

Thin struts, low COF



Tri-axial system with

braided shaft

(4F)

Low profile delivery

system



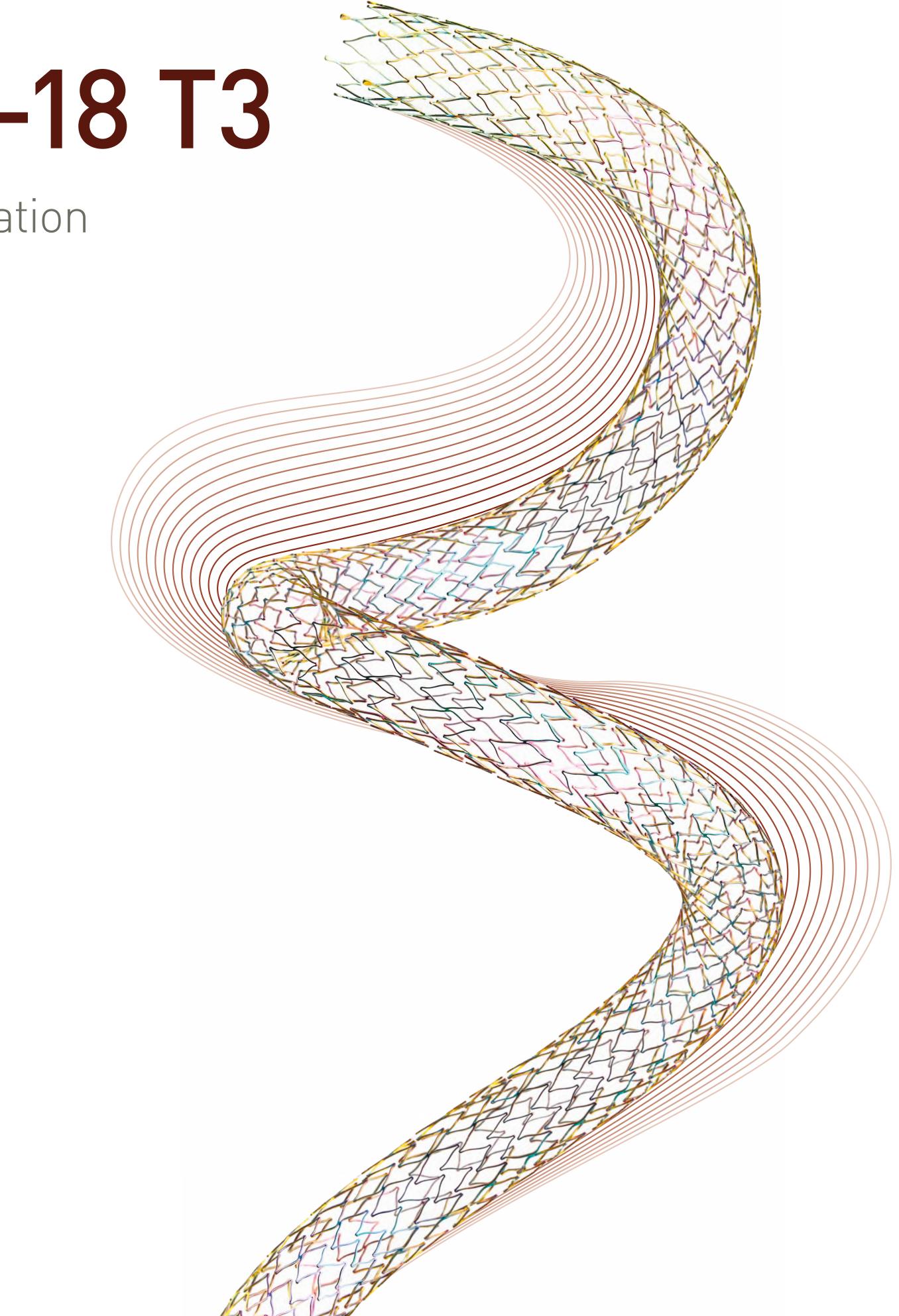
Technical data / ordering info

Vascular Intervention // Peripheral Self-Expanding Stent System/0.018"/OTW



# Pulsar<sup>®</sup>-18 T3

A unique combination of 3 technologies

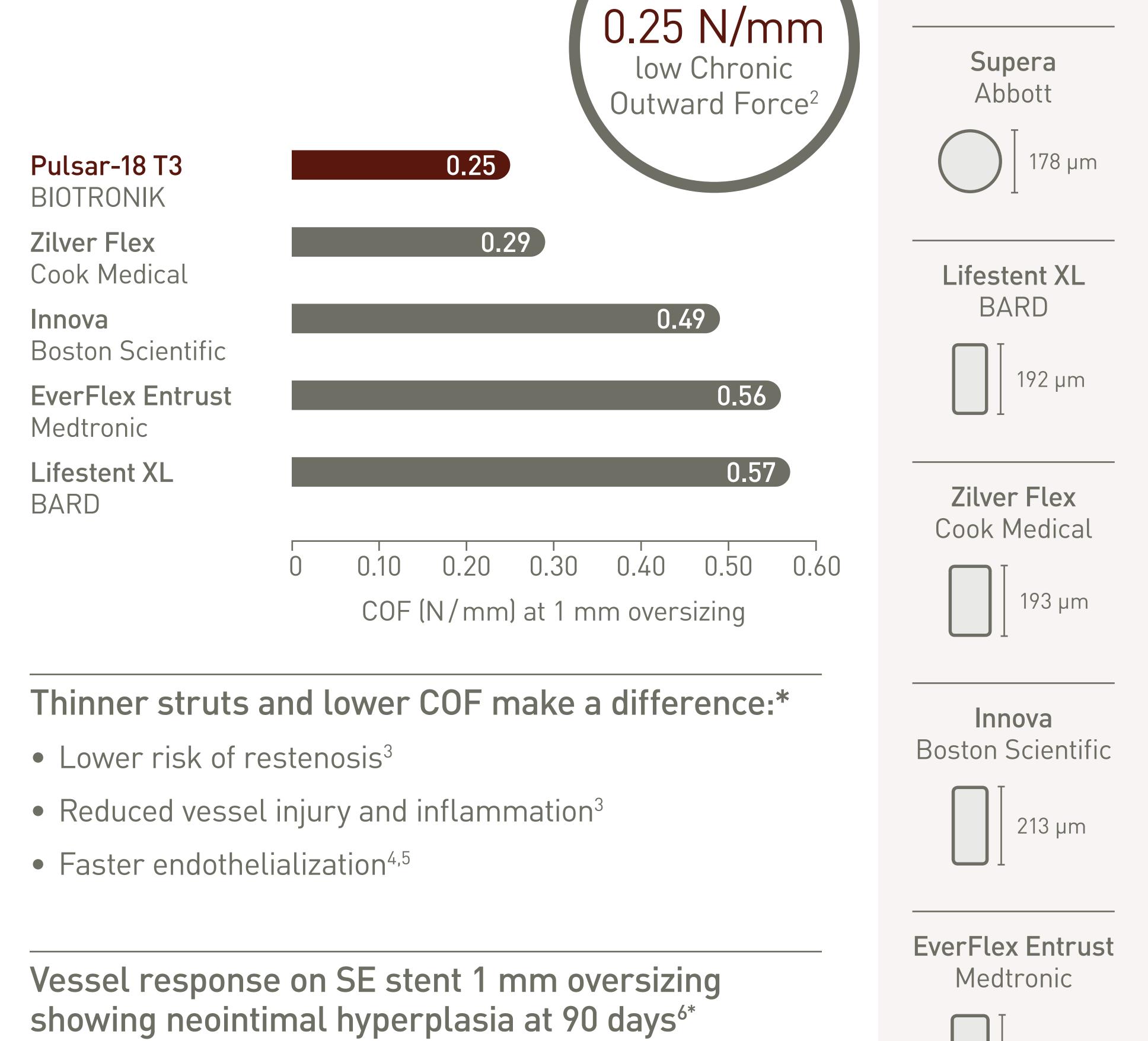


