Surgical Position, Cause of Extracranial Internal Carotid Artery Dissection, Presenting as Pourfour Du Petit Syndrome: Case Report and Literature Review

Cerrahi Pozisyon Nedeniyle Ekstrakraniyal İnternal Karotid Arter Diseksiyonunun Pourfour Du Petit Sendromu Olarak Ortaya Çıkması: Olgu Sunumu ve Literatür Derlemesi

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ABSTRACT
Dissection of the internal carotid artery is a rare cause of stroke overall, but causes 22% of strokes in younger patients. A common clinical presentation is as Claude Bernard Horner syndrome. We report a craniotomy with 30 degrees rotation of the neck (standard position) in a patient with no major risk factors for carotid dissection, who showed a Pourfour du Petit syndrome due to a dissection of the internal carotid artery. To the best of our knowledge, this is the first reported case in which a common surgical position causes an internal carotid artery dissection in a patient without relevant risk factors. The presentation with Pourfour du Petit syndrome is extremely unusual.

KEYWORDS: Carotid dissection, Pourfour du Petit syndrome, Neck rotation

ÖZ

ANAHTAR SÖZÇÜKLER: Karotid diseksiyonu, Pourfour du Petit sendromu, Boyun rotasyonu

INTRODUCTION
Carotid artery dissection is a rare cause of stroke. Although responsible for only about 2.5% of all strokes, it accounts for up to 22% of strokes in younger patients. The most common age group is between 35 and 50 years old, with a distinct peak incidence in the fifth decade; the prevalence is similar in both sexes. Extracranial internal carotid artery (ICA), from the carotid bulb to the cranial base, is more vulnerable to dissection compared to the intracranial portion; the most frequent dissection is 2-3 cm distal to the carotid bulb. The dissection can be classified as spontaneous or traumatic. Intrinsic factors associated with or predisposing to spontaneous dissection include the following: aortic root dilation, bicuspid aortic valve, increased arterial compliance and arterial wall stiffness, elevated CRP levels, fibromuscular dysplasia, genetic factors, hypertension, hypercholesterolemia; and connective tissue diseases (e.g., Ehler-Darlos syndrome type IV, Marfan syndrome, autosomal dominant polycystic kidney disease, osteogenesis imperfecta, alpha 1 antitrypsin deficit) (1, 5, 14, 25).

Traumatic dissection can occur by direct blow to the neck or by hyperextension, rotation or lateroverision of the neck. Sometimes the trauma is so mild that it is misclassified as spontaneous dissection. Minor traumas reported in literature...
include vomiting, painting a ceiling, a long telephone conversation, riding a roller coaster, and practicing yoga. Among the iatrogenic causes reported in literature are neck manipulations in anesthetized patients, aspiration with fine needles, and postcardiac resuscitation. Infection, particularly in the respiratory system, has been implicated as a trigger for carotid artery dissection (2, 6, 7, 10-13, 15, 17, 18, 23).

A large percentage of patients with carotid artery dissection are asymptomatic. In symptomatic patients, the clinical manifestations may be due to local complications. Facial or neck pain is the most common symptom and usually the first to appear, partial Claude Bernard Horner syndrome (CBHS) occurs in 50% of patients. Other symptoms include headache, unilateral or pulsatile tinnitus, amaurosis fugax, retinal infarct, and palsies of the lower cranial nerves (the hypoglossal being the most frequently affected). On the other hand, signs of ischemia may be present in 50% to 90% of cases. Most strokes occur in the first week of local symptoms onset and are located in the territory of the middle cerebral artery. Diagnosis can be made by angiography, computed tomography angiography (CTA), magnetic resonance angiography (MRA) and ultrasound (20). Differential diagnosis is required with respect to cluster headache, migraine, retinal artery occlusion, herpes zoster and musculoskeletal neck pain (16, 19, 24).

There is currently no consensus on the treatment of carotid artery dissection. In patients with an acute neurological deficit, intravenous thrombolysis or even endovascular treatment may be a possibility. In general, anticoagulation with heparin has been the treatment in the acute phase and subsequently acenocoumarol. More recently, and especially in asymptomatic patients, antiplatelet therapy has become the most commonly used treatment (4, 8, 13, 19, 22, 26).

**CASE REPORT**

In March 2012, a 43-year old woman with insulin-dependent diabetes mellitus and hypercholesterolemia had a subarachnoid hemorrhage due to the rupture of a left terminal carotid aneurysm. The aneurysm was embolized. As complications of subarachnoid haemorrhage, severe vasospasm of right middle cerebral artery (MCA) and right anterior cerebral artery (ACA) occurred, leading to a right frontal cerebral infarct and hydrocephalus that required ventriculoperitoneal shunt.

