Aneurysms of the cavernous segment of the internal carotid artery usually become symptomatic by mass effect on the adjacent cranial nerves, causing ophthalmoplegia and, sometimes, decreased vision or trigeminal neuralgia.1 Rupture of an adjacent cranial nerves, causing ophthalmoplegia and, some- possible,4 the preferred treatment is endovascular, either by aneurysm occlusion with detachable coils was performed in 841 an- and 932 aneurysms were treated by endovascular techniques: selective treatment at our institution. Surgery was performed in 948 aneurysms, and 932 aneurysms in 51 patients were located in the cavernous sinus. All 51 ane- ysm were treated by endovascular methods: selective occlusion with coils in 22 and parent-vessel occlusion in 29 (5 after bypass surgery). Of the 51 aneurysms in the cavernous sinus, 41 were symptomatic, 9 were additional to another symptomatic aneu- ysm, and one was an incidental finding. Presentation of 41 symp- tomatic cavernous sinus aneurysms was mass effect (ophthalmo- plegia, decreased vision) in 28 (68%), CCF in 10 (24%), SAH in 2 (5%), and epistaxis in one (2%).

During the study period, one patient with a ruptured cavernous sinus aneurysm causing CCF died of intracranial hemorrhage shortly before scheduled treatment. The present study group consists of 11 patients with ruptured cavernous sinus aneurysms causing CCF: 10 patients who were treated and the single patient who died before treatment (Table). The patients included 8 women and 3 men, with a mean age of 61.6 years (range, 27–77 years).

CCFs were classified according to flow rate into 3 categories: high, intermediate, or low flow. In high-flow CCFs, all the blood from the internal carotid artery entered the fistula without filling of intracra- nial vessels (Fig 1). In intermediate-flow CCFs, both the fistula and intracranial vessels received blood from the internal carotid artery (Fig 2), and in the low-flow CCFs only slow and sluggish filling of the cavernous sinus was apparent.

Treatment
Endovascular treatment was performed under general anesthesia on a biplane angiographic unit (Philips Integra V3000 Neuro; Philips

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Clinical presentation, treatment and outcome in 11 patients with ruptured cavernous sinus aneurysms causing CCF

<table>
<thead>
<tr>
<th>Patient No./Gender/Age (y)</th>
<th>Presentation</th>
<th>Diagnosis</th>
<th>Principal Venous Drainage</th>
<th>Test Occlusion</th>
<th>Treatment</th>
<th>Outcome; Duration of Follow-up in Months</th>
</tr>
</thead>
<tbody>
<tr>
<td>1/F/77</td>
<td>Bilateral exophthalmus and ophthalmoplegia, vision 0.1 bilateral, pulsatile bruit</td>
<td>High-flow left CCF</td>
<td>Bilateral superior ophthalmic vein, minor cortical venous drainage</td>
<td>–</td>
<td>Coiling aneurysm with balloon protection, carotid artery patent</td>
<td>Complete recovery; 9 mo</td>
</tr>
<tr>
<td>2/F/75</td>
<td>Left exophthalmus and ophthalmoplegia, pulsatile bruit, decreased vision</td>
<td>Intermediate-flow left CCF</td>
<td>Left superior ophthalmic vein, no cortical venous drainage</td>
<td>–</td>
<td>Coiling aneurysm ×2</td>
<td>Complete recovery; 72 mo</td>
</tr>
<tr>
<td>3/F/60</td>
<td>Left hemiplegia and aphasia, bilateral exophthalmus and ophthalmoplegia, decreased vision, pulsatile bruit</td>
<td>High-flow right CCF</td>
<td>Bilateral superior ophthalmic vein, major cortical venous drainage</td>
<td>+</td>
<td>Coiling aneurysm with balloon protection, carotid artery patent</td>
<td>Complete recovery; 6 mo</td>
</tr>
<tr>
<td>4/F/61</td>
<td>Bilateral exophthalmus and ophthalmoplegia, decreased vision, pulsatile bruit</td>
<td>Intermediate-flow left CCF</td>
<td>Bilateral superior ophthalmic vein, minor cortical venous drainage</td>
<td>+</td>
<td>Balloon occlusion aneurysm, later coiling pseudo-aneurysm</td>
<td>Complete recovery; 18 mo</td>
</tr>
<tr>
<td>5/M/27</td>
<td>Left abducens palsy, decreased vision, pulsatile bruit</td>
<td>Low-flow left CCF</td>
<td>Left superior ophthalmic vein, left petrosal sinus, no cortical venous drainage</td>
<td>Not done</td>
<td>Spontaneous closure, coiling aneurysm ×2</td>
<td>Complete recovery; 24 mo</td>
</tr>
<tr>
<td>6/F/70</td>
<td>Bilateral exophthalmus and ophthalmoplegia, decreased vision, pulsatile bruit</td>
<td>High-flow left CCF</td>
<td>Bilateral superior ophthalmic vein, minor cortical venous drainage</td>
<td>+</td>
<td>Coiling aneurysm with balloon protection, carotid artery patent</td>
<td>Complete recovery; 34 mo</td>
</tr>
<tr>
<td>7/F/51</td>
<td>Right exophthalmus and ophthalmoplegia, pulsatile bruit</td>
<td>High-flow right CCF</td>
<td>Right superior ophthalmic vein, no cortical venous drainage</td>
<td>+</td>
<td>Coiling aneurysm with balloon protection, carotid artery patent</td>
<td>Complete recovery; 28 mo</td>
</tr>
<tr>
<td>8/M/58</td>
<td>Left exophthalmus, left abducens palsy, pulsatile bruit</td>
<td>Low-flow left CCF</td>
<td>Left superior ophthalmic vein, no cortical venous drainage</td>
<td>+</td>
<td>Spontaneous closure CCF, carotid occlusion</td>
<td>Remaining VI palsy; 4 mo</td>
</tr>
<tr>
<td>9/F/74</td>
<td>Bilateral exophthalmus and ophthalmoplegia, decreased vision, pulsatile bruit</td>
<td>Intermediate-flow right CCF</td>
<td>Bilateral superior ophthalmic vein, minor cortical venous drainage</td>
<td>+</td>
<td>Coiling aneurysm + carotid artery</td>
<td>Remaining right VI palsy; 7 mo</td>
</tr>
<tr>
<td>10/M/65</td>
<td>Left exophthalmus and ophthalmoplegia, vision 0 left, pulsatile bruit</td>
<td>High-flow left CCF</td>
<td>Left superior ophthalmic vein</td>
<td>+</td>
<td>Coiling aneurysm + carotid artery</td>
<td>Enucleation left eye; 38 mo</td>
</tr>
<tr>
<td>11/F/59</td>
<td>Pulsatile bruit, left temporal hemorrhage ×2</td>
<td>High-flow right CCF</td>
<td>Major cortical venous drainage</td>
<td>Not done</td>
<td>None</td>
<td>Death</td>
</tr>
</tbody>
</table>

