Extracranial Internal Carotid Artery Aneurysms: case report of a saccular wide-necked aneurysm and review of the literature

Lukla Biasi, Matteo Azzarone, Alessandro De Troia, Pierfranco Salcuni, Tiziano Tecchio
Unit of Vascular Surgery, Department of Surgical Sciences, University of Parma, Parma, Italy

Abstract. Purpose: to describe a case of atherosclerotic Extracranial internal Carotid Artery Aneurysm (ECAA) with an unique “diverticulum-like” morphological structure, surgically treated and to review the experiences of major referral centers. Case Report: a 76-year-old woman presented with persistent dysphagia and transient hypophonia caused by a growing pulsatiling mass of the neck; duplex ultrasonography and angio-CT scan demonstrated a wide-necked, saccular aneurysm at the origin of the right Internal Carotid Artery (ICA). A total aneurysmectomy with patch angioplasty of the aneurysm of the neck on the carotid wall was successfully performed. No perioperative complications were encountered; at one year follow-up the patient was asymptomatic without any neurological complication and with carotid patency. Conclusions: open surgery remains the gold standard for the treatment of extracranial internal carotid artery aneurysms in terms of patency and reduced risk of adverse complications; endovascular procedures may, in selected cases, provide a valuable additional tool in the armoury of the physician. (www.actabiomedica.it)

Key words: Extracranial carotid artery, carotid aneurysm, surgery

Introduction

Extracranial internal Carotid Artery (ECA) Aneurysms are a rare entity, estimated at 0.1% to 2% of all carotid procedures performed at major referral centers (1). They can be classified according to their different physiopathology into true or false aneurysms and according to their etiology into atherosclerotic, dysplastic, infective, posttraumatic and iatrogenic (2). ECA Aneurysms may be asymptomatic but neurologic manifestations are the most frequent symptoms (3, 4).

The natural history of this pathology is associated with spontaneous progression of the aneurysm, most commonly associated with a high risk of neurological thromboembolic events, cranial nerve compression and, more rarely, rupture (5).

The surgical treatment of ECA aneurysms is determined by the aetiology, size and location of the aneurysm and any associated contraindications.

Whereas antiplatelet and anti-coagulant drugs are considered by many authors as the first-line approach for asymptomatic aneurysmal lesion, open surgery still remains the most valid option to prevent the most probable, severe and life-threatening complications, in particular embolisms (1).

Endovascular treatment, especially in the light of the recent progress in stent grafting, may be considered as a therapeutic choice but only in selected cases; more comparative studies and long-term follow up will be necessary to assess the role of endoluminal exclusion as a first choice treatment.

Case Report

A 76-year-old woman was admitted to our Surgical Unit with a diagnosis of Extracranial internal Carotid Artery aneurysm.
The patient presented with a large, growing, pulsatile mass in the right side of the neck (easily palpated due to the slimness of the patient’s neck). Auscultation evidenced no pathological carotid bruit. She reported persistent symptoms of dysphagia, and transient hypophonia; there was no history of cerebrovascular symptoms or cervical trauma.

The patient reported a medical history significant for hyperlipidemia with drug-controlled familial hypercholesterolemia; there was no familiarity for chronic arteriopathy.

The diagnosis was established through duplex ultrasonography followed by a cerebral and neck computed tomography: angio-CT scan revealed a wide-necked, saccular, “diverticulum-like” aneurysm at the origin of the right ICA, developing from the side of the carotid wall with a diverticular shape, with a longitudinal diameter of 15 mm and transversal diameters of 14 mm x13 mm (Fig. 1-2); in addition, a small atherosclerotic, calcific plaque of the bulbar ICA without hemodynamic flow impairment was shown; the left carotid axis appeared free from aneurysmatic lesions with an atherosclerotic, non-stenotic plaque at the carotid bifurcation. No other aneurysms were identified in the cerebral circulation. Further pre-operative angiographic investigation was deemed unnecessary.

