

Challenging treatment of in-stent restenosis in a coronary bifurcation by implantation of a bioresorbable scaffold under optical coherence tomography guidance

Grzegorz Zuk¹, Dariusz Ciecwierz¹, Piotr Drewla¹, Marcin Gruchala¹,
Juan Luis Gutiérrez-Chico^{2,3}, Milosz J. Jaguszewski^{1,3}

¹Interventional Cardiology, 1st Department of Cardiology, University Clinical Center, Gdansk, Poland

²Punta de Europa University Hospital, Algeciras (Cádiz), Spain

³Institute of Cardiovascular Translational Research of the Atlantic (ICTRA), Berlin, Germany

This paper was guest edited by Prof. Marek Koziński

A 67-year-old male patient with stable angina, hypertension and hypercholesterolemia who underwent bare metal stent (BMS) implantation in the distal right coronary artery (RCA) (Azule 3 × 9 mm) and everolimus-eluting stent (EES) implantation in the first diagonal branch (D1) (Xience 2.25 × 18 mm) and in the proximal circumflex branch (LCx) (Xience 3 × 28 mm). One year subsequent to the procedure the patient was readmitted for relapse of the angina Canadian Cardiovascular Society scale II, exhibiting a positive exercise test. The coronary angiography showed a distal-edge in-stent restenosis (ISR) in the distal RCA, extending to the posterior descending artery (PDA), Medina 110 bifurcation (Fig. 1A). Optical coherence tomography (OCT) showed predominantly fibro-lipidic restenotic tissue, with minimal lumen area (MLA) 1.95 mm², minimal lumen diameter (MLD) 1.57 mm, proximal reference vessel diameter (RVD) 3.1 mm, distal RVD 2.75 mm and lesion length 21.2 mm (Fig. 1B, C).

Optical coherence tomography-guided implantation of a bioresorbable scaffold (BRS) to treat the bifurcation ISR was performed through a radial approach, using a 6 french guiding-catheter. Guidewires were placed in the PDA and in the

posterolateral artery (PLA), in order to protect the side branch in case of an eventual occlusion. Predilation 1:1 with a non-compliant (NC) balloon 3.0 × 18 mm (16 atm) was performed until the balloon was completely expanded in angiography. A second OCT run verified fragmentation of restenotic tissue and sufficient luminal gain to ensure adequate scaffold expansion. A poly-lactide BRS (ABSORB 3 × 28 mm) was then slowly deployed at 12 atm, holding pressure for 60 s. Proximal-optimization-technique with an NC-balloon 3.25 × 15 mm (16 atm) was then performed by placing the proximal edge of the distal marker of the balloon at the carina of the PDA-PLA bifurcation, with an optimal angiographic result (Fig. 1D). A final OCT pullback showed optimal apposition and expansion (MLA 5.3 mm²/MLD 2.6 mm; Fig. 1E), structural integrity of the device and clear access to the PLA side branch through the scaffold struts (Fig. 1F). Three-month follow-up documented an optimal clinical and angiographical result (**Suppl. Video 1**).

Poly-lactide BRS are supposed to resorb completely [1–5], depending on the specific device and on patient/local conditions. The resorption restores vasomotion and eventually normal endothelial

Address for correspondence: Milosz J. Jaguszewski, MD, PhD, FESC, 1st Department of Cardiology, Medical University of Gdansk, ul. Dębinki 7, 80–210 Gdańsk, Poland, tel: +48 58 349 25 00, fax: +48 58 346 12 01, e-mail: mjaguszewski@escardio.com.pl