### 140 µm thin struts - thinner than leading brands<sup>1</sup>

#### Thinner struts for lower Chronic Outward Force (COF)<sup>2</sup>

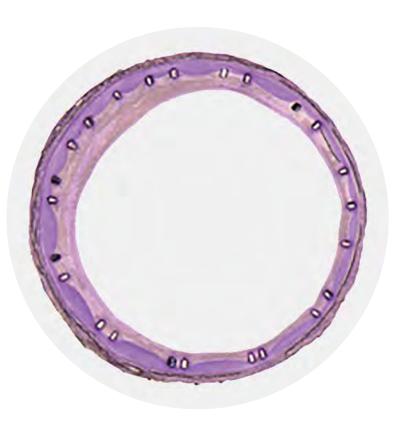
BIOTRONIK

Zilver Flex



Stent strut thickness in perspective<sup>1</sup> Pulsar-18 T3 BIOTRONIK 140 µm

228 µm

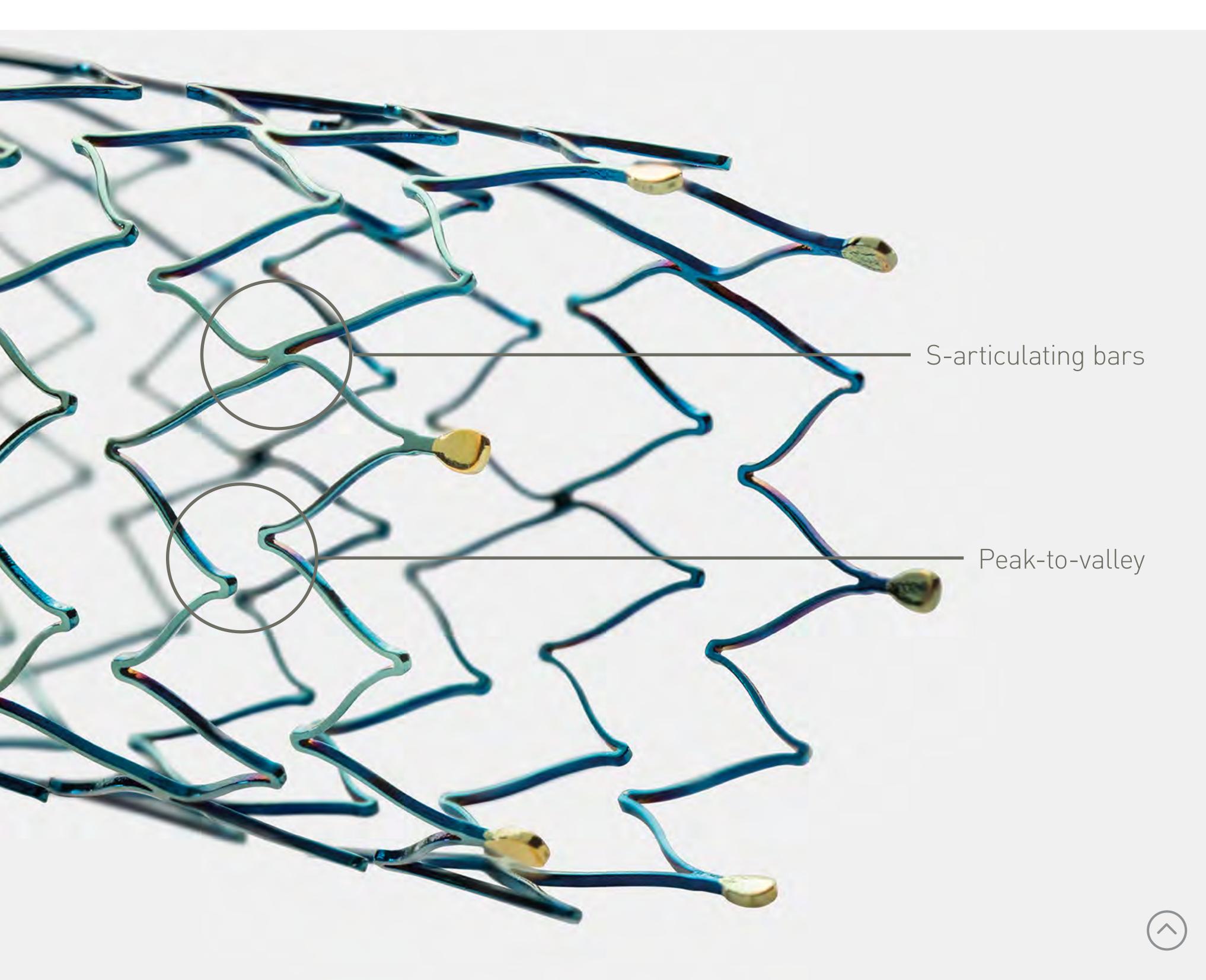




**Pulsar Stent** BIOTRONIK Low COF

Lifestent XL BARD High COF

