Patent ductus arteriosus stenting: ductal dissection and spasm in pulmonary atresia with intact ventricular septum

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Abstract
Ductal dissection is rarely mentioned but intractable complication of ductal stenting. We demonstrated the dissection image under fluoroscopy and the use of a relatively large stent as an applicable treatment.

A newborn (gestational age 37 + 5 days, birth weight 2676 g) was found to have cyanosis (pulse oximetry 75%) upon birth. Echocardiography resulted in diagnoses of pulmonary atresia with intact ventricular septum, right ventricle hypoplasia, and patent ductus arteriosus. Continuous prostaglandin E1 was administered immediately. CT angiography confirmed the diagnosis. Transcatheter valvotomy with sequential balloon valvuloplasty was performed at 7 days old. Through the femoral vein approach, a 0.014-inch wire (percutaneous transluminal coronary angioplasty guide wire, Sion) was advanced through the Judkins right guiding catheter to the right ventricle outflow tract, and then a microcatheter (Excelsior) was introduced to the right ventricle outflow tract as a supporting system. Pulmonary valvotomy was performed using coronary guidewire (Conquest Pro 8-20 wire) penetration, and then sequential balloon valvuloplasty was conducted. Saturation was improved to pulse oxymetry 90% by the procedure. Nevertheless, desaturation occurred when prostaglandin E1 administration was tapered. Recatheterization for patent ductus arteriosus stenting was performed at 20 days old. Prostaglandin E1 was discontinued before the procedure. Through the femoral artery approach, the 0.014-inch floppy run-through wire with Judkins right catheter guiding was advanced from the descending aorta side of the patent ductus arteriosus (Fig 1a). The wire could not smoothly pass the patent ductus arteriosus to the pulmonary artery side, with mild distortion occurring at the tip of the wire upon fluoroscopy (Fig 1b, Video 1). Contrast extravasation was found (Fig 1c, Video 2). As ductal dissection was diagnosed, the wire was withdrawn. Desaturation was noted (pulse oximetry reduced to 60%). Repeat angiography showed severe ductal spasm. Readministration of PGE1 (20 ng/kg/minute) was performed. The haemodynamics were stable, and the saturation returned to 85% after 20 min of prostaglandin E1 infusion. Finally, a patent ductus arteriosus stent (Energy 5 mm × 18 mm) was implanted (Fig 1d, Video 3). After oversized patent ductus arteriosus stenting, the patient had tachypnea and pulmonary congestion on chest radiography was noted. Diuretics was used (Furosemide 2 mg/kg/day), and the symptom improved after medication and gradually resolved. No further cyanosis episodes occurred, and the patient was soon discharged. The diuretics was then tapered and discontinued 2 months after procedure.

Conclusion
Compared with ductal spasm, ductal dissection is rarely mentioned in the literature. Ductal dissection mostly occurs when ducts are crossed by a stiff guidewire.1,2 However, even a soft wire may cause the complication. Use of a relatively large stent is an applicable treatment of such cases.
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References
