The ReCross dual-lumen microcatheter versatility during percutaneous coronary intervention of chronic total coronary occlusions

Versatilidad del microcatéter ReCross durante la angioplastia de oclusiones coronarias crónicas

To the Editor,

Microcatheters are essential tools to facilitate guidewire manipulation and exchange, thus enhancing the guidewire penetration force.1 The ReCross over-the-wire dual-lumen microcatheter [DLM] (IMDS, The Netherlands) is the latest evolution among DLM microcatheters. However, no data are available in the medical literature yet. From September 2020 through November 2020 a total of 8 patients undergoing percutaneous coronary intervention (PCI) of chronic total coronary occlusions (CTO) with the ReCross at 5 Italian high-volume PCI-capable centers were retrospectively identified. This study complied with the Declaration of Helsinki, and written informed consent was obtained from all participants.

Case 1. A proximal right coronary artery CTO underwent antegrade approach using the ReCross microcatheter (IMDS, The Netherlands) as a first choice due to the presence of a bifurcation at distal cap level (figure 1A). The antegrade wire escalation technique was used with an Ultimate Bros 3 guidewire (Asahi Intecc, Japan) followed by a Gaia Second guidewire (Asahi Intecc, Japan) that was able to cross the CTO body and reach the side branch (right ventricular branch) distal true lumen (figure 1B). The ReCross was advanced over the Gaia Second guidewire through the lesion close to the bifurcation; afterwards, another Gaia Second guidewire was advanced through the blue lumen and easily directed towards the distal main vessel (figure 1C). The ReCross was then advanced as a single-lumen microcatheter [SLM] into the distal right coronary artery true lumen.

Case 2. The CTO of a mid-left anterior descending coronary artery (LAD) underwent antegrade approach with the ReCross as the single-lumen microcatheter using the antegrade wire escalation strategy. However, all the guidewires went subintimal and finally the successful recanalization of the occluded artery was achieved.

Figure 1. Bifurcation at distal cap level. A: the first chronic total coronary occlusion guidewire (Gaia Second) went into the side branch; B: a second chronic total coronary occlusion guidewire (Gaia Second) was inserted through the hub of the stylet lumen to engage the main vessel; C: a contralateral injection confirmed the correct positioning of the second guidewire into the main distal true lumen.
system for the tip lumen; b) an additional exit port (the third one) in the tip lumen at a 180° angle of the exit port of the stylet lumen. The second OTW lumen gives the possibility of exchanging and using 2 different guidewires simultaneously and the additional exit port facilitates the redirection of the guidewire [figure 2]. In the following paragraphs we detail the versatile use of the ReCross in different anatomical settings during the PCI of a CTO.

Combination of CTOs and bifurcations: the presence of bifurcation lesions in the context of a CTO can be one of the most complex subsets during the PCI. In case of a bifurcation close to the proximal cap, the ReCross can be selected as a frontline microcatheter as the procedure is expected to require dual guidewire access. The ReCross is advanced over the workhorse guidewire inside the side branch until the proximal cap. Then, a CTO dedicated guidewire can be advanced through the second OTW lumen to negotiate the occlusion. This technique allows precise manipulations of the CTO guidewire and increases support to be able to penetrate the proximal cap. Notably, when the use of intravascular ultrasound-guided puncture is required, the small profile of the ReCross allows the simultaneous use of both devices in a 7-Fr guiding catheter with a large lumen. After successful puncture of the proximal cap the ReCross microcatheter should be removed from the side branch using the trapping technique. Afterwards, it can be re-advanced as a SLM to support the guidewire advance into the distal target true lumen. Finally, when facing a bifurcation inside the CTO body like at distal cap level, the use of the ReCross can be essential for the CTO guidewires to engage the side branch. In this scenario, the ReCross can be advanced at bifurcation level and a second CTO guidewire can be inserted through the second OTW lumen to engage the main distal vessel (case 1).

Unintentional antegrade guidewire subintimal tracking: in many antegrade procedures when the first guidewire goes subintimal, the most useful strategy is to use the parallel guidewire technique. The first guidewire is left in place as a marker occluding the false lumen and modifying the anatomy of the vessel. The first microcatheter must be replaced by a DLM, which is advanced over the subintimal guidewire. Afterwards, a second dedicated CTO guidewire can be used to re-engage the cap for intentional intimal plaque tracking.

### Table 1. Case series

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ADR, antegrade dissection and re-entry; CTO, chronic total coronary occlusion; DLM, dual-lumen microcatheter; LAD, left anterior descending coronary artery; LCX, left circumflex artery; RCA, right coronary artery.
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-dilatation 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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**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. Oreglia, and A. Gagnor contributed to the design, and writing of this manuscript too.

**CONFLICTS OF INTEREST**

None reported.

**REFERENCES**


The congenitally corrected transposition of the great arteries is a rare congenital defect characterized by atrophicventricular 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. 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

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**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**

Alejandra Salinas Gallegos, Eduardo Pozo Osinalde, Luis Nombela-Franco, Pilar Jiménez Quevedo, Rodrigo Estévez-Loureiro, and José Alberto de Agustín

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