#### \*As demonstrated in pre-clinical studies





# Unique tri-axial shaft design on 4F low profile

# Tri-axial system with braided retractable shaft

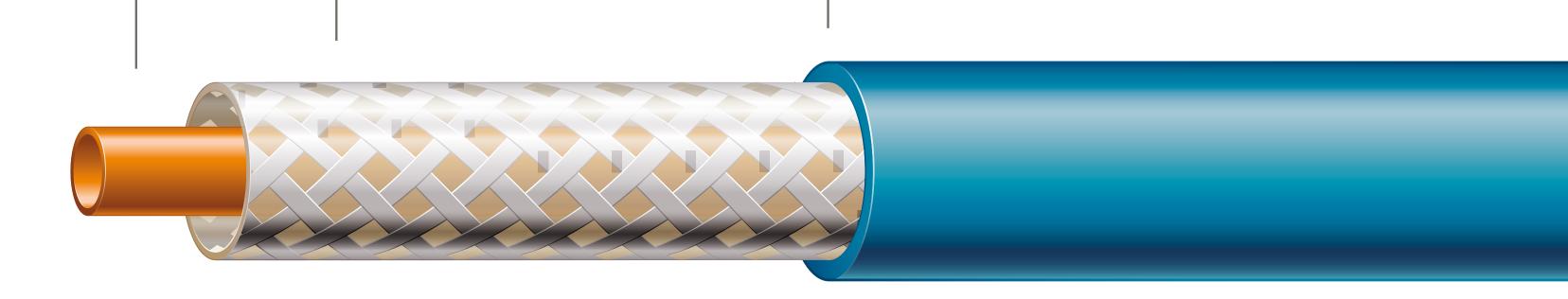
#### Accurate stent deployment

The outer stabilizing shaft isolates the retractable shaft from friction caused by the introducer valve to ensure accurate stent deployment.

