Andreas S. Triantafyllis,^{a,*} Andreas S. Kalogeropoulos,^b Keir McCutcheon,^a Walter Desmet,^a and Johan Bennett^a

^aDepartment of Cardiovascular Medicine, University Hospitals Leuven, Leuven, Belgium ^bDepartment of Cardiology, Barts NHS Trust, London, United Kingdom

* Corresponding author: E-mail address: andtridoc@yahoo.gr (A.S. Triantafyllis).

Available online 11 November 2017

REFERENCES

 Shovlin CL, Awan I, Cahilog Z, Abdulla FN, Guttmacher AE. Reported cardiac phenotypes in hereditary hemorrhagic telangiectasia emphasize burdens from arrhythmias, anemia and its treatments, but suggest reduced rates of myocardial infarction. *Int J Cardiol.* 2016;215:179–185. http://dx.doi.org/ 10.1016/j.ijcard.2016.04.006.

- Clark K, Pyeritz RE, Trerotola SO. Angina pectoris or myocardial infarctions, pulmonary arteriovenous malformations, hereditary hemorrhagic telangiectasia, and paradoxical emboli. Am J Cardiol. 2013;112:731–734.
- Mieghem CA, Ligthart JM, Cademartiri F. Spontaneous dissection of the left main coronary artery in a patient with Osler-Weber-Rendu disease. *Heart.* 2006;92: 195–196.
- Talha S, Brandt C, Maamari G, Mossard JM, Germain P, Andres E. Myocardial infarction in a patient with normal coronary arteries and hereditary haemorrhagic telangiectasia. *QJM*. 2006;99:195–196.
- Jerkic M, Letarte M. Contribution of oxidative stress to endothelial dysfunction in hereditary hemorrhagic telangiectasia. Front Genet. 2015;6:. http://dx.doi.org/ 10.3389/fgene.2015.00034.
- Fernández LA, Sanz-Rodriguez F, Blanco FJ, Bernabéu C, Botella LM. Hereditary hemorrhagic telangiectasia, a vascular dysplasia affecting the TGF-beta signaling pathway. *Clin Med Res.* 2006;4:66–78.

https://doi.org/10.1016/j.rec.2017.10.010

1885-5857/

© 2017 Sociedad Española de Cardiología. Published by Elsevier España, S.L.U. All rights reserved.

Pulmonary Valve Perforation and Stent Implantation in the Right Ventricle Outflow Tract Using a Hybrid Procedure as an Alternative to a Central Shunt

Perforación de válvula pulmonar e implante de stent en el tracto de salida del ventrículo derecho por procedimiento híbrido como alternativa a la fístula central

To the Editor,

Pulmonary atresia with a ventricular septal defect is a complex congenital condition associated with a poor prognosis in patients with hypoplastic or nonexistent pulmonary branches and large aortopulmonary collateral vessels. In this situation, a staged strategy is preferred to provide pulmonary flow as promptly as possible and achieve growth and development of the pulmonary arteries before corrective surgery is undertaken.¹ The most commonly used traditional palliative surgery techniques were a central systemic-to-pulmonary shunt¹ and placement of a transannular patch or a synthetic conduit between the right ventricle and the pulmonary artery. Later, a percutaneous approach was described, consisting of pulmonary valve perforation by radiofrequency² and stent implantation in the right ventricular

outflow tract, as an alternative to surgery. A hybrid approach is used in low-weight newborns, such as the patient described here.

A term neonate weighing 2.6 kg was diagnosed with a double outlet right ventricle and an unrelated ventricular septal defect, overriding tricuspid valve, and pulmonary atresia, with compensatory circulation through multiple aortopulmonary collateral vessels. The pulmonary trunk and both branches were hypoplastic (trunk, 2.5 mm; branches, 1.3 mm). A central shunt and partial correction were ruled out, and a decision was taken to perform a hybrid procedure consisting of a surgical sternotomy and perforation of the pulmonary valve, with periventricular stent placement. Periventricular puncture of the right ventricle was carried out using a 22-G needle. A 0.018-inch guidewire was inserted, enabling passage of a 5-Fr Cook radial introducer, which was placed just below the pulmonary valve (Figure A and video 1 of the supplementary material). Contrast material was injected and minimal anterograde pulmonary flow was observed. Passage of a 0.014-inch Whisper coronary guidewire (Abbot) was achieved and the introducer was then advanced through the dilator. Patency of the pulmonary trunk and branches was verified. A premounted 6×16 -mm Formula stent (Cook) was then implanted in the pulmonary valve. The procedure was successful, achieving adequate perfusion of the pulmonary tree and showing hypoplasia of the left pulmonary branch (Figure B



Figure. A, Right ventriculogram with cranial angulation in the posteroanterior view shows minimal anterograde pulmonary flow and the confluence of the trunk and the hypoplastic pulmonary branches. B, Pulmonary arteriogram with cranial angulation in the posteroanterior view depicts correct stent placement and adequate perfusion of both pulmonary branches, although with hypoplasia of the left pulmonary branch. C, The posteroanterior pulmonary arteriography image with cranial angulation at the second catheterization shows that both pulmonary branches have grown and there is adequate perfusion of the lung segments.

and video 2 of the supplementary material). The patient was transferred to the intensive care unit, but extubation was not possible because of excessive pulmonary blood flow. A new catheterization was carried out 12 days after the first, and growth of both pulmonary branches was observed (Figure C and video 3 of the supplementary material). The collaterals were successfully embolized with Interlock coils (Boston), thereby allowing extubation and hospital discharge at 20 days following the second procedure. At 6 months of follow-up, the patient was asymptomatic, with arterial oxygen saturation at 82%, and awaiting a partial cavopulmonary shunt.