The ECA aneurysm was surgically treated under loco-regional anesthesia; the carotid branches and the aneurysm were exposed using a short standard lateral cervical incision obtaining limited isolation and control of the cervical carotid aneurysm (CCA), ECA, and proximal and distal ICA (Fig. 3). Intraluminal shunting was unnecessary as no neurological symptoms presented after carotid clamping.

Due to the unusual saccular, wide necked, “diverticulum-like” shape of the aneurysm, we performed an

Figure 1. Angio-CT scan: frontal view of carotid axis

Figure 2. Angio-CT scan: lateral view of the right carotid axis

Figure 3. Intraoperative view: wide-necked, saccular, “diverticulum like” aneurysm at the origin of the right ICA
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Extracranial Internal Carotid Artery aneurysms are defined as localised increases of the calibre of more than 50% as compared with the reference values of the ICA (0.55±0.06 cm in men; 0.49±0.07 in women).

They may be classified like true or pseudo-aneurysms according to their different aetiologies. Pseudoaneurysms (PAs) comprise only 14% of ECA aneurysms in the reviewed literature (7) even though they exceptionally represented the 57% of these causes in the THI series (El Sabrout et al.). PAs are mainly secondary to previous endarterectomy (CEA) with the onset of symptoms ranging from 1 month to 15 years after the original carotid procedure; other contributing factors of PAs may be trauma (car accidents, stab-wounds, iatrogenic central venous cannulation) and infections: patch complications or recurrent tonsillitis or pharingitis reaching the carotid artery wall following peritonsillar abscess with subsequent ischemia of the wall leading to acute rupture or causing septicemia and invasion of vasa vasorum (8).

The usual infectious agents are staphylococcus aureus and streptococcus pyogenes. This etiology was firstly described in 1843 by Liston (9) In 1933 Salinger and Pearlman reviewed 228 cases(10). Before the introduction of antibiotics, infection was the main cause of ECA aneurysms but since then, only 31 additional cases have been reported (11, 12).

True aneurysms are most commonly due to atherosclerosis or fibromuscular dysplasia.

Although Moreou reported dysplastic lesions (single or associated with spontaneous dissection) as a major cause of ICA aneurysms (13), atherosclerotic degeneration is assumed to be the most frequent etiology (14, 15): up to 53% in a review of the literature by McCann (72-85) (2).

Atherosclerotic Aneurysms (AAs) are usually found in patients aged 50 to 70 years old with a male-female ratio of 1.9:1 (16).
AAs may involve various portions of the carotid artery: they tend to be located in the common artery bifurcation and in the proximal ICA, size ranging from 1.5 to 5.0 cm, whereas dysplastic aneurysms, often associated with chronic dissection are more distally located (17).

Carotid aneurysms of the external carotid artery are less frequent but should also be considered in the differential diagnosis of patients presenting with a pulsating mass in this area.

Traumatic aneurysms are either due to compression and stretching of the artery on the lateral mass of the 1st cervical vertebra or the shearing of the artery at the level of its intracranial entry (18,19).

Other rare contributing factors to aneurysm formation include neck irradiation, neurofibromatosis, Marfan’s Syndrome, Bechet’s Syndrome and Takayasu’s arteritis (20); 30% to 60% of ECA aneurysms are symptomatic for thrombo-embolic focal or non-focal symptoms; more rarely cranial nerve compression may result in neurological deficits (as shown in our patient’s clinical presentation).

Spontaneous progression of ECA aneurysms is associated with a higher risk of mainly thrombo-embolic neurological complications; other adverse events may be rupture, with massive haemorrhage, and nerve compression (3).

Conservative treatment is based on anticoagulant or anti-aggregant therapy. A non-surgical approach may be indicated in young patients with asymptomatic, traumatic or spontaneous dissecting aneurysms (1).

In many cases, however, medical treatment is considered to be ineffective or even dangerous, with a persistent risk of damage due to haemorrhage (21).

Before 1950, 71% of the patients non-surgically treated later died because of complications due to the aneurysm (22). More recently one-fifth of ECA aneurysms were non-surgically managed with an overall stroke rate of 50% (23).

Surgical treatment of ECA aneurysms may therefore be seen as a first-line approach, even in asymptomatic patients.