Inner shaft

Retractable braided shaft

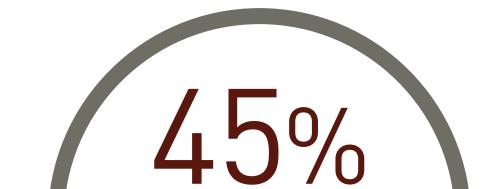
Outer stabilizing shaft



# 4F low profile - improved acute outcomes\* vs. 6F<sup>7</sup>

Potential for safer, faster and simpler procedures than 6F

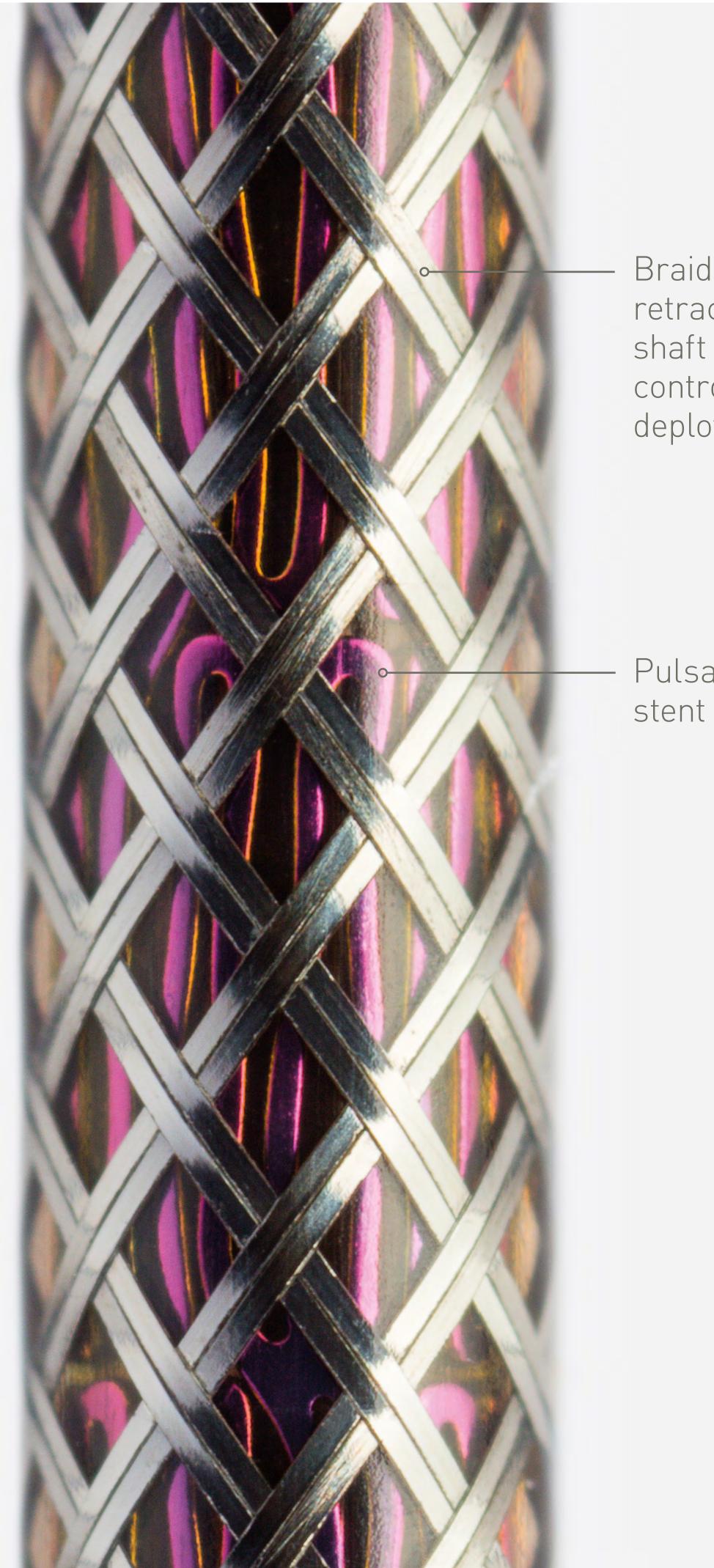
- Clinically proven lower access site complication rates<sup>7</sup>
- Shorter compression time<sup>7</sup>
- 45% smaller puncture site than 6F<sup>8</sup>
- No need for a closure device<sup>7</sup>