The first arteriography showed the existence of an unruptured polylobulated aneurysm of the right MCA. Three months later, a new arteriography showed a left terminal carotid embolized aneurysm that was totally excluded, and a right middle cerebral artery aneurysm, proposed for open surgery. No carotid artery dissection or other problems were observed. In November 2012, a preoperative MRA was performed; no change was observed. In January 2013, the patient had a seizure, and a CTA was performed. It showed no changes from previous images and normal ICA diameters (Figure 1).

Surgery was performed in March 2013 under general anesthesia and neurophysiological monitoring, with the patient in supine position, head slightly hyperextended and lateralized to the left 30 degrees. We performed a right frontotemporal craniotomy, dissected the Sylvian valley and exposed the aneurysm. The right middle cerebral artery was temporarily clipped for 18 minutes and then definitively, with two clips. The surgery lasted 5 hours. No anesthesiological or neurophysiological problems were detected (Figure 2).

The patient had a delay in awakening (30 minutes), and showed left anisocoria, (which reversed in about 24 hours). Postoperative CT showed no complications.

Postoperative arteriography was performed 48 hours after surgery, showing complete occlusion of the left ICA in its extracranial portion and leading to the diagnosis of carotid dissection. The clipped aneurysm was excluded from circulation (Figure 3).

Antiplatelet therapy with acetylsalicylic acid 100 mg was started. Cranial MRI did not show any acute ischemic lesions.

![Figure 1: Cerebral tomography angiography of the supraaortic trunks. The internal carotid arteries are permeable, with normal diameters.](image-url)
DISCUSSION

This is the first case reported in the literature of carotid dissection due to a common surgical position. The patient had no relevant risk factors, such as fibromuscular dysplasia, that predispose to carotid dissection. The 30 degrees rotation of the head used in this case is the standard access to MCA aneurysms. There is a clear temporal association between the anisocoria and the surgery. Therefore, we can say that carotid dissection was not spontaneous. It likely resulted from the 5 hours of neck rotation.

The literature describes cases of dissection due to forced positions of the neck, especially in patients with risk factors (9, 11-13, 21), but there are no case reports for a common surgical position.

In this case, carotid dissection was asymptomatic from the ischemic cerebral standpoint, probably due to compensation from the right hemisphere. Interestingly, the 18-minute temporary clipping of the MCA during the surgical procedure did not interfere with that compensation. Anesthetic monitoring revealed no changes during surgery. Intraoperative neurophysiological monitoring was performed including an electroencephalogram, somatosensory evoked and motor evoked potentials after stimulating the median and tibial nerve. The recordings were stable throughout surgery. We wonder whether such compensation would have been possible if the temporary clipping had been maintained longer.

The clinical presentation of carotid dissection with mydriasis is unusual. If there is pupillary involvement, it is more common in the context of CBHS due to injury of the sympathetic nerve fibers, with its classic triad of miosis, enophthalmos and ptosis (16, 19, 22, 24). Pourfour du Petit syndrome (PDPS), a little known entity and the inverse of CBHS, is characterized in its complete form by mydriasis, exophthalmos, upper eyelid retraction with increased lid fissure and periorbital hyperhidrosis. All of these are evidence of irritation (hyperexcitation) of the ipsilateral cervical sympathetic nerve chain. Other causes of PDPS described in the literature include cervicofacial trauma, thyroid tumors, primitive carotid aneurysms, and iatrogenic causes. Recognition of this syndrome is valuable for its high locator value, making possible the early diagnosis of various potentially serious pathologies (3).

Therapeutic management of these patients varies from antithrombotic drugs to complex endovascular and surgical techniques in the series and observational studies reported in the literature (6, 19, 26). Choosing the best drug treatment remains controversial. In the absence of randomized studies, the superiority of anticoagulant therapy compared to antiplatelet therapy has not yet been demonstrated. Antiplatelet therapy was chosen in our patient, after assessing both treatments for effectiveness and safety (given her history of recent surgery).

CONCLUSION

Standard surgical position in neurosurgery that involves prolonged neck rotation may cause carotid dissection. Therefore, postoperative neurological examination, especially if the signs and symptoms have no correlation with the surgical site, must always take into account this possible diagnosis, especially in patients with risk factors (e.g., fibromuscular dysplasia).

Pupillary dilation must be considered as a clinical manifestation of ICA dissection in the context of PDPS.
REFERENCES