Note: + indicates tolerance; –, nontolerance.
Medical Systems, Best, the Netherlands). In cases with rapid shunt to the cavernous sinus, additional filming was done, at 6 frames per second, to visualize the aneurysm in an early arterial phase before the cavernous sinus was opacified (Fig 1). After completion of the diagnostic work-up, in most cases a test occlusion of the ipsilateral carotid artery was performed to establish whether the patient could tolerate permanent occlusion of the carotid artery in case carotid sacrifice would be necessary to occlude the fistula. For that purpose, an over-the-wire balloon catheter (Sentry 15 mm; Boston Scientific, Fremont, Calif) was positioned close to or over the neck of the aneurysm and inflated to occlude the carotid artery and a second catheter was used for internal carotid and/or vertebral artery angiography to assess collateral circulation with emphasis on synchronous opacification of cortical veins of both hemispheres. After the test occlusion, the balloon catheter was deflated and left in place to assist in coiling of the ruptured aneurysm. The second catheter was subsequently also positioned in the ipsilateral carotid artery and served as a guide for the microcatheter that was placed inside the aneurysm. After positioning of the balloon and the microcatheter, coiling was started with protection of the internal carotid artery by the balloon and continued until the fistula was closed. In one patient (9), the neck of the aneurysm was wider than the protective balloon and, because test occlusion indicated tolerance, both the aneurysm and carotid artery were occluded with coils (Fig 2). In another patient (10), with a left CCF, tortuosity of vessels precluded a femoral approach to the left carotid artery and only one catheter could be positioned via direct carotid artery puncture. Test occlusion indicated tolerance, and both the aneurysm and carotid artery were occluded with coils. In one patient (4), a small ruptured cavernous sinus aneurysm was occluded by placement of an intrasaccular detachable balloon (Fig 3). Angiographic follow-up after 4–6 months was performed for all treated patients.

Results

The incidence of ruptured cavernous sinus aneurysms causing CCF was 1.5% (10 of 689 endovascular treated ruptured aneurysms). Ruptured cavernous sinus aneurysms causing CCF accounted for 24.4% (10 of 41) of treated symptomatic cavernous sinus aneurysms.