• Potential for ambulatory treatment

#### smaller puncture site area than 6F<sup>8</sup>

#### \*Less access site complications



Braided retractable shaft for controlled deployment

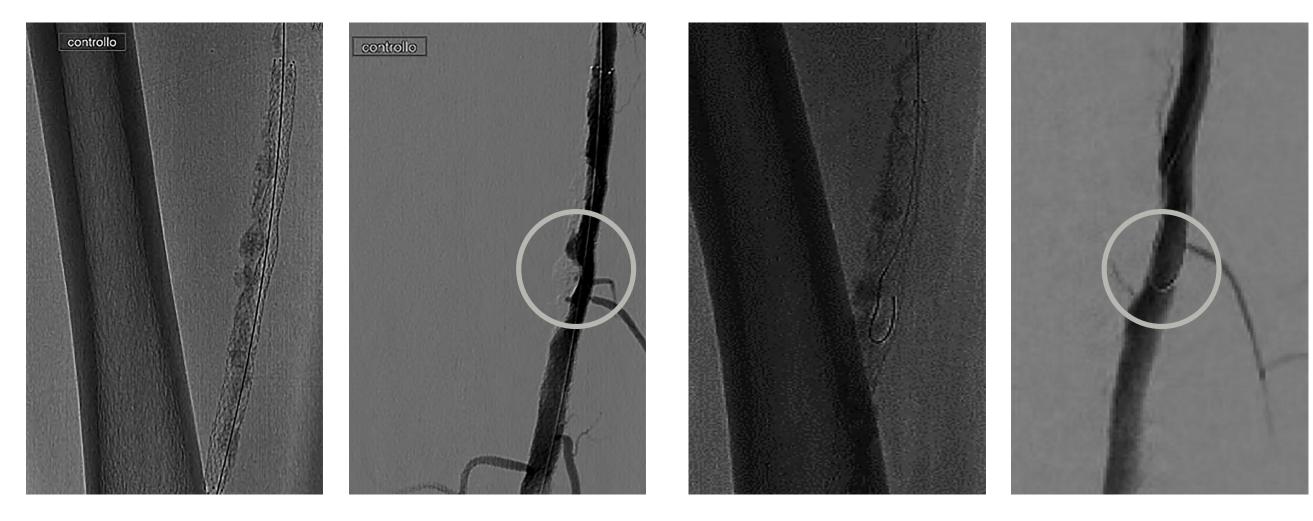
stent

 $\bigcirc$ 

Clinically pro	ven	
Safety and efficacy at	12 months	
<b>4F INTERVENTIONS</b> 4EVER <sup>7</sup>		
FTLR:** <b>89.3%</b>	PP: <b>* 81.4%</b>	A.L.L: <sup>++</sup> 7.1 cm
LONG & OCCLUDED TASC D <sup>9</sup>		
FTLR: <b>86%</b>	PP: <b>77%</b>	A.L.L: <b>24.5 cm</b>
ALL-COMERS BIOFLEX PEACE	E <sup>10</sup> (stent only)	
FTLR: <b>89.3%</b>	PP: <b>84.7%</b>	A.L.L: <b>8.2 cm</b>

\*\* FTLR - Freedom from Target Lesion Revascularization; \*PP - Primary Patency; \*\*A.L.L. - Average Lesion Length

### Sufficient radial force for long term vessel support, even in calcified lesions

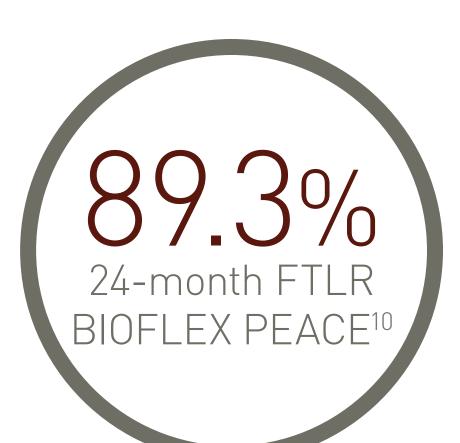


2016

After the treatment 2011 (Courtesy of Prof. van den Berg<sup>8</sup>)

With a constant low chronic outward force applied to the vessel, patency can be achieved and maintained over a long term follow up even in calcified lesions.

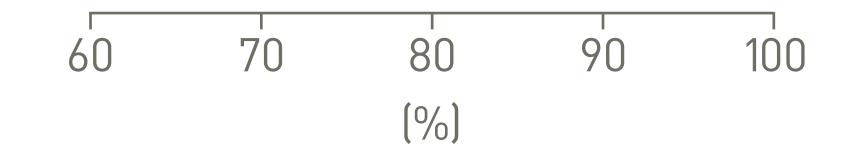
24-month outcomes of Pulsar stent, highlighting the long term safety and efficacy



Stuc	dy, Product	Manufacturer	A.L.L. <sup>++</sup>	PP <sup>+</sup>	
	FLEX PEACE <sup>10</sup> sar (stent only)	BIOTRONIK	8.2 cm	78.4%	FTLR** 89.3%
SUP Sup	PERB <sup>11</sup> era	Abbott	7.8 cm	N/A	FTLR 83.3%
4EV Puls	ER <sup>12</sup> sar	BIOTRONIK	<b>7.1 cm</b>	72.3%	FTLR 82.7%
	OLL <sup>13</sup> A.R.T Control	Cardinal Health/Cordis	7.7 cm	74.9%	FTLR 80.3%

RESILIENT <sup>14</sup> Lifestent	BD/Bard	7.0 cm	N/A	FTLR 77.8%
ZILVER PTX <sup>15</sup> Zilver BMS provisional	Cook Medical	6.3 cm	65.8%	FTLR 76.7%
DURABILITY II <sup>16</sup> EverFlex	Medtronic	10.9 cm	66.1%	FTLR 75.3%

