

IMAGES IN INTERVENTION

Very Late Restenosis After Bioresorbable Scaffold Implantation Due to Simultaneous External Compression of the Scaffold and Intrasc scaffold Tissue Growth



Akihito Tanaka, MD,*† Neil Ruparelia, MBBS, DPHIL,*†† Hiroyoshi Kawamoto, MD,*† Azeem Latib, MD,*† Antonio Colombo, MD*†

A 78-year-old man underwent percutaneous coronary intervention with implantation of a 2.5 × 28-mm Absorb bioresorbable scaffold (BRS) (Abbott Vascular, Santa Clara, California) in the mid-left anterior descending coronary artery, and treatment with a drug-coated balloon for recurrent metallic stent restenosis in the proximal segment (Figures 1A and 1B) (1).

Follow-up angiography at 7 months demonstrated no restenosis at the BRS site (Figure 1C). After 34 months, the patient returned with a recurrence of angina and underwent angiography, which demonstrated restenosis at the BRS site as well as recurrent metallic stent restenosis (Figure 1D). Intravascular ultrasound was performed at baseline, showing some residual scaffold struts. Although the vessel area was similar to that at the index procedure, the minimal lumen area was 1.73 mm² with a lumen loss of 3.68 mm², likely caused by simultaneous external compression (1.97 mm²) and intrasc scaffold tissue growth (1.71 mm²) (Figure 2).

By virtue of its bioresorption, the Absorb BRS begins to lose its radial strength approximately 6 months after implantation, and completely loses it by 12 months (2,3). Taking these processes into account,

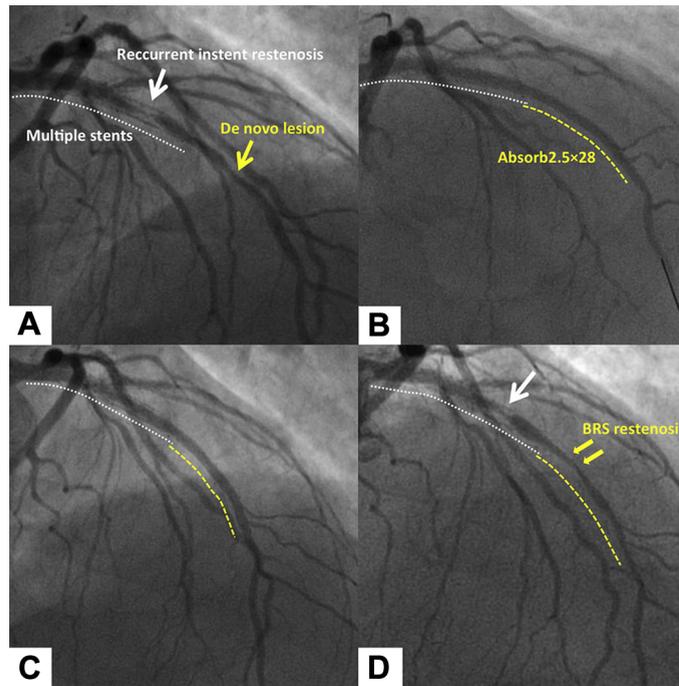
the mechanisms of BRS restenosis may be different to those in the setting of metallic stents, especially restenosis occurring in the very late phase.

On the basis of reports describing BRS restenosis (3,4), in the short-term, scaffold restenosis seems to be a result of intrasc scaffold tissue growth similar to that seen with metallic stents, and would be reasonable because the scaffold still maintains some degree of its radial strength during the period. On the other hand, little data are available regarding scaffold restenosis occurring in the very late phase. Although a case series suggested very late restenosis could be also attributed to intrasc scaffold tissue growth (3), data remain sparse.

In our case, very late restenosis appeared to be a result of the simultaneous external compression of the scaffold and intrasc scaffold tissue growth. This may suggest another potential mechanism of very late scaffold restenosis with progression of plaque behind incompletely resorbed scaffold struts.