Clinical presentation, treatment, and outcome are displayed in the Table. All patients had an audible pulsatile bruit at presentation. Other clinical symptoms correlated well with venous drainage of the fistula: uni- or bilateral exophthalmus, ophthalmoplegia, and decreased vision were the key clinical symptoms when drainage was to the ipsilateral superior ophthalmic vein(s). Major retrograde drainage into the cerebral cortical veins in 2 patients was associated with intracranial hemorrhage in both.

Two low-flow CCFs closed spontaneously between diagnosis and planned treatment with resolution of symptoms. In one of these patients the cavernous sinus aneurysm was occluded with coils, and in the other patient the aneurysm was treated with balloon occlusion of the parent carotid artery.
Eight CCFs were successfully closed: in 7 patients with coils and in one patient with a detachable balloon. In 2 patients both the aneurysm and carotid artery were occluded with coils, and in 6 patients the CCF was closed with sparing of the carotid artery. During follow-up, 2 aneurysms that were occluded with coils showed reopening by compaction and were coiled for a second time. The aneurysm that was occluded with a detachable balloon reopened at follow-up and subsequently was occluded with coils (Fig 3). Clinical follow-up showed complete recovery in 7 patients, remaining abducens nerve palsy in 2 patients (in both cases treated with surgical shortening of lateral rectus muscle), death in one patient, and in one patient the left eye had to be enucleated.

Discussion

Although carotid cavernous fistulas as a result of a ruptured cavernous sinus aneurysm are rare and comprised 1.5% of endovascular-treated ruptured aneurysms in our institution, they accounted for 24.4% of all symptomatic cavernous sinus aneurysms. Clinical presentation is usually an acute onset of ocular symptoms and a pulsatile bruit is invariably present. Clinical diagnosis of a spontaneous CCF (caused by a ruptured aneurysm or a dural fistula) is simple by auscultation on the skull where the pulsatile bruit can be heard. Other symptoms correlate well with venous drainage: when drainage is to the superior ophthalmic veins this leads to ocular motor palsy, exophthalmus, conjunctival injection, and decreased visual acuity. Major cortical venous drainage was present in 2 of 11 patients and was associated with venous hemorrhages, a phenomenon analogous to dural arteriovenous fistulas or traumatic CCFs with cortical venous drainage.

When the clinical diagnosis of CCF is established with auscultation, adjunctive MR imaging or contrast-enhanced CT scanning is of little value but may reveal dilated superficial veins, an ominous sign of cortical venous drainage. Angiography establishes the diagnosis and can differentiate between a CCF caused by a ruptured aneurysm and a dural fistula. Although CCFs from ruptured cavernous sinus aneurysms may close spontaneously, as in 2 of our patients with low-flow CCFs, angiography and endovascular closure of the fistula should follow promptly to prevent possible hemorrhage from engorged cortical veins. After closure of the fistula, exophthalmus and conjunctival injection quickly resolve. Except for the one patient in whom an eye had to be enucleated because of extreme exophthalmus, visual acuity was restored in all and ocular motor palsy was cured in 6 of 8 patients. In 2 patients, complete ophthalmoplegia improved to abducens palsy.

Endovascular treatment is aimed at closing the fistula and preferably sparing the patency of the internal carotid artery. We found it useful to begin with a test occlusion: in patients with tolerance, one can proceed to carotid occlusion if technical problems occur, and in patients with nontolerance patency of the carotid artery must be maintained to prevent ischemic deficits. The over-the-wire balloon used for test occlusion was routinely left in place to assist in coiling of the aneurysm by protection of the carotid artery lumen. We used coils to occlude the ruptured cavernous aneurysm in all but one patient, because coils can be easily controlled with balloon protection of the carotid artery and occlude the aneurysm and the fistula gradually. In the 2 patients with simultaneous occlusion of the aneurysm and carotid artery, the use of coils was straightforward without the need for trapping the fistula.
tachable balloons can also be used to occlude the aneurysm and CCF. Balloons, however, have inherent disadvantages: premature detachment may occur, or, when the balloon does not occlude the fistula but blocks part of the venous drainage, rerouting of venous drainage to cortical veins may lead to hemorrhage. If the balloon is not congruent with the aneurysm, the fistula remains patent and placement of additional balloons may become problematic. Moreover, after balloon occlusion of the ruptured cavernous aneurysm, over time the aneurysm is likely to reopen after deflation of the balloon, necessitating additional coil occlusion.13

In conclusion, the incidence of CCF by a ruptured cavernous sinus aneurysm was 1.5%. CCF was the presenting symptom in 24.4% of treated symptomatic cavernous sinus aneurysms. Clinical symptoms correlate with venous drainage. Drainage to cortical veins may lead to intracranial hemorrhage. Endovascular treatment with coils is effective in occluding the fistula.

References