Giant major aortopulmonary collateral artery: A rare cause of heart murmur in newborns

Major aortopulmonary collateral arteries (MAPCAs) are anomalous vessels that develop from the aorta or its main branches and supply the pulmonary parenchyma with oxygenated blood. They are usually associated with congenital heart disease and are rare in isolation.

We report the case of a newborn girl, full term, with no relevant history, in whom a III/VI murmur was detected on the left sternal border on the first day of life. At seven days a remote cardiology consultation raised the suspicion of right inferior pulmonary vein stenosis despite normal pulmonary artery systolic pressure. Echocardiography in the Hospital Pediátrico Carmona da Mota showed "an anomalous vessel draining into the right inferior pulmonary vein".

At one month of age computed tomography angiography revealed a single MAPCA emerging from the celiac trunk to the right lung, draining into the right inferior pulmonary vein (Figure 1).

Diagnostic and therapeutic catheterization at five months showed normal systemic and pulmonary pressures. Angiography of the pulmonary artery revealed branches of reasonable caliber and normal venous return with mixed perfusion of the right inferior pulmonary field by the pulmonary artery and the MAPCA (Figure 2), while angiography of the aorta showed a wide collateral (7 mm), emerging from the celiac trunk to the right inferior lung (Figure 3). It was decided to embolize the collateral with a 10/7-mm Amplatzer Vascular Plug II, leaving a residual shunt (Figure 4).

The patient had a good clinical course with regular cardiological follow-up. She is currently two years old and has no shunt after three months of embolization, normal height, weight and development, and no clinical or echocardiographic signs of heart failure.


* Corresponding author.
E-mail address: lvmartins@hotmail.com (L. Martins).

2174-2049/$ - see front matter © 2014 Sociedade Portuguesa de Cardiologia. Published by Elsevier España, S.L.U. All rights reserved.
Figure 1  Computed tomography angiography (three-dimensional reconstruction). Green arrow: major aortopulmonary collateral artery (MAPCA) emerging from the celiac trunk.

Figure 2  Angiography of the pulmonary artery showing vascularization of the right inferior pulmonary artery branch (red arrow).

Figure 3  Angiography of the aorta showing giant MAPCA (red arrow) emerging from the aorta and vascularizing the lower right lung, and venous drainage (blue arrow) to the right inferior pulmonary vein.

Figure 4  Post-procedural angiography of the aorta showing Amplatzer Vascular Plug II in the MAPCA.
Giant major aortopulmonary collateral artery: A rare cause of heart murmur in newborns

Ethical disclosures

Protection of human and animal subjects. The authors declare that no experiments were performed on humans or animals for this study.

Confidentiality of data. The authors declare that they have followed the protocols of their work center on the publication of patient data.

Right to privacy and informed consent. The authors have obtained the written informed consent of the patients or subjects mentioned in the article. The corresponding author is in possession of this document.

Conflicts of interest

The authors have no conflicts of interest to declare.