Use of coronary stent grafts for the treatment of high-flow carotid cavernous fistula

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SUMMARY
Carotid cavernous fistulas are abnormal communications between the carotid artery or its branches and the cavernous sinus. It can be traumatic or spontaneous. The widely accepted treatment is by detachable balloons. Advancements in the field of endovascular medicine made available other options for the treatment of this condition. Covered stents are widely available and offer preservation of the parent artery while occluding the fistula.

BACKGROUND
The carotid cavernous fistula (CCF) could cause life-threatening bleeding, and the endovascular management is considered to be the gold standard. Thanks to improvements in techniques and devices in recent years, covered stents provide a widely available alternative treatment option for the CCF. Further data are needed about the neuroendovascular usefulness of the covered stents.

CASE PRESENTATION
A 20-year-old man was transferred to our clinic for the treatment of life-threatening CCF. The patient had suffered a car accident a month earlier and has been in coma and on ventilator since. He had skull base fractures on both sides in the basal temporal region, fracture of petrous part of the right temporal bone, fractures of the right zygomatic bone, right part of the maxilla, and blood collections in the maxillary, ethmoidal and sphenoidal sinus. CT of the head had visualised a CCF on the right side. Since the crash, the patient has had several episodes of profuse bleeding from both ears and the nose. They were managed conservatively with placement of temporary tamponade. He required more than 10 hemotransfusions during the hospital stay.

On presentation, the patient was in severe general condition, intubated but with normal vital signs (HR 84 bpm, blood pressure 123/77 mm Hg). He had a quite tense right eye proptosis. The left eye was spontaneously opened. Glasgow-Liege Coma Scale (GLCS) Score was 11. The left corneal and glabellar (frontalorbicular) reflex were absent. The direct pupillary reaction to light was preserved as well as oculocephalic reflexes. A quadriparesis with spastic paralysis of the left arm, anisoreflexia and bilateral negative Babinski’s signs were evident. On pain stimulation, there was minimal flexor response in the right limbs. The head was deviated to the right side, with spontaneous mild movements. Verbal contact was missing.

INVESTIGATIONS
The CT scan on presentation confirmed the previously described bone fractures, CCF and multiple old haemorrhagic contusions in the right temporal, frontal and parietal zones (figure 1). The right superior ophthalmic vein was dilated up to 7.5 mm, the right inferior ophthalmic vein was dilated up to 2.6 mm. Due to the high-flow CCF and the backflow into the ophthalmic veins, the right facial vein was arterialised and visible on the three-dimensional angioreconstruction (figure 2). The CCF was classified as traumatic high-flow fistula type A.

MRI scan on admission showed multiple small haemorrhagic contusions in the right hemisphere of the brain and cerebellum, which have greatly reduced in size compared with the MRI 3 weeks earlier. In this case, we see multiple lesions in the cerebrum, cerebellum and brainstem with perivascular haemorrhages, contusions and lacerations of the brain parenchyma with necrotic, dystrophic and oedematous changes, which all lead to damage to the upper right motor neuron and the right
abducens nerve. The upper motor neuron syndrome may or may not be accompanied by the Babinski sign. The sign itself simply implies a pyramidal tract lesion and may not be associated with any of the upper motor neuron signs, which are extrapyramidal in origin (weakness, spasticity, clonus, hyperreflexia). The lesion in the right part of the cerebellum is responsible of the ataxia in the right limbs and the contusion in the right hemisphere and the brainstem are responsible for the weakness, hyperreflexia in the right limbs plus the central lesions of the facial nerve.

