In those cases in which the parallel guidewire technique is required, the ReCross microcatheter allows the insertion of a second CTO guidewire through the hub of the stylet lumen in order to re-engage the cap. By using the ReCross device operators can advance simultaneously 2 CTO guidewires through 2 different OTW lumens, somehow similar to the pioneering see-saw wiring technique with 2 SLMs.

Antegrade dissection and re-entry (ADR): ADR techniques are characterized by the intentional use of the subintimal space to cross coronary CTOs followed by the subsequent re-entry into the distal true lumen. Several devices have been developed to facilitate a controlled ADR (CrossBoss microcatheter and the Stingray balloon; Boston Scientific, United States). The main limitations of these devices are their costs and crossing-profile, which often requires prior balloon dilatation with the corresponding increase of subintimal hematomas. Conversely, the ReCross microcatheter can be advanced into the subintimal space distally to the occlusion often without the need for vessel pre-dilation to perform a subintimal guidewire redirection. The operator can advance a stiff guidewire through the appropriate lumen to perform a controlled re-entry puncture from the subintimal space towards the true lumen.

Additionally, with the ReCross device it is possible to use 2 CTO guidewires simultaneously to achieve re-entry from the subintimal space towards the true lumen with an high success rate. Moreover, the first lumen can be used for vessel decompression of the subintimal hematoma, thus facilitating the re-entry of the second guidewire. In conclusion, the ReCross device provides a versatile and attractive alternative to standard DLM when performing the PCI of a CTO potentially reducing procedural costs and time.

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https://doi.org/10.24875/RECICE.M21000231

**Percutaneous edge-to-edge tricuspid valve repair in congenitally corrected transposition of the great arteries**

*Reparación percutánea borde a borde de la válvula tricúspide en transposición de grandes vasos congénitamente corregida*

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To the Editor,

The congenitally corrected transposition of the great arteries is a rare congenital defect characterized by ativoventricular and ventriculoarterial discordance. As a result, the tricuspid valve and the anatomical right ventricle sustain the systemic circulation. Typically, the patient remains asymptomatic at an early age, but the right ventricle and the tricuspid valve deteriorate with the passing of time. The only curative treatment for this condition is heart transplant. In this setting, percutaneous edge-to-edge tricuspid valve repair has been traditionally used to treat tricuspid regurgitation in patients who are ineligible for heart transplantation; however, to this date, the evidence available is scarce and based on case reporting in heterogeneous clinical settings. 1-3

This is the case of a young male patient with congenitally corrected transposition of the great arteries, advanced heart failure, and torrential tricuspid regurgitation considered ineligible for heart transplant.

**AUTHORS’ CONTRIBUTIONS**

R. Garbo, M. Iannaccone, J. Sanz Sánchez, and G.L. Gasparini contributed to the design, analysis, and writing of this manuscript. J.A.Orellia, and A.Gagnor contributed to the design, and writing of this manuscript too.

**CONFLICTS OF INTEREST**

None reported.

**FUNDING**

None reported.

**AUTHORS’ CONTRIBUTIONS**

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Online: 09-08-2021.

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transplantation due to irreversible severe pulmonary hypertension, but eligible for percutaneous edge-to-edge tricuspid valve repair. The patient signed an informed consent form authorizing the publication of his case that was eventually approved by our center ethics committee.

This is the case of a male diagnosed with congenitally corrected transposition of the great arteries and congenital atrioventricular block at the early age of 7 months. The patient remained asymptomatic until he was 29 years-old when he required a pacemaker due to presence of chronotropic incompetence. Afterwards, he was lost to follow-up until he was admitted to the intensive care unit with signs of pulmonary edema at the age of 35 when he was diagnosed with biventricular systolic dysfunction, severe systemic atrioventricular valve regurgitation, and pulmonary hypertension. Due to the occurrence of a cardiac arrest, an implantable cardioverter-defibrillator with resynchronization therapy was indicated followed by the optimal medical therapy.