Treatment of pulmonary atresia with a ventricular septal defect, hypoplastic pulmonary arteries, and major aortopulmonary collateral vessels remains a challenge. The conventional surgical approach is a systemic-to-pulmonary artery shunt, which, in lowweight neonates, may imply major complications, such as shunt obstruction or thrombosis, pulmonary hyperflow, and infection.¹ Another surgical option is partial correction, which requires the use of extracorporeal circulation and can lead to greater morbidity and mortality.

In the cardiac catheterization laboratory, radiofrequency perforation of the pulmonary valve with stent implantation in the right ventricular outflow tract has become a valid option for treating patients with pulmonary atresia, although it is associated with a need for follow-up surgery in 33% to 75% of patients.³ The most commonly described complications related to this technique are perforation of the cardiac wall and fracture or dislocation of the stent, among others.⁴

The hybrid procedure approach averts the need for extracorporeal circulation and is safer and faster than an interventional procedure, as it allows adequate anatomical inspection and enables the cardiovascular surgeon to act promptly if there are incidents. Other advantages are a shorter fluoroscopy time and no weight limitations in neonatal patients.⁵ Therefore, it can be concluded that valve perforation with stent implantation through the transventricular route using a median sternotomy is an effective palliative technique, especially for low-weight premature infants, that offers results similar to those of other therapeutic strategies with lower in-hospital morbidity and mortality rates and satisfactory long-term survival (Table, supplementary material). This technique achieves adequate development of the pulmonary arterial tree, so that corrective surgery can be performed in a second stage when the patient is older and has gained weight.⁶

SUPPLEMENTARY MATERIAL



Supplementary material associated with this article can be found in the online version available at doi:10.1016/j. rec.2017.10.032.

Gertrudis Parody-Cuerda,^{a,*} Félix Coserria-Sánchez,^b Alejandro Adsuar Gómez,^c Juan Antonio García-Hernández,^d Begoña Manso-García,^b and A. Reza Hosseinpour^c

^aSección de Cirugía Cardiovascular, Hospital Universitario
Virgen Macarena, Sevilla, Spain
^bSección de Cardiología Pediátrica y Hemodinámica Infantil, Hospital
Universitario Virgen del Rocío, Sevilla, Spain
^cServicio de Cirugía Cardiovascular, Área del Corazón, Hospital
Universitario Virgen del Rocío, Sevilla, Spain
^dServicio de Cuidados Críticos y Urgencias Pediátricas, Hospital
Universitario Virgen del Rocío, Sevilla, Spain

* Corresponding author:

E-mail address: gerparcue@gmail.com (G. Parody-Cuerda).

Available online 7 December 2017

REFERENCES

- 1. Watterson KG, Wilkinson JL, Karl TR, Mee RBB. Very small pulmonary arteries: centralend-to-side shunt. Ann Thorac Surg. 1991;52:1132–1137.
- Medrano C, Guía JM, Rueda F, Moruno A. Actualización en cardiología pediátrica y cardiopatías congénitas. *Rev Esp Cardiol*. 2009;62:39–52.
- Zampi JD, Hirsch-Romano JC, Goldstein BH, Shaya JA, Amstrong AK. Hybrid approach for pulmonary atresia with intact ventricular septum: early single center results and comparison to the standard surgical approach. *Catheter Cardiovasc Interv.* 2014;83:753–761.
- 4. Cools B, Boshoff D, Heying R, Rega F, Meyns B, Gewillig M. Transventricular balloon dilation and stenting of the RVOT in small infants with tetralogy of Fallot with pulmonary atresia. *Catheter Cardiovasc Interv.* 2013;82:260–265.
- Park JY, Seo DM, Shin HJ, Kim SJ, Son JS. Hybrid procedure for pulmonary atresia with ventricular septal defect in a low birth weight neonate. *Korean J Thorac Cardiovasc* Surg. 2013;46:56–59.
- 6. Zampi JD, Armstrong AK, Hirsch-Romano JC. Hybrid perventricular pulmonary valve perforation and right ventricular outflow stent placement: a case report of a premature, 1.3-kg neonate with tetralogy of Fallot and pulmonary atresia. *World J Pediatr Congenit Heart Surg.* 2014;5:338–341.

https://doi.org/10.1016/j.rec.2017.10.032

1885-5857/

© 2017 Sociedad Española de Cardiología. Published by Elsevier España, S.L.U. All rights reserved.

Volumetric Quantification of Coronary Flow by Using a Monorail Infusion Catheter: Initial Experience

Cuantificación volumétrica de flujo coronario mediante catéter de infusión monorraíl: experiencia inicial

To the Editor,

Absolute volumetric coronary blood flow can now be measured invasively by thermodilution due to a new purposely designed monorail microcatheter that allows homogeneous and continuous mixing of room-temperature saline solution with intracoronary blood.¹ This microcatheter (Rayflow; Hexacath) has 4 openings at its distal end, and the terminal distal opening is occluded by the guidewire, permitting homogeneous infusion.¹ The intracoronary guidewire has a temperature sensor linked to dedicated software (RadiView; Abbott), allowing thermodilution-based measurements of the maximal volume of intracoronary blood per unit of time, according to the formula:

Qb = 1.08Ti/TQi

where T_i is the temperature of the infused saline exiting the infusion catheter, T is the temperature of the blood-saline mixture in the distal portion of the coronary artery, and Q_i is the preset saline infusion rate through the microcatheter (Figure).¹ This procedure can yield quantitative estimates not only of the maximal coronary blood flow (MCF), but also of the corresponding minimal microvascular resistance in the irrigated territory, since this is the ratio of coronary pressure to coronary blood flow, both of which are known.

We prospectively recruited 14 patients in whom quantitative coronary angiography revealed no significant coronary lesions (stenosis < 30% of the luminal diameter). In most study