TREATMENT

Using right femoral approach, a diagnostic angiography was performed, revealing the CCF on the right side (figure 3 and video 1). The angio confirmed that the diagnosis of high-flow CCF type A. There was neither arterial steal, nor the fistula affected the rest of the arterial intracranial circulation in any way. The diagnostic catheter was exchanged for a 90 cm 6F Flexor Shuttle Guiding Sheath (Cook medical, USA). A 0.014 Runthrough NS PTCA guide wire (Terumo, Japan) was positioned in the distal part of anterior cerebral artery and the long sheath was placed in the right internal carotid artery (ICA), as high as possible. A guiding catheter was placed just below in CCF. Two covered stents, PK Papyrus 3.0/20 mm (Biotronik, Germany) and PK Papyrus 3.5/20 mm, were consecutively implanted in the ICA, covering the fistula orifice (figures 4 and 5; videos 2 and 3). Small residual filling of the fistula was visualised (figure 6).
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and video 4) and post dilation with NC Sprinter 4.0/20 mm (Medtronic, Ireland) was performed with excellent final result (figures 7–9, videos 5 and 6). Neither artery straightening nor vessel spasm was evident at the end of the procedure. The patient received dual antiplatelet therapy after the procedure as is the in-hospital protocol in such procedures. The patient was started on aspirin 100 mg once daily and clopidogrel 75 mg once daily for 1 year, followed by the discontinuation of clopidogrel.

The tamponades in the ears and nose were removed 6 hours after the procedure with no residual bleeding. Previous attempts to remove the tamponade have led to recurrent severe episodes of epistaxis. We must emphasise that this time the patient was even receiving dual antiplatelet therapy, which furthermore gave us confidence that the fistula was isolated. The patient was kept in induced coma after which he was extubated and gradually started rehabilitation. The neurological status on discharge 2 months after the admission was significantly improved with GLCS of 15. The right eye exophthalmos/proptosis was significantly reduced. The patient was able to execute elementary commands and to move even to a limited extent his right limbs with ataxia. The left spastic paresis remained as well as moderate right abducens nerve lesion and left central facial palsy.

OUTCOME AND FOLLOW-UP

In the 12-month follow-up period, no bleeding of embolic incidents. There has been a great progress in the rehabilitation of the patient, with improvement of both his motor and speaking skills.

DISCUSSION

In 1985, Barrow et al published a classification of CCFs by their anatomical location on angiography. CCF was divided into four categories: type A fistula—a direct communication between the ICA and the cavernous sinus (CS), type B—between the meningeal branches of ICA and the CS, type C—between meningeal branches of the external carotid artery and the CS and type D—between the meningeal arteries of both carotid arteries and the CS. The CCF can also be classified as direct (type A) and
indirect (types B, C and D), low-flow and high-flow, traumatic and spontaneous.1

The most common presentations of CCF are proptosis of the eye on the ipsilateral side of the fistula, cephalic bruit, headache, diminished visual acuity and conjunctival chemosis.2 The typical presentation of a type A CCF is the development of Dandy’s triad—exophthalmos, bruit and conjunctival chemosis. Otorrhagia and epistaxis may be seen in up to 3% of the cases of CCF.3

The superior and inferior veins of the eye both drain into the CS, the high-flow fistulas increase the pressure in the CS and cause backflow into the ophthalmic veins, leading to stasis of blood in the eye. That may cause proptosis, episcleral and conjunctival arterialisations, and enlargement of the extraocular muscles. Increased episcleral pressure and venous pressure may result in elevated intraocular pressure and secondary glaucoma.4

A spontaneous resolution of high-flow CCF has been described in the literature but it is extremely rare. Clinical signs such as increased intracranial pressure, rapidly progressive proptosis, haemorrhage, diminished visual activity and transient ischaemic attacks are associated with poor prognosis and warrant emergent and aggressive interventional treatment to improve outcome.5

The first endovascular treatment of CCF with a detachable balloon done by Serbinenko was described in 1974.6 Since the 1980s, this has been the golden standard for the treatment of CCFs. Due to advancement in endovascular devices and techniques, and the unavailability of detachable balloons in recent years, coils, alcohol and stent graft embolisation of CCF have been an emerging treatment options.5

The endovascular treatment options of type A CCF device mainly in two groups—transarterial and transvenous. Different

Figure 7 Final fluoroscopy showed patient’s internal carotid artery and completely isolated fistula. Left oblique projection.