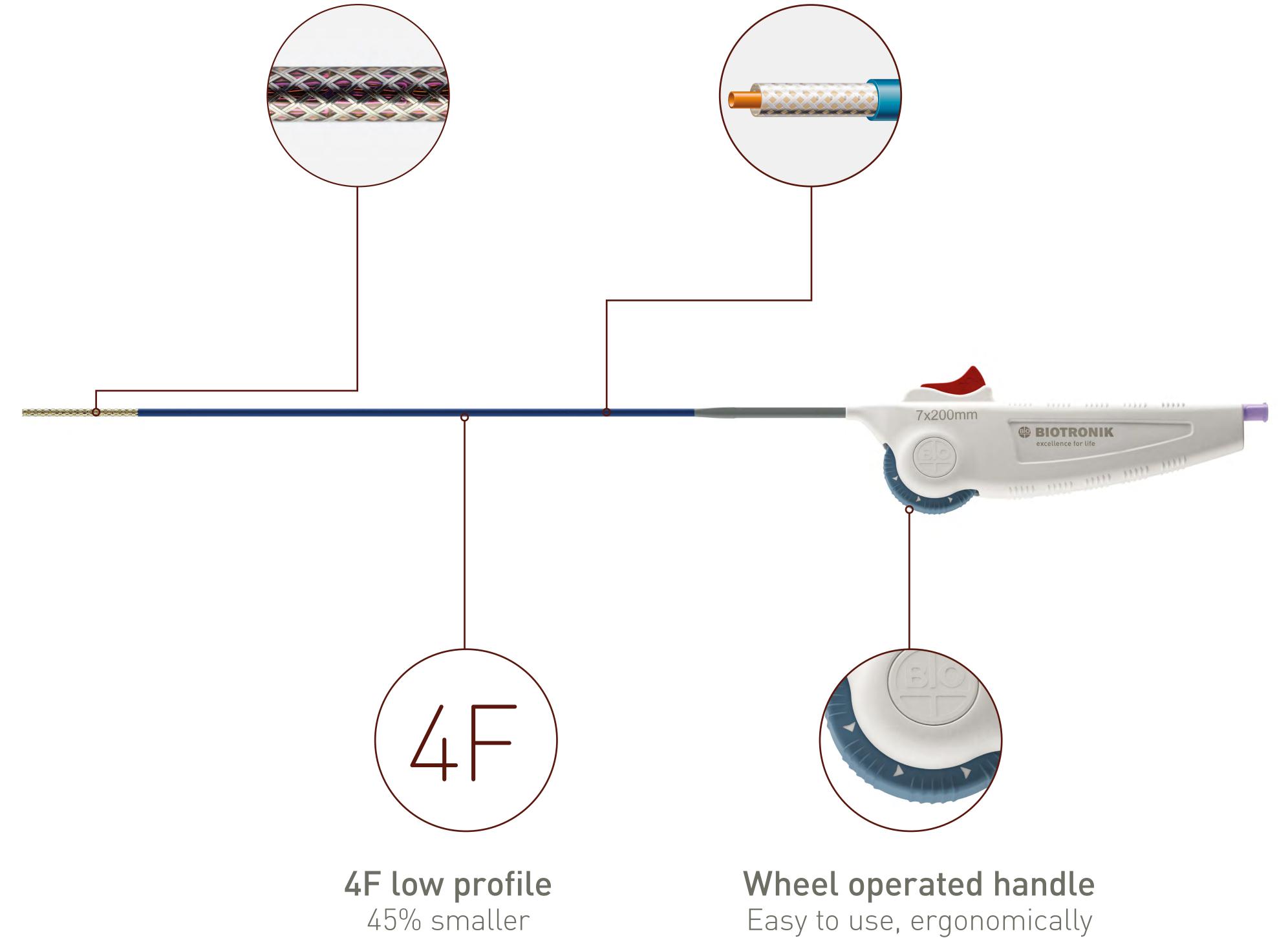
Results from different trials are not directly comparable. Differences in outcomes may be the result of differences in protocol design, patient populations or other factors. Astron Pulsar, Pulsar-18, Pulsar-18 T3 and Pulsar-35 have equivalent stent platforms, therefore the clinical results are valid for the Pulsar range.



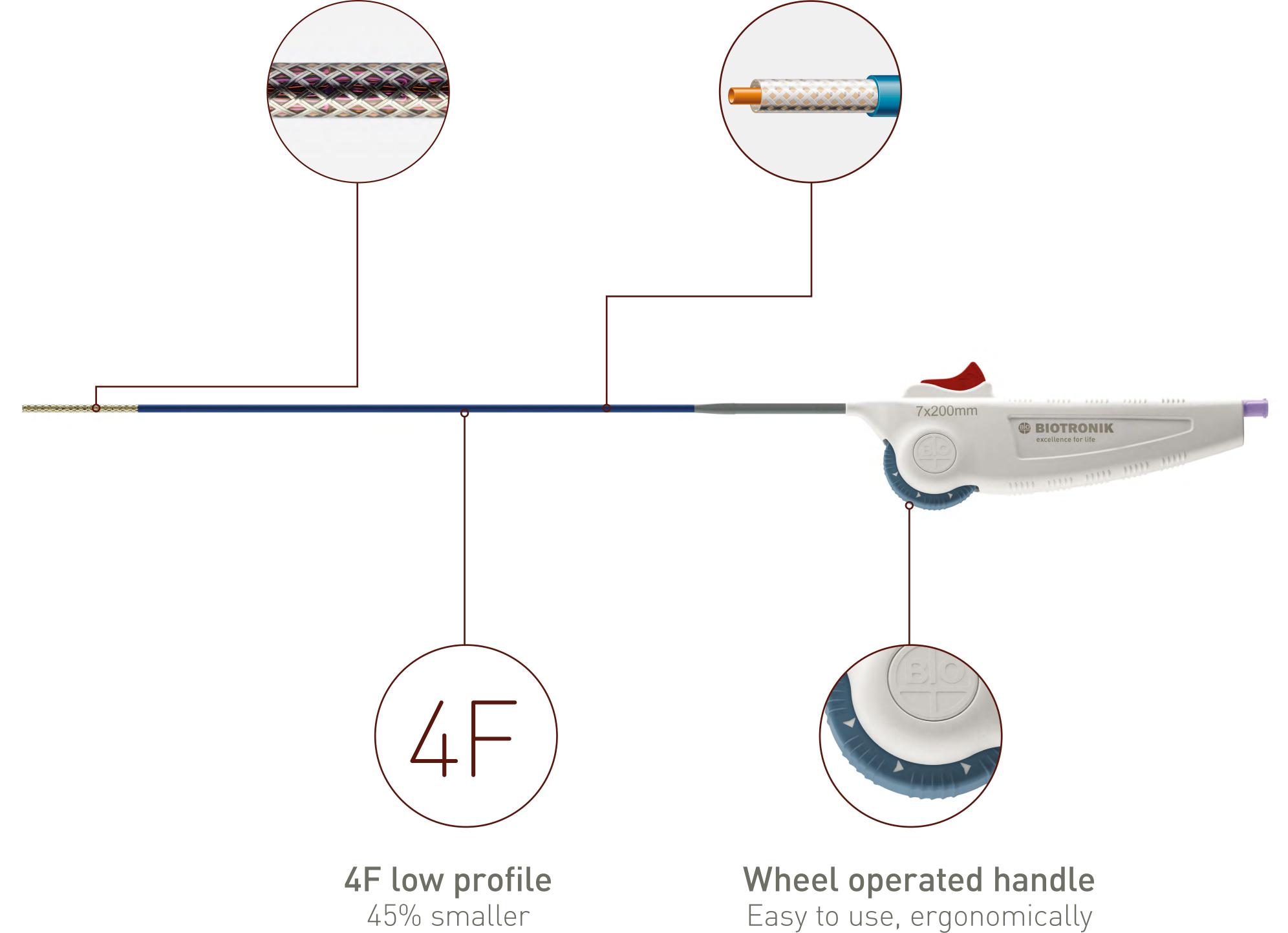
#### Pulsar-18 T3 A unique combination of 3 technologies



