hemorrhagic complications reported in this series. Acute in-stent thrombosis occurred in 2 cases, both of which were successfully resolved by thrombolysis. The parent artery occluded during the subacute phase in 1 case and later recanalized completely. In-stent thrombosis was attributed to the patient’s noncompliance in taking antiplatelets. An ICA occluded in a delayed fashion and could not be recanalized, which was attributed to premature cessation of clopidogrel 3 months after treatment of a giant and (more important) fusiform aneurysm. Such aneurysms likely require much longer times to endothelialize compared with berry aneurysms.

CONCLUSIONS
In this small series, FD treatment was highly effective in eliminating radiologic mass effect and resolving related clinical symptoms. Clinical improvement is expected only after aneurysm shrinkage or collapse. Angiographic occlusion indicates a lack of flow through the aneurysm orifice, but complete thrombosis may take considerably longer. MR imaging studies are strongly suggested in the follow-up of FD-treated aneurysms, especially when aneurysm-associated symptoms do not resolve.

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