Figure 8 Final fluoroscopy showed patient’s internal carotid artery and completely isolated fistula. Right oblique projection.

Figure 9 Final fluoroscopy showed patient’s internal carotid artery and completely isolated fistula. Left cranial projection.

Video 5 Final fluoroscopy left oblique projection—final fluoroscopy showed patient’s internal carotid artery and completely isolated fistula. Neither artery straightening nor spasm is evident.
sources give different data, which one is the preferred method but anyway endovascular repair has excellent results. Besides stent grafts, the another transarterial options are placement of detachable balloons, placement of coils or use of ethylene vinyl alcohol copolymer (EVOH) to occlude the direct communication. The coils or balloons may migrate through the defect into the parent vessel if it the tear is big enough (like in our case), potentially causing brain ischaemia. The transvenous approach can be used as a main one or when the ICA cannot be accessed due to traumatic injury, severe tortuosity or inability to catheterise the tear. The route to the CCF in this case goes usually through the internal jugular vein and the inferior petrosal sinus. The goal is to discontinuette the venous outflow from the CS using coils or liquid embolic agents. In both transarterial and transvenous approach, there are a number of times when coils are not enough for the full embolisation of the fistula and EVOH or Onyx is used as an additional agent in order to penetrate and fully occlude the communication. Because of the wide orifice of the fistula and the hugely dilated superior ophthalmic vein, we estimated the risk of coil migration as high. Transvenous approach was planned as a backup option in case of failure of the transarterial approach due to the high risk of coil migration and the lack of experience of our team with this approach for intracranial procedures. Embolisation with Onyx (Medtronic, Ireland) was considered but due to insufficient experience of our team with it and the hazard of postembolisation CS thrombosis and swelling, we deemed it inappropriate. We choose to repair the vessel defect with stent graft PK Papyrus deployment as our main strategy.

The four largest studies to date of CCF, treated with stent graft, involve a total of 25 patients and report excellent results with mean follow-up of 12 months and 84% patency of the ICA. Covered stents are widely available, offered in a great range of sizes and offer the possibility of ICA patency preservation, while sealing the fistula. The main problem of these covered stents is the fact that they are still not approved for intracranial use in some neurological conditions. Another disadvantage is the limited flexibility of the graft, complicating their navigation through the tortuous anatomy of the intracranial vessels and also the risk of ‘straightening’ of the artery. The stent we used (PK Papyrus) offers improved flexibility thanks to its single stent design and made possible the perfect final result we managed, without any straightening of the artery at the end.

A problem of the Onyx and coil treatment of CCF is the mass effect. Fewer mass effects have been observed with stent graft compared with coils and Onyx in the treatment of cerebral aneurysms. Aneurysms treated with covered stents have greatly reduced recurrence rates compared with those treated with coils or liquid embolic materials. Additional data are needed to see if the same applies for the CCFs.

**Patient’s perspective**

Patient perspective—I am greatly thankful to everyone involved in my treatment. I am fully motivated continue rehabilitation until I get back to my old self.

Family’s perspective—We cannot imagine our life right now if our son has not received the treatment, he did. Without the fully committed staff of the hospital, we would not be where we are. After the procedure, the condition of our son improved minute by minute and eventually, he was out of come, which gave is hope that everything will be OK. We believe that with active rehabilitation he can go back to his life.

**Learning points**

- Carotid cavernous fistulas can be a life-threatening condition.
- Covered stents are a feasible treatment option for CCF.
- Approved neuroendovascular covered stent and additional data are warranted.
- Double antiplatelet therapy after stenting was not causing problems in this patient with post-traumatic and haemorrhagic brain injuries, and previous severe cranial bleeding.

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Case reports provide a valuable learning resource for the scientific community and can indicate areas of interest for future research. They should not be used in isolation to guide treatment choices or public health policy.

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