#### Braided retractable shaft Controlled deployment



#### Tri-axial shaft Accurate stent deployment



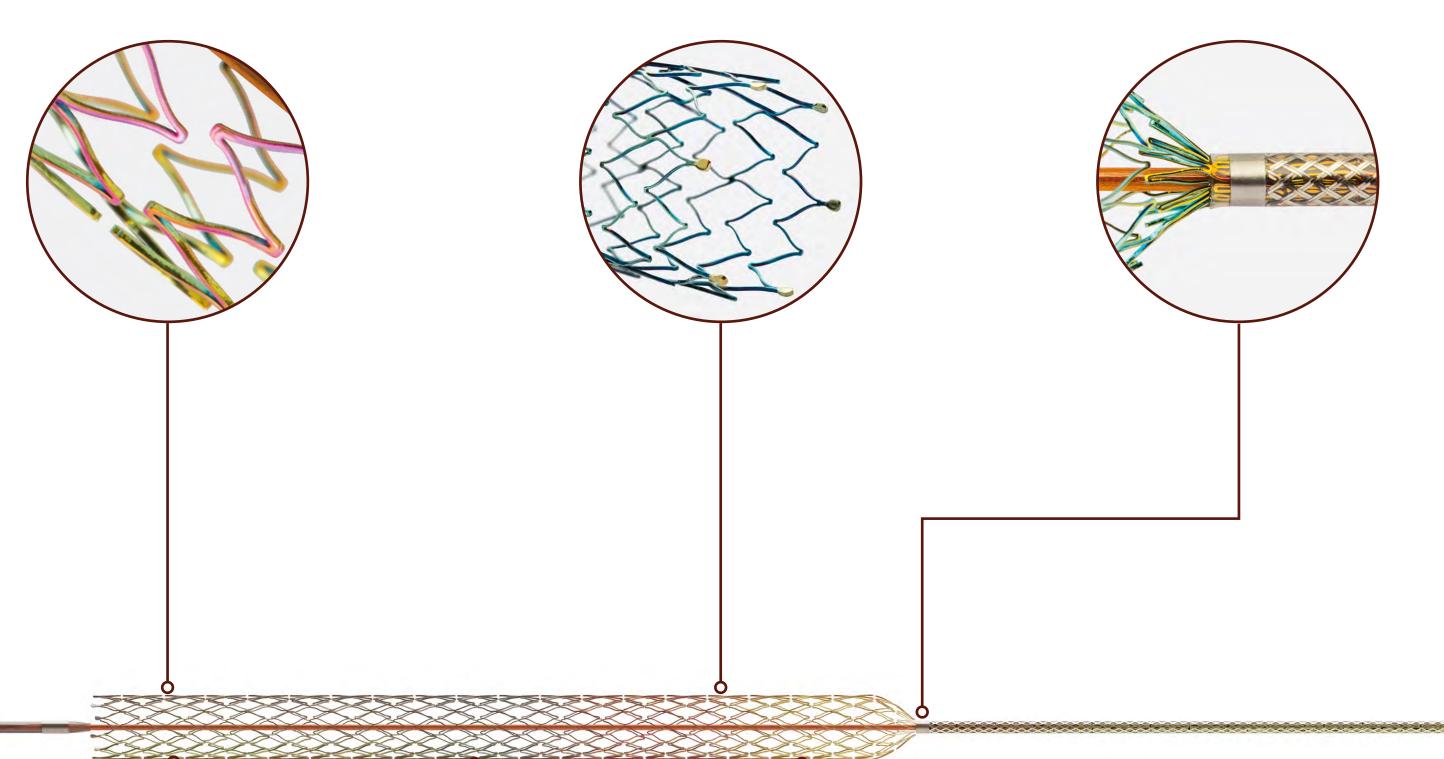
puncture site vs. 6F<sup>8</sup>

designed handle.

