

# MASTER THE COMPLEX

Optimizing revascularization through innovation, training, and education.





A TALE OF
4 BIFURCATIONS

## Study and Images Courtesy of Dr. Colum Owens Royal Victoria Hospital Belfast

Clinical Interests include: Percutaneous Coronary Intervention, Adult Congenital Heart Disease and Structural Heart Disease Intervention

#### Patient History

- 57-year-old male. Ex-smoker. 1st presentation of ischemic heart disease
- Presented with a one week history of progressive angina and an anterolateral ST-elevation myocardial infarct complicated by cardiogenic shock

#### Clinical Presentation

- Unwell on arrival with blood pressure of 78/42 mmHg
- Echo: severe LV dysfunction with LVEF of 30%
- · Right radial access, 7Fr sheath
- Diagnostic angiography showed severe 3-vessel disease with complex bifurcations involving the distal right coronary artery
  (RCA) (click on video 1), left main stem (LM/LMS), left circumflex bifurcation (LCX) (click on video 2) and an occluded
  left anterior descending/2nd diagonal artery bifurcation (LAD/D2) (click on video 3)
- · After surgical discussion, decision taken to re-vascularise with percutaneous revascularisation

#### Techniques Used

- Primary PCI
- . Multiple complex bifurcation techniques including modified crush, mini crush and culotte with use of anchoring balloons

### SYNERGY Platform characteristics

- The two connector design on the stent body, vessel conformability<sup>4</sup> and large side-branch access<sup>3</sup> allowed for multiple bifurcation techniques
- Ability of stent overexpansion, when called for during post dilatation, to match the diameter of larger vessels, and permits complex LM intervention. (The SYNERGY 4.0 mm dedicated stent has a labeled overexpansion capability of up to 5.75 mm<sup>5</sup>)
- Its deliverability, visibility and conformability<sup>4</sup> allowed for full revascularisation of complex disease

#### SYNERGY Drug & polymer characteristics

- Abluminal coating and synergistic drug release<sup>1</sup> with polymer degradation\* facilitates early healing<sup>6</sup> from day one and thus allows for flexibility of DAPT duration<sup>7</sup>. Polymer is gone after completion of drug elution at 3 months. This is an important consideration when baseline morbidity is unknown as in this case where the patient presents with cardiogenic shock
- \* The polymer is gone when it's no longer needed, shortly after the drug is completely eluted at three months, which minimizes polymer exposure in the vessel. 1.2



Distal RCA Bifurcation Stenosis



Left Main, Left Circumflex & Left Anterior Descending Disease



Occluded Left Anterior Descending/2nd Diagonal Artery Bifurcation

#### **KEY LEARNINGS**

This case highlights:

- Interventional strategies to deal with complex bifurcation disease in a patient presenting with cardiogenic shock
- · How a robust knowledge of stent characteristics allow for full revascularisation in patients with complex disease
- The importance of stent choice where DAPT duration considerations are important
   To view complete, narrated case video, click here: A Tale of 4 Bifurcations





Final Result in LAD/D2



Residual Left Main, Proximal LAD and Circumflex Disease



LMS, LAD and OM1 Bifurcation Stent Result



PLB/PDA Bifurcation Result



Balloon Anchoring & OM1/OM2 Culotte



Final Result in Left Main and Left Circumflex

#### Acute Procedure (Primary PCI to LAD/D2 bifurcation)

- Flow re-established in LAD/D2 with a 2.5 mm compliant balloon but D2 dissected
- A "crush" bifurcation strategy chosen for 2 reasons: vessel size disparity and maintaining flow in dissected diagonal by stenting it first with a 2.25 x 32 mm SYNERGY DES placed from mid LAD to D2 and optimised with a 2.5 mm NC EMERGE halloon
- Crushed with 3.0 x 15 mm compliant balloon in LAD (modified crush technique using a balloon rather than a stent)
- LAD then stented with 3.0 x 32 mm SYNERGY DES optimising with a 3.5 NC EMERGE balloon
- Final kissing balloon inflation in LAD/D2 with 3.5 and 2.5 mm NC EMERGE balloon respectively with TIMI III flow restored (click on video 4)

