Vascular ring related to Kommerell diverticula: case report

Anel vascular associado a divertículo de Kommerell: relato de caso

Décio Cavalet Soares ABUCHAIM¹, Martin BURGER², Silvana Agnoletto BERWANGER², Djalma Luis FARACO³

Abstract

Report of a surgical treatment for vascular ring (right aortic arch and the anomalous origin of the left subclavian artery) related to Kommerell diverticulum with resection of the ligamentum arteriosum (ductus arteriosus), suture of the Kommerell diverticulum, and reimplantation of left subclavian artery in the ipsilateral carotid artery through left thoracotomy in a 13-year-old female.

Descriptors: Aorta, thoracic, abnormalities. Subclavian artery, abnormalities. Aortic diseases, surgery.

Resumo

Relato do tratamento cirúrgico de anel vascular (arco aórtico à direita e origem anômala de artéria subclávia esquerda) relacionado a divertículo de Kommerel, com realização de seção de ligamento arterial, rafia de divertículo e reimplante de artéria subclávia esquerda em carótida ipsilateral, por toracotomia esquerda, em uma paciente de 13 anos.


1. MCMSc in Clinical Surgical by UFPR; cardiovascular surgeon; manager of Surgical Center of Hospital Santa Catarina – Blumenau, SC.
2. Cardiovascular Surgeon at Hospital Santa Catarina – Blumenau, SC.
3. MCMSc in Clinical Surgical by UFPR; cirurgião cardiovascular; manager of Cardiovascular Surgery Service of Hospital Santa Catarina – Blumenau, SC.

Work done at Hospital Santa Catarina, Blumenau, SC.

Correspondence address:
Décio Cavalet-Soares Abuchaim
Rua Amazonas, 301 – Blumenau, SC - CEP 89020-900.
E-mail: decioabu@terra.com.br

Article received in April 20º, 2007
Article accepted in July 20º, 2007
INTRODUCTION

Kommerell’s diverticulum, which is a rare condition, involves the left fourth aortic arch (LAA) and the anomalous origin of the right subclavian artery (RSA) (0.5%-2%), or the right aortic arch (RAA) with the anomalous origin of the left subclavian artery (LSA) (0.05%-0.1%) [1]. A cause of aberrant subclavian artery origin can be an abnormal regression of the primitive fourth aortic arch during embryogenesis. The left fourth aortic arch persists as the aortic arch, while the right fourth aortic arch remains as the right subclavian artery (RSA) and the innominate artery (obsolete term for brachiocephalic (arterial) trunk) [2].

The right aortic arch (RAA) with ligamentum arteriosum to the descending aorta is one of the two vascular rings that cause tracheoesophageal compression; usually, children with a double aortic arch had earlier onset of symptoms (stridor and dysphagia) than children with a right aortic arch and ligamentum arteriosum.

A diverticulum may become aneurysmal even after the ligamentum arteriosum resection [1], the procedure of choice usually performed.

CASE REPORT

A 13-year-old female patient with wheezing or stridor and dysphagia in the last 5 years was, at present, progressive for pasty food. She reported weight loss in the last few months, unmeasured.

It was observed, on a chest radiograph, a right aortic arch (Figure 1); on the esophagram, an extrinsic compression suggestive of vascular ring. An angiotomography showed a right aortic arch with Kommerell diverticulum at the anomalous origin of the left subclavian artery (Figure 2).

The patient underwent left thoracotomy, opening of the pleura, and exposure of the aorta through lung retraction. After the identification of the anatomical elements, the following procedures were performed: resection of the ligamentum arteriosum; suture of the Kommerell diverticulum (Figure 3), and resection and reimplant of the left subclavian artery into the left carotid artery (Figure 4). Afterwards, a procedure to free the fibrous bands adjacent (adhesive) to the esophagus and trachea was performed. The pleura was left open with the drainage tube placed in the normal position.
REFERENCES


DISCUSSION

The Kommerell diverticulum is an important vascular structure that through dilatation can cause symptoms, such as dysphagia, dyspnea, stridor, chest pain, emphysema, and pneumonia [2]. Left subclavian artery originating from Kommerell’s diverticulum which results in absence of pulse with the arm in a supine position is very unusual [4]. By the recurrence of symptoms in patients with a Kommerell diverticulum who had simple ligamentum resection, it is recommended surgical repair and primary reimplantation of the subclavian artery [2, 3], whenever the patients present with a diameter > 50 mm [2].

There were no further complications and the patient was transferred to the ward the next day without the drain. The patient was discharged on postoperative day 5, eating solid food, with normal voice, and absence of differential blood pressure between upper limbs.

CONCLUSION

The resection of the ligamentum arteriosum, the resection and reimplantation of the left subclavian artery into the left carotid artery, and the repair of Kommerell’s diverticulum, in spite of being a more complex procedure than the simple resection of the ligamentum arteriosum, the procedure of choice, has satisfactory result and can prevent the recurrence of symptoms by extrinsic compression.

REFERENCES