The main indication for a surgical approach is the prevention of permanent neurological deficits arising as sequelae of thromboembolism.

Historically, the first successful operation was performed by Sir Astley Cooper in London, in 1808: the patient was a 50-year-old labourer with a bulbar aneurysm of the ICA that was “about the size of a pullet’s egg” with a “remarkably strong pulsation of the tumour (24).

Until the 1950s surgical treatment of ECA aneurysms only consisted of carotid ligation, with an alarming mortality/major stroke rate of 20% to 60% (nowadays ligation is only performed in about 10% of the cases, mainly if a distal ICA plaque occlusion is present) (25).

By the 1970s, reconstruction of the carotid axis had definitely replaced ligation as a treatment option.

The success of open surgery is dependant on the aetiology, size, location of the aneurysm and associated co-morbidities.

The most frequent surgical procedures for Atherosclerotic Aneurysms are represented by: total resection of the aneurysm with direct end-to-end anastomosis or interposition of a prosthetic graft or the saphenous vein; rarely performed, it may be indicated for fusiform aneurysms; partial aneurysmorraphy and patch (synthetic or venous) closure; this may be indicated in saccular aneurysms. The aim is the preservation of the posterior carotid wall, thus reducing the risk of cranial nerve injuries; extracranial – intracranial by-pass graft: this is a very difficult and rarely performed technique, moreover it does not exclude the aneurysm from the blood stream and therefore does not eliminate the risk of embolization or rupture unless followed by carotid ligation; in the presented case, due to the unique morphological structure of the aneurysm (‘diverticulum-like’), we performed a total aneurysmectomy with patch angioplasty of the aneurysm of the neck on the carotid wall to restore the normal lumen of native artery; in our review of the literature we did not find any similar presentation justifying this surgical approach.

A review of the literature suggests that surgical repair has proven to be effective and safe in the following situations: THI (1): 67 aneurysms in 35 years with a 9% overall mortality/major stroke incidence and a 6% rate of cranial nerve injury; Zhang (26): 66 aneurysms in 17 years with a 1.6% immediate death rate and a 6% rate of adverse events; D’addato (14):
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24 aneurysms in 21 years with no perioperative deaths and 4.5% immediate major or minor adverse events.

This is compared with an estimated complication rate of between 20 to 71% in non surgically treated ECA. However it should be noted that open surgery may be invasive, disabling and extremely difficult when lesions are located in the distal portion of the ICA, at the base of the skull with a high risk of cranial nerve damages (permanent nerve palsy up to 42% in Mokri et al. (16).

This may be the reason why endovascular procedures have been recently used in the treatment of ECA aneurysms by means of embolization with detachable coils, endografts or covered stents. The largest series up to date was reported by Bergeron et al. (6 cases endovascularly treated) (27).

The use of covered stenting allows simultaneous exclusion of the aneurysm and dilatation of an eventually stenosed distal ICA, with excellent results in terms of patency and absence of migration (28).

Very few neurological complications following endovascular exclusion and little nerve damage have been reported.

By contrast, it should be noted that stented segments of the carotid artery have been shown to have less flexibility when the patient moves his head (29). In addition, endoluminal treatment of distal ICA aneurysms does not offer the possibility of surgical conversion in case of complications durino or after the procedure.

In conclusion, studies of larger series of patients with extensive post-operative follow-up will be required to prove the safety and efficacy of endovascular procedures.

In this case, the presence of a wide neck (saccular, “diverticulum-like” aneurysm), suggested that the open surgery approach was the best treatment for the patient, in terms of patency and the reduced risk of adverse complications (the patient’s thin neck made the task easier).

In conclusion, we believe that, while open surgery remains the gold standard to treat ECA aneurysms, endovascular procedures may, in the future and for selected cases, provide a valuable additional tool in the armoury of the physician.

References

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Accepted: September 1st 2008
Correspondence: Lukla Biaisi, M.D.
Unit of Vascular Surgery,
Department of Surgical Sciences,
University of Parma
Via Gramsci, 14
43100 Parma, Italy
Tel. 00390521702272
E-mail: luklabiasi@hotmail.com