Axillary Aneurysm Due to Behcet’s Disease Revealed by Nerve Compression

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Introduction

Aneurysms of the axillary artery are rare, most often they are pseudo-anneurysms of post-traumatic origin, true aneurysms are rare. We report the case of a patient who consulted for signs of nerve compression of the left upper limb due to an axillary aneurysm, the patient suffers also of Behcet's disease.

Materials and methods

We report the case of a 44-year-old patient with a history of Behcet's disease since adolescence, who consulted in orthopedic surgery unit for paresthesia of the left upper limb with amyotrophy of the thenar and hypothenar eminence that had been evolving for 6 months without concept of trauma.

An electromyogram, revealed a block of conduction. Diagnosis of axillary artery aneurysm was confirmed by echodoppler findings and CT angiography, which showed the existence of an aneurysm of the left axillary artery (Figure A and B).

Discussion

Aneurysms of the axillary artery are rare. Etiologies include atherosclerosis [1], post-stenotic dilatation related to thoracic outlet syndrome [2], elastopathy (Marfan, Ehler Danlos), infections, and penetrating or closed trauma [3].

The pathophysiology of arterial vascular disease in Behcet disease remains poorly understood. Some authors believe that the anatomic substratum of these arterial lesions is the abnormalities of the arterial wall involving in particular parietal factors [4,5]. The role promoting local trauma has been highlighted, such as the occurrence of aneurysm at the point of arterial puncture when performing arteriography, measurement of blood gases or trauma [6]. The involvement of blood hypercoagulability is not clearly established [4]. The treatment of these lesions is the surgical excision of the aneurysm.
With the significant development of endovascular treatment, endovascular excision is now the treatment of choice for this type of lesion [7].

**Conclusion**

Aneurysmal lesions of the axillary artery and its branches are rare and may be related to several etiologies, but they may also be the consequence of systemic vasculitis such as Behçet’s disease. The physician must always bear in mind this diagnosis in order not to venture to perform biopsies or punctures of any axillary mass before performing a minimum morphological assessment including at least one ultrasound coupled to Doppler.

**Declaration**

The authors declare that they have no conflict of interest.

**References:**