

Transcatheter closure of right coronary artery fistula to the right ventricle

A. Abu Haweleh^a, Luna Baangood^{a,*}, J.V. DeGiovanni^a

^a Royal Medical Services, Queen Alia Heart Institute, Amman

^a Jordan

Coronary artery fistula (CAF) is an uncommon anomaly that is usually congenital but can be acquired. Although most patients are asymptomatic, some may present with congestive heart failure, infective endocarditis, myocardial ischemia or rupture. In the past, surgical ligation was the only option in the management of CAF, but since 1983, transcatheter closure of CAF has been increasing as an alternative to surgery. We report a 3-year-old boy, presented in Queen Alia Heart Institute, who underwent successful transcatheter closure of a large fistula communicating the distal part of the right coronary artery to the right ventricle. Our case differs from other CAFs in that the fistula was communicating the right coronary artery itself to the right ventricle.

© 2017 Production and hosting by Elsevier B.V. on behalf of King Saud University. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

Keywords: Coronary artery fistula, Right ventricle, Vascular plug

Introduction

Coronary artery fistula (CAF) is a direct communication between a coronary artery and a cardiac chamber, great vessels as well as vascular structure [1]. It is a rare disease of coronary arteries, which can cause a significant hemodynamic problems depending on the size, exit chamber and its relationship to the native coronary artery. It occurs in around 0.002% of general population [2].

Around 50% of CAF patients are asymptomatic, picked up with an incidental heart murmur

whereas others may present with acute myocardial ischemia, angina pectoris, and infective endocarditis [3].

In this case report, we present a percutaneous closure of right coronary artery fistula to the right ventricle in a 3-year-old male patient.

Case report

The patient was a 3-year-old boy, born at full term, one of identical twins following an uneventful pregnancy and cesarean delivery. He presented in the neonatal period with attack of tachypnea and feeding difficulties with normal

Disclosure: Authors have nothing to disclose with regard to commercial support.

Received 1 March 2016; revised 16 November 2016; accepted 26 January 2017.

* Corresponding author at: Royal Medical Services, Queen Alia Heart Institute, Amman, Jordan.
E-mail address: lunabaan@yahoo.com (L. Baangood).



P.O. Box 2925 Riyadh – 11461KSA
Tel: +966 1 2520088 ext 40151
Fax: +966 1 2520718
Email: sha@sha.org.sa
URL: www.sha.org.sa



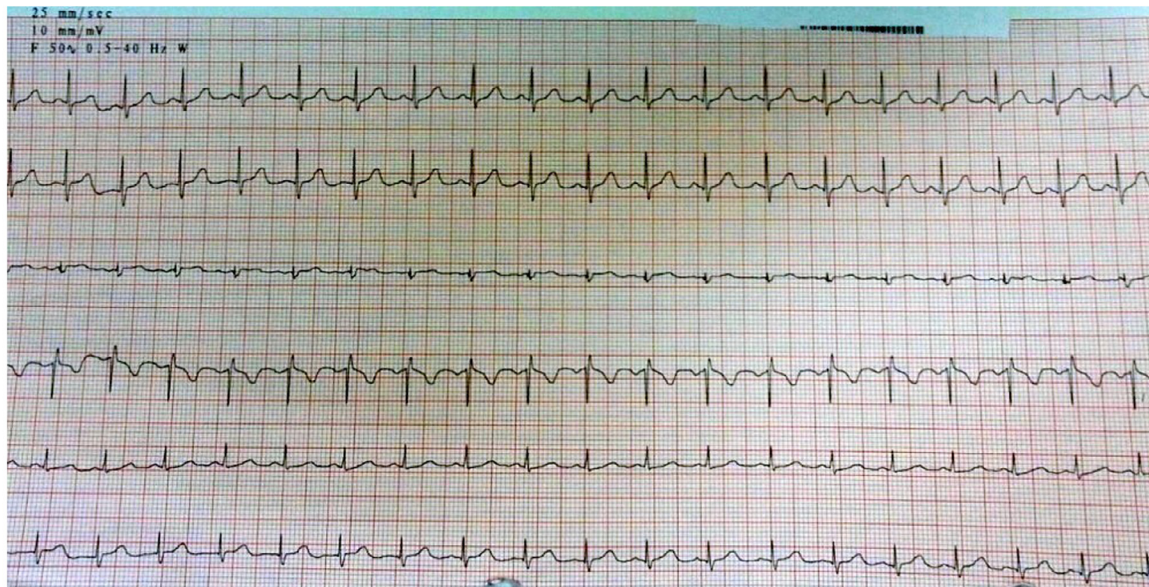
1016-7315 © 2017 Production and hosting by Elsevier B.V. on behalf of King Saud University. This is an open access article under the CC BY-NC-ND license (<http://creativecommons.org/licenses/by-nc-nd/4.0/>).