Despite treatment, the patient remained symptomatic with New York Heart Association functional class III, and INTERMACS 4. The echocardiographic assessment revealed the presence of severe systemic ventricular systolic dysfunction (right ventricular ejection fraction = 35%) with severe regurgitation of the systemic atrioventricular valve. The valve showed an Ebstein-like anomaly (8.3 mm²/m²), abnormal chordae structures, and thickened leaflets with restriction of motion causing a wide coaptation defect, mainly between the septal and posterior leaflets triggering torrential regurgitation (V/V) (Figure 1). Cardiac catheterization revealed the presence of severe pulmonary hypertension (mean pulmonary artery pressure of 55 mmHg) with pre- and post-capillary components (transpulmonary gradient of 30 mmHg, and pulmonary vascular resistance of 6.4 WU). The vasodilator test with nitric oxide resulted in a maximum response, but without any significant changes. Considering all this information, the heart team decided that the patient remained ineligible for heart transplantation and suggested the percutaneous edge-to-edge tricuspid valve repair of the tricuspid valve with a MitraClip device (Abbott Vascular, United States) as palliative treatment.

During the procedure the presence of torrential tricuspid regurgitation was confirmed (Figure 2A,B) with consistent invasive...
hemodynamic findings [figure 2C]. An early MitraClip XTR device was implanted at the origin of the regurgitant jet at the center of the septal-posterior coaptation line that was able to reduce regurgitation significantly. However, due to the presence of moderate-to-severe persistent regurgitation (II-III/V) without significant stenosis a second device had to be implanted between the anterior and septal valves for being the area with the greatest residual regurgitation. The outcome assessment confirmed the presence of mild-to-moderate residual tricuspid regurgitation (II/V) [figure 2D,E] without stenosis. This positive outcome was also confirmed on the invasive assessment [figure 2F], which is why the procedure was considered terminated.

Despite the slight worsening of tricuspid regurgitation at the 6-month follow-up (grade III/V), and the presence of systemic ventricular dysfunction (right ventricular ejection fraction = 35%) and severe pulmonary hypertension (pulmonary artery systolic pressure > 60 mmHg) the patient showed a maintained functional class improvement (New York Heart Association II). Also, the values of the amino-terminal fraction of B-type brain natriuretic propeptide dropped significantly (from 9787 pg/mL to 2083 pg/mL), and fewer diuretics were required. This translated into a significant improvement of the patient’s quality of life, a better functional capacity, and no rehospitalizations 1 year after the procedure.

This case reinforces the role of percutaneous edge-to-edge tricuspid valve repair even in such an adverse setting as the congenitally corrected transposition of the great arteries. Very few case reports have previously described this indication, and always in more favorable clinical situations. Our case has various technical limitations that make it extra interesting, especially the presence of a dysplastic valve with significant restriction of motion, and a large coaptation defect. Therefore, the largest device available was used to target the area with greater regurgitation (septal-posterior). Deep leaflet capture followed due to the significant bulge present even at the risk of inadvertently capturing the chordae tendineae. However, a second device was required to reduce tricuspid regurgitation significantly. Despite the inherent empiricism of the clinical situation and the unfavorable hemodynamic conditions described with severe pulmonary hypertension and severe systemic right ventricular failure, the patient improved significantly and consistently through time regardless of the lack of improvement reported in the numbers of pulmonary artery pressure and right ventricular ejection fraction. That is why we believe that the maintained reduction of tricuspid regurgitation had a clinical impact. In this sense, pulmonary hypertension has been reported to improve in 1 case only, which allowed to reassess the eligibility of heart transplantation. Therefore, percutaneous edge-to-edge tricuspid valve repair should be considered an effective option in patients with congenitally corrected transposition of the great arteries who remain ineligible for heart transplantation or surgery.

FUNDING
None whatsoever.

AUTHORS’ CONTRIBUTIONS
A. Salinas Gallegos, and E. Pozo Osinalde: study design, and writing of the manuscript. L. Nombela-Franco, P. Jiménez Quevedo, and R. Estevez-Loureiro: management of the patient, and manuscript review. J.A. de Agustín: manuscript review.

CONFLICTS OF INTEREST
L. Nombela-Franco, and R. Estévez-Loureiro are consultors, and proctors, and have received speaking fees from Abbott Vascular, Edwards Lifesciences, and Boston Scientific.

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