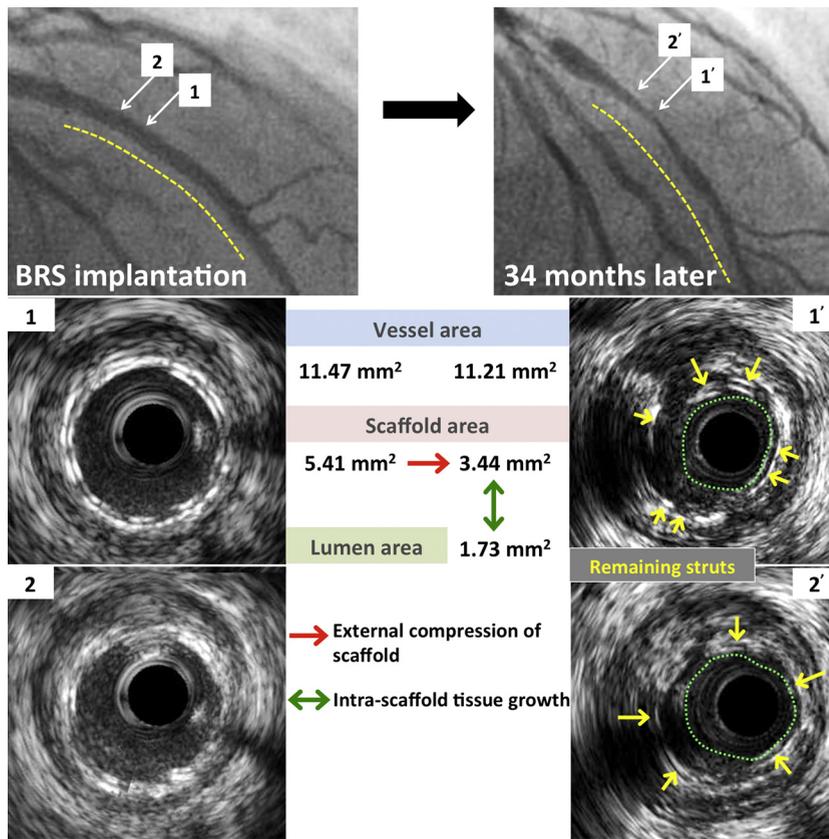
REPRINT REQUESTS AND CORRESPONDENCE: Dr. Antonio Colombo, EMO-GVM Centro Cuore Columbus, 48 Via M. Buonarroti, 20145 Milan, Italy. E-mail: info@emocolumbus.it.

From the *Interventional Cardiology Unit, EMO-GVM Centro Cuore Columbus, Milan, Italy; †Interventional Cardiology Unit, San Raffaele Scientific Institute, Milan, Italy; and the †Department of Cardiology, Imperial College, London, United Kingdom. Dr. Latib serves on the advisory board for Medtronic; and receives honoraria from Boston Scientific and Abbott Vascular. All other authors have reported that they have no relationships relevant to the contents of this paper to disclose.

FIGURE 1 Angiography of the Index Procedure and Follow-Up

(A) Baseline angiogram demonstrating a de novo lesion (**yellow arrow**) in the mid-left anterior descending coronary artery (LAD) and recurrent restenosis (**white arrow**) within multiple metallic stents in the proximal LAD. **(B)** Final angiogram showing a good angiographic result following implantation of 1 bioresorbable scaffold (BRS) into the mid-LAD as well as drug-coated balloon treatment in the proximal LAD. **(C)** Follow-up angiogram after 7 months showing no restenosis at the BRS segment. **(D)** Follow-up angiogram after 34 months showing BRS restenosis (**yellow arrows**) as well as recurrent metallic stent restenosis (**white arrow**). The **white dotted line** indicates multiple stents and the **yellow dashed line** indicates the BRS.

FIGURE 2 IVUS Imaging at the Index BRS Implantation and at 34 Months Follow-Up



At the site of minimal lumen area within the bioresorbable scaffold (BRS), the vessel area was similar (11.47 mm² at BRS implantation and 11.21 mm² at 34 months follow-up). IVUS imaging showed some remaining struts (yellow arrow) with the scaffold area decreasing at 34 months (5.41 mm² at implantation and 3.44 mm² at 34 months). Lumen area at 34 month was 1.73 mm² (the lumen loss during 34 months was 3.68 mm²). This lumen loss (3.68 mm²) was likely caused by the simultaneous external compression of the scaffold (5.41 mm² – 3.44 mm² = 1.97 mm²) and intrascaffold tissue growth (3.44 mm² – 1.73 mm² = 1.71 mm²). The yellow dashed line indicates the BRS. IVUS = intravascular ultrasound. The green dotted line indicates lumen area.

REFERENCES

- Naganuma T, Latib A, Ielasi A, et al. No more metallic cages: an attractive hybrid strategy with bioresorbable vascular scaffold and drug-eluting balloon for diffuse or tandem lesions in the same vessel. *Int J Cardiol* 2014;172:618-9.
- Onuma Y, Serruys PW, Muramatsu T, et al. Incidence and imaging outcomes of acute scaffold disruption and late structural discontinuity after implantation of the absorb everolimus-eluting fully bioresorbable vascular scaffold: optical coherence tomography assessment in the ABSORB cohort B trial (A Clinical Evaluation of the Bioabsorbable Everolimus Eluting Coronary Stent System in the Treatment of Patients With De Novo Native Coronary Artery Lesions). *J Am Coll Cardiol Intv* 2014;7:1400-11.
- Nakatani S, Onuma Y, Ishibashi Y, et al. Early (before 6 months), late (6-12 months) and very late (after 12 months) angiographic scaffold restenosis in the ABSORB Cohort B trial. *Euro-Intervention* 2015;10:1288-98.
- Longo G, Granata F, Capodanno D, et al. Anatomical features and management of bioresorbable vascular scaffolds failure: a case series from the GHOST registry. *Catheter Cardiovasc Interv* 2015;85:1150-61.

KEY WORDS bioresorbable scaffold, percutaneous coronary intervention, restenosis