Peer review under responsibility of King Saud University.
URL: www.ksu.edu.sa
<http://dx.doi.org/10.1016/j.jsha.2017.01.005>



Production and hosting by Elsevier

(A)



(B)

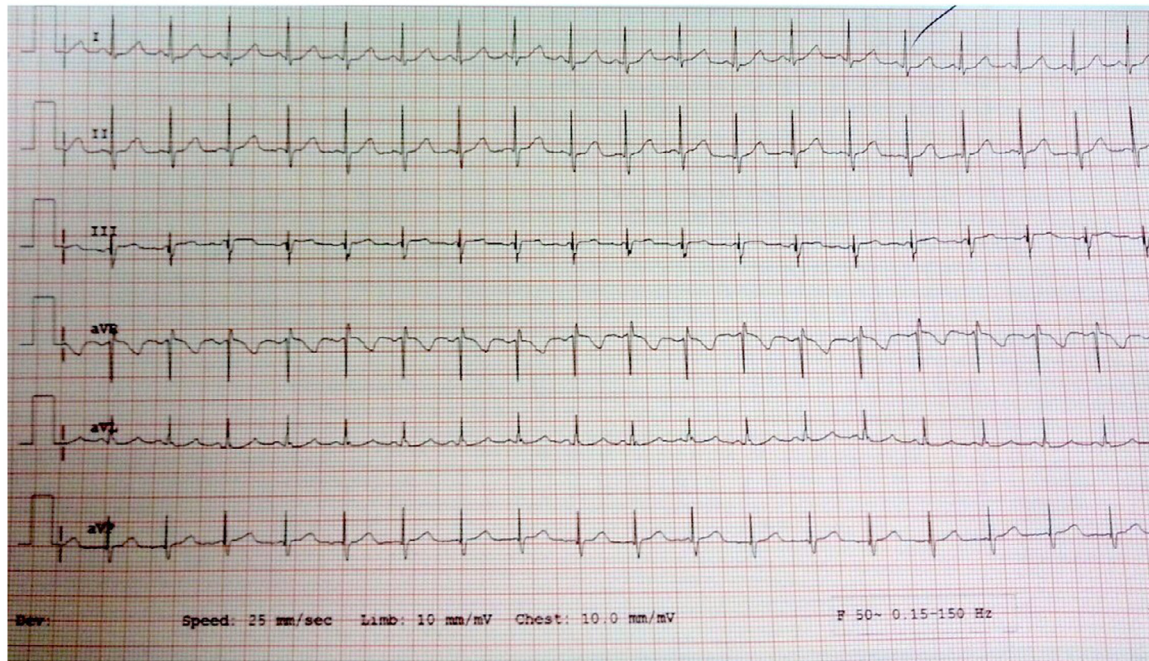


Figure 1. (A). Echocardiography before coronary artery fistula closure. (B) Echocardiography after coronary artery fistula closure, shows no changes in ST and T waves.

O₂ saturation. Cross sectional echocardiogram at the neonatal period showed right coronary artery (RCA) fistula to the right ventricle (RV). The patient was asymptomatic during the regular follow-up until age 3 years; he started complaining of effort intolerance with profuse sweating after minimal activity. Echocardiography (ECG) showed the same CAF with mild dilatation of the

right heart. The patient was started on furosemide and planned for cardiac catheter.

On clinical examination, the patient looked well, not in failure, with no visible impulse, and with soft S1 and S2 and continuous murmur grade 3/6, at the third intercostal space. ECG showed sinus tachycardia with no evidence of myocardial ischemia (Figure 1). Coronary arteries computed

Download English Version:

<https://daneshyari.com/en/article/8669809>

Download Persian Version:

<https://daneshyari.com/article/8669809>

[Daneshyari.com](https://daneshyari.com)