Percutaneous coronary intervention of a tortuous and complex circumflex lesion using the robotic CorPath GRX system

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Aleksander Zelias, MD, Center for Invasive Cardiology, Electrotherapy and Angiology, Kilińskiego 68, 33–300 Nowy Sącz, Poland, phone: +48 18 540 02 00. e-mail: aa.zelias@gmail.com Copyright by the Author(s),

Kardiol Pol. 2021; 79 (9): 1044–1045; DOI: 10.33963/KP.a2021.0057

Received:

2021

April 22, 2021

Revision accepted: July 5, 2021

Published online: July 6, 2021 Robotic-assisted percutaneous coronary interventions (R-PCI) dramatically reduce physician radiation exposure and potential musculo-skeletal injuries [1]. In addition, accumulating evidence has demonstrated R-PCI safety and efficacy in a broad range of lesion types [2, 3].

We report the first case of R-PCI performed in Poland using the CorPath GRX (Corindus Vascular Robotics) system (Supplementary material, *Figures S1* and *S2*) to treat a complex tortuous lesion of the left circumflex (Cx) artery.

A 47-year-old male with previous PCI to the left anterior descending artery (LAD) was referred with worsening typical angina (Canadian Cardiovascular Society class III). Echocardiography revealed hypokinesia in the basal and mid-segments of the inferior and posterior wall. Diagnostic angiography revealed a short left main stem (LMS) and a tortuous circumflex (Cx) artery with a critical lesion in the mid-vessel (Figure 1A) associated with a separate critical lesion in the 1st obtuse marginal (OM) branch (Figure 1B). The right coronary artery was hypoplastic and the LAD stent was patent with no significant other lesions. A provisional strategy was planned to treat the disease in the Cx/OM1.

Six Fr right radial access was secured and the operator manually cannulated the LMS with a 6 Fr AL1 guiding catheter. Following successful and stable cannulation, the guiding catheter was connected to the robotic arm, and the rest of the procedure was completed from the remote workstation (Supplementary material, *Figure S3*). A Runthrough NS Floppy wire (Terumo systems, Somerset, NJ, USA)

was selected and robotically advanced to the distal vessel, with the tortuosity and diseased segment successfully navigated using a combination of manual joystick controls and pre-set automation techniques (Figure 1C). A 2.5×27 mm non-compliant (NC) balloon was used to pre-dilate the lesion. Precise measurement (1 mm precision) of the lesion length was performed using the robotic system and accordingly a 3.0 × 38 mm drug-eluting stent (Promus PREMIER, Boston Scientific, Marlborough, MA, USA) was advanced and successfully implanted at the intended site (Figure 1D). Post-dilatation of the proximal portion of the stent was performed with a 3.5×15 mm NC balloon. Following main vessel stenting, the main vessel wire was retracted and robotically advanced into the OM1 branch which was treated with balloon-only angioplasty using a 2.0×12 mm NC inflated at 10 atm (Figure 1E).

Final angiography revealed a good angiographic result, optimal stent expansion with no complications (Figure 1F). Fluoroscopy time was 22 minutes, radiation dose was 943 mGy and total contrast volume was 150 ml.

This case demonstrates how the R-PCI system can be used to safely and successfully treat complex lesions. Despite the lack of haptic feedback, wiring of this tortuous vessel was achieved using the joystick controls manually aided by the built-in automated robotic movements. The CorPath GRX system can accommodate multiple coronary wires and devices simultaneously. In such instances, one wire and one device are allocated to the active

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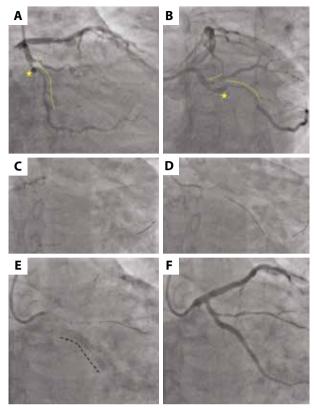


Figure 1. Robotic-assisted percutaneous coronary interventions (R-PCI) of the tortuous complex circumflex lesion. Baseline angiography (A-B) demonstrates the critical lesions (yellow dotted lines) in the mid circumflex artery and proximal segment of the 1st marginal branch. The tortuous segment of the vessel (yellow star) arises just before the critical lesion in the mid-vessel. Through a 6 Fr amplatz left guiding catheter a 0.014" coronary wire was advanced distally (C) using the robotic controls and following pre-dilatation, the stent was advanced and deployed (D-E) in the intended position (dashed black line). The marginal lesion was wired and treated with balloon angioplasty (E) using the robotic controls. The final result (F) was optimal without any angiographic complications

drive and can be controlled from the console, whilst the remaining wires and devices cannot be maneuvered but remained fixed in the passive drive. This can enable operators to treat complex lesions including bifurcations and perform final kissing inflations when required. The presence of a short LMS required repeated repositioning and stabilization maneuvers of the guiding catheter, which were all performed using the guide catheter joystick control. During initial wiring, the guide catheter disengaged into

the aorta with subsequent loss of wire position. With the robotic controls, the guide catheter was safely manipulated back into a more stable position achieving semi-selective cannulation of the Cx artery.

Worldwide experience with R-PCI systems is growing, enabling increasingly complex coronary lesions to be treated safely and effectively, without compromising procedural time, and with improved operator safety [4, 5]. In our case, the primary operator completed the entire procedure without wearing any radioprotection sat at the robotic console, which was located within the operating room.

Supplementary material

Supplementary material is available at https://journals.viamedica.pl/kardiologia_polska.

Article information

Conflict of interest: None declared.

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How to cite: Zelias A, Khokhar AA, Proniewska K, et al. A Percutaneous coronary intervention of a tortuous and complex circumflex lesion using the robotic CorPath GRX system. Kardiol Pol. 2021; 79(9): 1044–1045, doi: 10.33963/KP.a2021.0057.

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