Received: 24.07.2018

Accepted: 15.02.2019

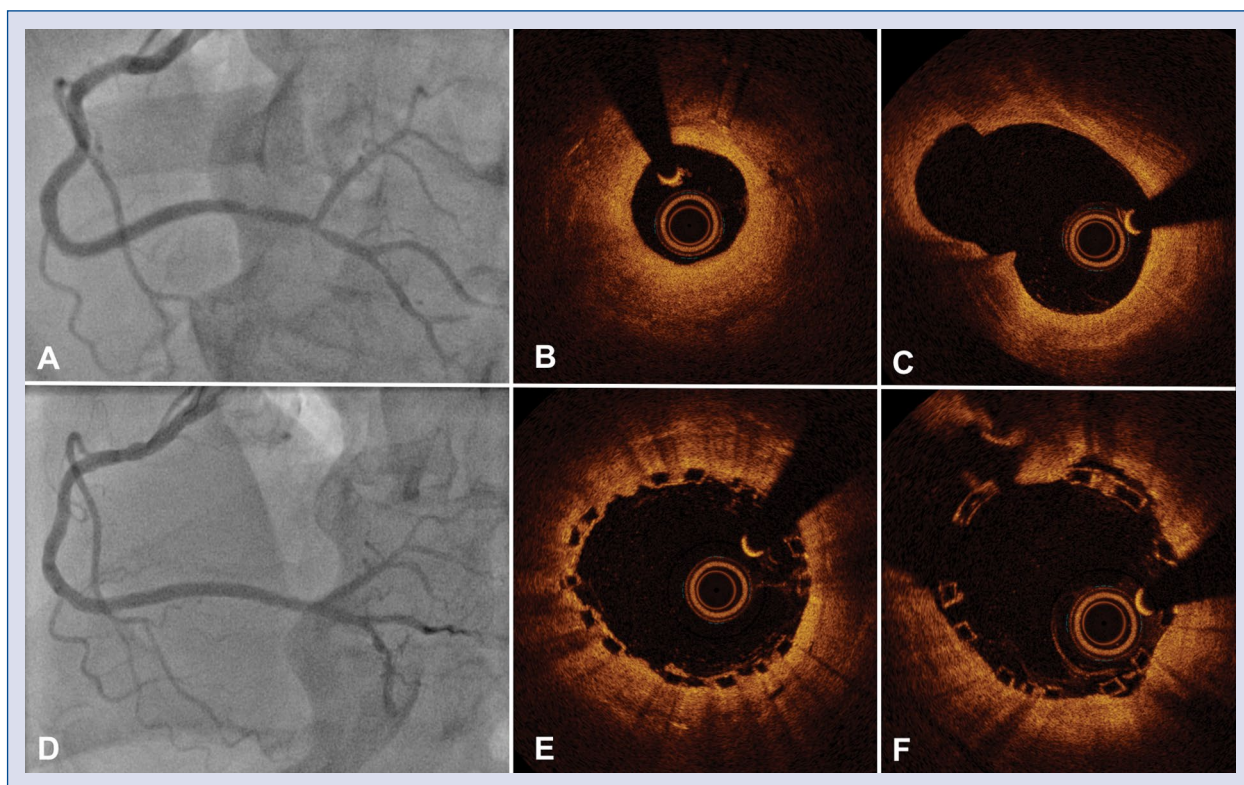


Figure 1. **A, D.** The coronary angiography shows a distal-edge in-stent restenosis in the distal right coronary artery, extending to the PDA, Medina 110 bifurcation; **B, C.** Optical coherence tomography (OCT) shows predominantly fibrolipidic restenotic tissue; **D.** An optimal angiographic result after proximal-optimization-technique with a non-compliant-balloon 3.25×15 mm (16 atm) performed by placing the proximal edge of the distal marker of the balloon at the carina of the PDA-PLA bifurcation; **E, F.** Optimal apposition, expansion and structural integrity of the device and clear access to the PLA side branch through the scaffold struts as assessed by OCT; PDA — posterior descending artery; PLA — posterolateral artery.

function [2, 6, 7]. Moreover, the disappearance of a permanent foreign body in the vessel wall is also intended to minimize inflammation and risk of device failure, i.e. very late BRS-thrombosis, neoatherosclerosis, restenosis and catch-up phenomenon. Nonetheless, the suitability of polylactide BRS for bifurcations is currently a matter of debate, with reported higher risks of side branch occlusion [8] and of scaffold rupture following some bifurcation techniques [9, 10]. Some scientific reports however, focus on dedicating interventional techniques to minimize these risks [10, 11]. ISR is also a challenging scenario for BRS, because the expansion of the scaffold is sensibly inferior than in on-label indications [12] and reported clinical outcomes are inconsistent to date [13, 14]. The current case reports the successful treatment of a lesion combining both bifurcation and ISR challenges, by implanting a BRS. OCT-guidance played an instrumental role in achieving an optimal result

and it may be considered for all off-label indications of BRS devices.

Conflict of interest: None declared

References

1. Oberhauser JP, Hossainy S, Rapoza RJ. Design principles and performance of bioresorbable polymeric vascular scaffolds. *Euro-Intervention*. 2009; 5 Suppl F: F15–F22, doi: [10.4244/EIJV5IFA3](https://doi.org/10.4244/EIJV5IFA3), indexed in Pubmed: [22100671](https://pubmed.ncbi.nlm.nih.gov/22100671/).
2. Serruys PW, Ormiston JA, Onuma Y, et al. A bioabsorbable everolimus-eluting coronary stent system (ABSORB): 2-year outcomes and results from multiple imaging methods. *Lancet*. 2009; 373(9667): 897–910, doi: [10.1016/S0140-6736\(09\)60325-1](https://doi.org/10.1016/S0140-6736(09)60325-1), indexed in Pubmed: [19286089](https://pubmed.ncbi.nlm.nih.gov/19286089/).
3. Onuma Y, Serruys PW, Perkins LEL, et al. Intracoronary optical coherence tomography and histology at 1 month and 2, 3, and 4 years after implantation of everolimus-eluting bioresorbable vascular scaffolds in a porcine coronary artery model: an attempt to decipher the human optical coherence tomography images

