

Revista Portuguesa de **Cardiologia**Portuguese Journal of Cardiology www.revportcardiol.org



CASE REPORT

Anomalous origin of left coronary artery: A malignant interarterial variant with a benign clinical course*

Maria Salomé Carvalho^{a,*}, Pedro De Araújo Gonçalves^{a,c}, Hugo Marques^b, Pedro Jerónimo De Sousa^a, Rita Calé^a, Helder Dores^a, Daniel Ferreira^a, Francisco Pereira Machado^a, Ana Aleixo^c, Miguel Mota Carmo^c, José Roquette^a

Received 3 October 2011; accepted 30 November 2011 Available online 29 May 2012

KEYWORDS

Anomalous origin of coronary arteries; Cardiac CT; Sudden death **Abstract** Anomalous origin of coronary arteries represents a clinical challenge because of the anatomical variability and possible functional consequences, the pathophysiological mechanisms involved, and the lack of large published series that would provide evidence to guide the clinical and therapeutic approach. The authors describe the case of a 55-year-old male patient with a long history of atypical chest pain who was considered to have a low to intermediate likelihood of coronary artery disease. Therefore, and also bearing in mind his physical limitations (congenital left leg atrophy), he was referred for cardiac CT to rule out coronary artery disease. The exam showed a left coronary artery arising from the right coronary cusp and with an interarterial course, between the aorta and pulmonary trunk. Although this is a potentially malignant anatomical variant with surgical indication, a conservative approach was chosen, considering the late diagnosis and particular risk-benefit profile.

 $\ \odot$ 2011 Sociedade Portuguesa de Cardiologia. Published by Elsevier España, S.L. All rights reserved.

PALAVRAS-CHAVE

Origem anómala das coronárias; AngioTC cardíaca; Morte súbita

Origem anómala da coronária esquerda: variante interarterial maligna com evolução clínica benigna

Resumo A origem anómala das artérias coronárias constitui um importante desafio clínico pela variabilidade anatómica, possíveis repercussões funcionais, mecanismos fisiopatológicos implicados e também pela ausência de grandes séries na literatura que forneçam sólida evidência científica para a sua orientação clínica e terapêutica.

E-mail address: mariasalomecarvalho@gmail.com (M.S. Carvalho).

^a Centro Cardiovascular, Hospital da Luz, Lisboa, Portugal

^b Centro de Imagiologia, Hospital da Luz, Lisboa, Portugal

c Investigador do CEDOC/FCM-UNL, Centro de Estudos de Doenças Crónicas da FCM-UNL, Lisboa, Portugal

[†] Please cite this article as: Carvalho, S., et al. Origem anómala da coronária esquerda: variante interarterial maligna com evolução clínica benigna. Rev Port Cardiol 2012. doi:10.1016/j.repc.2011.11.010

^{*} Corresponding author.

456 M.S. Carvalho et al.

Os autores descrevem o caso de um doente de 55 anos, com antecedentes de hipertensão, dislipidemia e atrofia congénita da perna esquerda, que se apresentou com quadro de dor retroesternal atípica com um ano de evolução. Tendo em conta os fatores de risco cardiovasculares e exames complementares de diagnóstico previamente realizados, considerou-se ser um doente com probabilidade baixa a intermédia de doença coronária. Assim, e pela sua limitação funcional, realizou angioTC cardíaca que revelou uma origem anómala da coronária esquerda na cúspide coronária direita e com trajeto entre a aorta e a artéria pulmonar. Embora esta seja uma variante anatómica com potencial evolução maligna, optou-se por uma atitude conservadora após ponderação do risco-benefício no contexto clínico deste doente.

© 2011 Sociedade Portuguesa de Cardiologia. Publicado por Elsevier España, S.L. Todos os direitos reservados.

Introduction

Coronary artery anomalies are the third leading cause of sudden death in young athletes¹; their incidence in healthy individuals is estimated at 0.3-1%.²

Various different anomalies have been described, and classifications have been proposed based on anatomy (anomalies of origin, course and termination) and on prognosis (with or without clinical significance).² With regard to the latter, most anomalies have no clinical significance and thus require no specific treatment.³

However, congenital coronary artery fistulas, myocardial bridging and anomalous origin in the pulmonary artery can be associated with malignant evolution.

Another potentially serious anomaly is origin of a coronary artery from the contralateral coronary sinus, which can lead to myocardial ischemia and sudden death when it courses between the aorta and pulmonary artery (interarterial course). Other courses—retroaortic, prepulmonary and subpulmonary, and septal—are considered benign.^{4,5}

Origin of the left coronary artery in the right sinus of Valsalva is rare, with an estimated incidence of 0.09–0.11% in patients undergoing coronary angiography,⁶ but it is frequently associated with sudden cardiac death during intense physical activity and in adolescents and young adults, half of whom were previously asymptomatic.^{7,8} An interarterial course is seen in most of these patients⁹; their poor prognosis may be due to the acute angle of the artery from the ostium, stretching of the intramural segment, or compression between the commissure of the right and left coronary cusps.⁵

Cardiac CT angiography is a recent and valuable complementary exam that has high spatial resolution and provides three-dimensional views of the coronary arteries, with better definition of their origin and proximal portion, course, and relations with other cardiac structures than conventional angiography. The increasing popularity of this technique has greatly improved detection of coronary artery anomalies, which has made clinical decision-making more difficult, since there is, and is unlikely to be, sufficient evidence on which to base therapeutic strategies in doubtful cases such as the one presented.

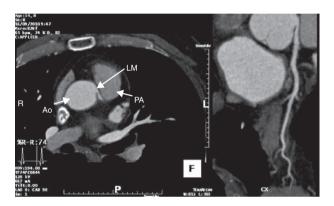


Figure 1 Multiplanar reconstruction. Ao: aorta; PA: pulmonary artery; LM: left main.

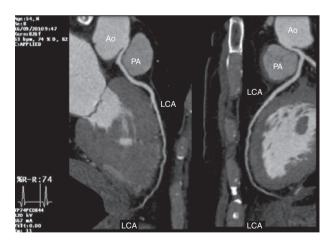


Figure 2 Multiplanar reconstruction. Ao: aorta; PA: pulmonary artery; LCA: left coronary artery.

Case report

A 54-year-old male patient was assessed in the cardiology outpatient clinic for atypical chest pain of about one year's evolution. He reported no other relevant cardiovascular symptoms, including fatigue, dyspnea, palpitations or syncope.

His cardiovascular risk factors included hypertension and dyslipidemia; he also had congenital left leg atrophy. He was

Download English Version:

https://daneshyari.com/en/article/3020775

Download Persian Version:

https://daneshyari.com/article/3020775

<u>Daneshyari.com</u>