### Acute Procedure (Left main/circumflex bifurcation)

- Complex residual disease involving LM, proximal LAD and LCX (click on video 5)
- Second marginal circumflex (OM2) directly stented with a 2.25 x 24 mm SYNERGY DES optimising with a 2.5 mm NC EMERGE balloon
- 1st marginal circumflex (OM1) stented directly with a 2.5 x 24 mm SYNERGY DES back into left main/LCX os in a "mini crush" strategy, optimised with a 3.0 mm NC balloon and crushed with a 3.5 mm balloon in the LMS
- LMS to LAD stented with a 4.0 x 28 mm SYNERGY DES and optimised with a 4.0 mm NC balloon. The LMS was proximally
  optimised with a 5.0 mm NC balloon ensuring both stent apposition and side branch access to the LCX, and demonstrating
  stent over-expansion characteristics
- Final kissing balloon inflation in LMS/OM1 bifurcation with 4.0 and 3.0 mm NC balloons resulting in resolved cardiogenic shock and improved ST segments (click on video 6)

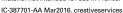
#### Elective Procedure (Distal RCA bifurcation-Culotte)

- Residual disease addressed in order to complete revascularisation
- Distal RCA bifurcation treated using culotte technique as branches comparable in size (click on video 1)
- PLB stented with a 2.5 x 28 mm SYNERGY DES, optimising with a 3.0 and 3.5 mm NC EMERGE balloon
- Distal RCA stented into the PDA with a 2.5 x 24 mm SYNERGY DES and optimised with a 3.0 mm NC EMERGE balloon
- Distal RCA to PL bifurcation culotted with a 3.0 x 38 mm and a 4.0 x 20 mm SYNERGY DES in an overlapping fashion, optimising with 3.5 and 4.0 mm NC EMERGE balloons
- Final kissing balloon inflation into PL/PDA bifurcation with 3.5 and 3.0 NC EMERGE balloons respectively (click on video 7)

### OM1/OM2 bifurcation (Culotte)

- Check angiogram showed patent stents but significant residual OM2/OM1 bifurcation disease
- To address this complex residual bifurcation disease, a 7Fr guide and anchoring balloon technique was used in OM1 to facilitate stent delivery in OM2
- Using a 2.5 mm EMERGE in the OM1for anchoring a 2.5 x 24 mm SYNERGY DES was delivered into the OM2 and optimised with a 3.0 mm NC EMERGE balloon to high pressure in a culotte technique (click on video 8)
- A final kissing balloon inflation undertaken in the OM1/OM2 bifurcation with a 2.5 and 3.0 mm NC EMERGE balloon respectively
- The LMS/OM2 bifurcation finally optimised with a kissing balloon inflation 5.0 and 3.0 mm NC EMERGE balloons
  respectively with excellent final result (click on video 9)

CAUTION: The law restricts these devices to sale by or on the order of a physician. Indications, contraindications, warnings and instructions for use can be found in the product labelling supplied with each device. Information for the use only in countries with applicable health authority product registrations. Material not intended for use in France.





<sup>&</sup>lt;sup>1</sup> Eppihimer M, PhD. Impact of Polymer Type and Location on Stent Thrombogenicity and Endothelial Cell Coverage. EuroPCR 2014

<sup>&</sup>lt;sup>2</sup> Chen YL, PhD, Foss A, PhD, Eppihimer M, PhD, et al. Characterization of In Vivo Poly(DL-lactic-co-glycolic acid) Bioabsorption from a Drug-Eluting Stent. EuroPCR 2012

<sup>&</sup>lt;sup>3</sup> Stent\_MECD\_Technical \_Report; PDM doc.91019938

<sup>&</sup>lt;sup>4</sup> Data on file at Boston Scientific. Bench testing performed by Boston Scientific Corporation. Bench test results not necessarily indicative of clinical performance

performance
5 SYNERGY Stent Systems DELI

SORT OUT VIII presentation by Ida Riise Balleby, MD at PCR 2015; 2 month OCT Analysis presentation by J. M. de la Torre, MD at TCT 2014; TIMELESS presentation by J. un Granada, MD, at CRT 2015; Burgos-Santander study presentation by J. M. de la Torre, MD at PCR 2015

<sup>&</sup>lt;sup>7</sup> SYNERGY is designed with a low initial polymer load, abluminal coating and bioabsorbable polymer which may reduce the risk of thrombosis and the need for prolonged dual antiplatelet therapy. In selected higher risk patients where the physician determines that the risks outweigh the benefits of continued DAPT, it may be reasonable to interrupt or discontinue therapy after 1 month based on low stent thrombosis rates and no observed increased risk for stent thrombosis as shown in the current literature.