- in the ABSORB trial. *Circulation*. 2010; 122(22): 2288–2300, doi: [10.1161/CIRCULATIONAHA.109.921528](https://doi.org/10.1161/CIRCULATIONAHA.109.921528), indexed in Pubmed: 20975003.
4. Verheye S, Ormiston JA, Stewart J, et al. A next-generation bioresorbable coronary scaffold system: from bench to first clinical evaluation: 6- and 12-month clinical and multimodality imaging results. *JACC Cardiovasc Interv*. 2014; 7(1): 89–99, doi: [10.1016/j.jcin.2013.07.007](https://doi.org/10.1016/j.jcin.2013.07.007), indexed in Pubmed: 24139932.
 5. Campos CM, Muramatsu T, Iqbal J, et al. Bioresorbable drug-eluting magnesium-alloy scaffold for treatment of coronary artery disease. *Int J Mol Sci*. 2013; 14(12): 24492–24500, doi: [10.3390/ijms141224492](https://doi.org/10.3390/ijms141224492), indexed in Pubmed: 24351829.
 6. Serruys PW, Onuma Y, Dudek D, et al. Evaluation of the second generation of a bioresorbable everolimus-eluting vascular scaffold for the treatment of de novo coronary artery stenosis: 12-month clinical and imaging outcomes. *J Am Coll Cardiol*. 2011; 58(15): 1578–1588, doi: [10.1016/j.jacc.2011.05.050](https://doi.org/10.1016/j.jacc.2011.05.050), indexed in Pubmed: 21958884.
 7. Serruys PW, Chevalier B, Dudek D, et al. A bioresorbable everolimus-eluting scaffold versus a metallic everolimus-eluting stent for ischaemic heart disease caused by de-novo native coronary artery lesions (ABSORB II): an interim 1-year analysis of clinical and procedural secondary outcomes from a randomised controlled trial. *Lancet*. 2015; 385(9962): 43–54, doi: [10.1016/S0140-6736\(14\)61455-0](https://doi.org/10.1016/S0140-6736(14)61455-0), indexed in Pubmed: 25230593.
 8. Muramatsu T, Onuma Y, García-García HM, et al. Incidence and short-term clinical outcomes of small side branch occlusion after implantation of an everolimus-eluting bioresorbable vascular scaffold: an interim report of 435 patients in the ABSORB-EXTEND single-arm trial in comparison with an everolimus-eluting metallic stent in the SPIRIT first and II trials. *JACC Cardiovasc Interv*. 2013; 6(3): 247–257, doi: [10.1016/j.jcin.2012.10.013](https://doi.org/10.1016/j.jcin.2012.10.013), indexed in Pubmed: 23517836.
 9. Ormiston JA, Webber B, Ubod B, et al. Absorb everolimus-eluting bioresorbable scaffolds in coronary bifurcations: a bench study of deployment, side branch dilatation and post-dilatation strategies. *EuroIntervention*. 2015; 10(10): 1169–1177, doi: [10.4244/EIJY14M05_08](https://doi.org/10.4244/EIJY14M05_08), indexed in Pubmed: 24835848.
 10. Ormiston JA, Webber B, Ubod B, et al. An independent bench comparison of two bioresorbable drug-eluting coronary scaffolds (Absorb and DESolve) with a durable metallic drug-eluting stent (ML8/Xpedition). *EuroIntervention*. 2015; 11(1): 60–67, doi: [10.4244/EIJY15M02_03](https://doi.org/10.4244/EIJY15M02_03), indexed in Pubmed: 25680225.
 11. Derimay F, Souteyrand G, Motreff P, et al. Sequential proximal optimizing technique in provisional bifurcation stenting with everolimus-eluting bioresorbable vascular scaffold: fractal coronary bifurcation bench for comparative test between absorb and XIENCE xpedition. *JACC Cardiovasc Interv*. 2016; 9(13): 1397–1406, doi: [10.1016/j.jcin.2016.04.021](https://doi.org/10.1016/j.jcin.2016.04.021), indexed in Pubmed: 27388830.
 12. Rivero F, Bastante T, Cuesta J, et al. Treatment of in-stent restenosis with bioresorbable vascular scaffolds: optical coherence tomography insights. *Can J Cardiol*. 2015; 31(3): 255–259, doi: [10.1016/j.cjca.2014.11.017](https://doi.org/10.1016/j.cjca.2014.11.017), indexed in Pubmed: 25660152.
 13. Moscarella E, Ielasi A, Granata F, et al. Long-Term Clinical Outcomes After Bioresorbable Vascular Scaffold Implantation for the Treatment of Coronary In-Stent Restenosis: A Multicenter Italian Experience. *Circ Cardiovasc Interv*. 2016; 9(4): e003148, doi: [10.1161/CIRCINTERVENTIONS.115.003148](https://doi.org/10.1161/CIRCINTERVENTIONS.115.003148), indexed in Pubmed: 27059683.
 14. Jamshidi P, Nyffenegger T, Sabti Z, et al. A novel approach to treat in-stent restenosis: 6- and 12-month results using the everolimus-eluting bioresorbable vascular scaffold. *EuroIntervention*. 2016; 11(13): 1479–1486, doi: [10.4244/EIJV11I13A287](https://doi.org/10.4244/EIJV11I13A287), indexed in Pubmed: 27107313.