#### proBIO<sup>®</sup> coating Reduces ion release

#### 140 µm thin struts Thinner than leading brands

Additional radiopaque marker Improved visibility





#### Designed for the SFA\*

Multi-directional flexibility to conform to vessel movement Low chronic outward force For lower risk of restenosis<sup>3</sup>

\*Superficial Femoral Artery

### Pulsar-18 T3

Indicated for use in patients with atherosclerotic disease of the superficial femoral, proximal popliteal and infrapopliteal arteries and for the treatment of insufficient results after Percutaneous Transluminal Angioplasty (PTA), e.g. residual stenosis and dissection.\* Vascular Intervention Peripheral

Technical Data	Stent	Stent						
	Catheter type	OTW						
	Recommended guide wire	0.018"						
	Stent material	Nitinol						
	Strut thickness	140 µm						
	Strut width	85 µm						
	Stent coating	proBIO® (Amorphous Silicon Carbide)						
	Stent Markers	6 gold markers each end						
	Sizes	ø 4.0 - 7.0 mm: L:20 - 200 mm						
	Shaft	4F, hydrophobic coating, tri-axial						
	Usable length	90 cm and 135 cm						

Ordering Information	<b>Stent</b> ø (mm)	<b>Catheter length 90 cm</b> (Stent length mm)									
		20**	30	40	60	80	100	120	150	170	200
	4.0	430437	430438	430439	430440	430441	430442	430443	430444	430445	430446
	5.0	430447	430448	430449	430450	430451	430452	430453	430454	430455	430456
<b>4F</b>	6.0	430457	430458	430459	430460	430461	430462	430463	430464	430465	430466
	7.0	430467	430468	430469	430470	430471	430472	430473	430474	430475	430476
	Stent ø (mm)	<b>Catheter length 135 cm</b> (Stent length mm)									
		20**	30	40	60	80	100	120	150	170	200
	4.0	430477	430478	430479	430480	430481	430482	430483	430484	430485	430486
	5.0	430487	430488	430489	430490	430491	430492	430493	430494	430495	430496
<b>4F</b>	6.0	430497	430498	430499	430500	430501	430502	430503	430504	430505	430506
	7.0	430507	430508	430509	430510	430511	430512	430513	430514	430515	430516
								*:	*8 week	s pre-or	der only

1. BIOTRONIK data on file. 6.0 mm diameters; 2. BIOTRONIK data on file. 6.0 mm diameters. Supera stent not possible to test due to its design and applied test method ; 3. Zhao HQ Late stent expansion and neointimal proliferation of oversized nitinol stents in peripheral arteries. Cardiovasc. Interv. Radiol. 2009; 32(4); 720-6; 4. Koskinas C. Role of endothelial shear stress in stent restenosis and thrombosis: pathophysiologic mechanisms and implications for clinical translation. JACC 2012 10;59(15):1337-49; 5. Koppara T. Thrombogenicity and early vascular healing response in metallic biodegradable polymer-based and fully bioabsorbable drug-eluting stents. Circ Cardiovasc Interv. 2015 8(6):e002427; 6. Funovics M. Correlation between chronic outward force (COF) and neointimal hyperplasia in self-expanding nitinol stents in swine in clinically relevant oversizing ranges. Presented at: LINC, Jan 26, 2017; Leipzig, Germany; 7. Bosiers M et al. 4-French – compatible endovascular material is safe & effective in the treatment of femoropopliteal occlusive disease: Results of the 4EVER Trial. ENDOVASC THER 2013; 20: 746-756; 8. BIOTRONIK data on file; 9. Lichtenberg M. Superficial Femoral Artery TASC D registry: 12-month effectiveness analysis of the Pulsar-18 SE nitinol stent in patients with critical limb ischemia. J Cardiovasc Surg (Torino). 2013 ; 54(4):433-9; 10. Lichtenberg et al. Effectiveness of the Pulsar-18 self-expanding stent with optional drug-coated balloon angioplasty in the treatment of femoropopliteal lesions - the BIOFLEX PEACE All-Comers Registry.Vasa (2019), 1-9. doi\_10.10240301-1526a000785; 11. Supera IFU, EL2100430 (2016-03-23); 12. Bosiers M. 4EVER 24 month results: long-term results of 4F Pulsar stent in femoropopliteal lesions. Presented at: CIRSE 2013; Barcelona, Spain; 13. Bunte Metal. in STROLL Catheterization and Cardiovascular Interventions 2018; 92:106-114; 14. Laird J et al. RESILIENT SFA nitinol stenting. JET 2012;19:1-9; 15. Dake M et al. Durable clinical effectiveness with paclitaxel-eluting stents in the femoropopliteal artery: 5-year results of the Zilver PTX randomized trial. Am Heart Assoc 133.15 (2016): 1472-1483. doi: 10.1161/CIRCULATIONAHA.115.016900; 16. Rocha-Singh et al. DURABILITY II Three-Year Follow-up. Catheterization and Cardiovascular Interventions 2015; 86:164-170.

Leading competitors have been selected based on the PV Stent Revenue Market Shares EU, 2017 and PV Revenue Market Shares APAC 2015; (Source: Millennium Research Group Inc.). Latest SFA self expanding stents for each manufacturer; Zilver and Zilver Flex are trademarks or registered trademarks of Cook Medical Technologies or its affiliates. Innova is a trademark or registered trademark of Boston Scientific or its affiliates. Everflex and Entrust are trademarks or registered trademarks or registered trademark of C. R. Bard or its affiliates. Supera is a trademark or registered trademark of the Abbott Group of Companies. S.M.A.R.T. Control is a trademark or registered trademark of Cardinal Health or its affiliates.

\*Indication as per IFU.

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