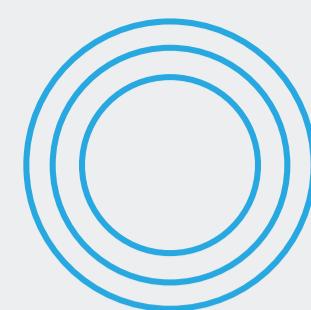




140 μm
thin struts



Clinically proven



Tri-axial delivery
system



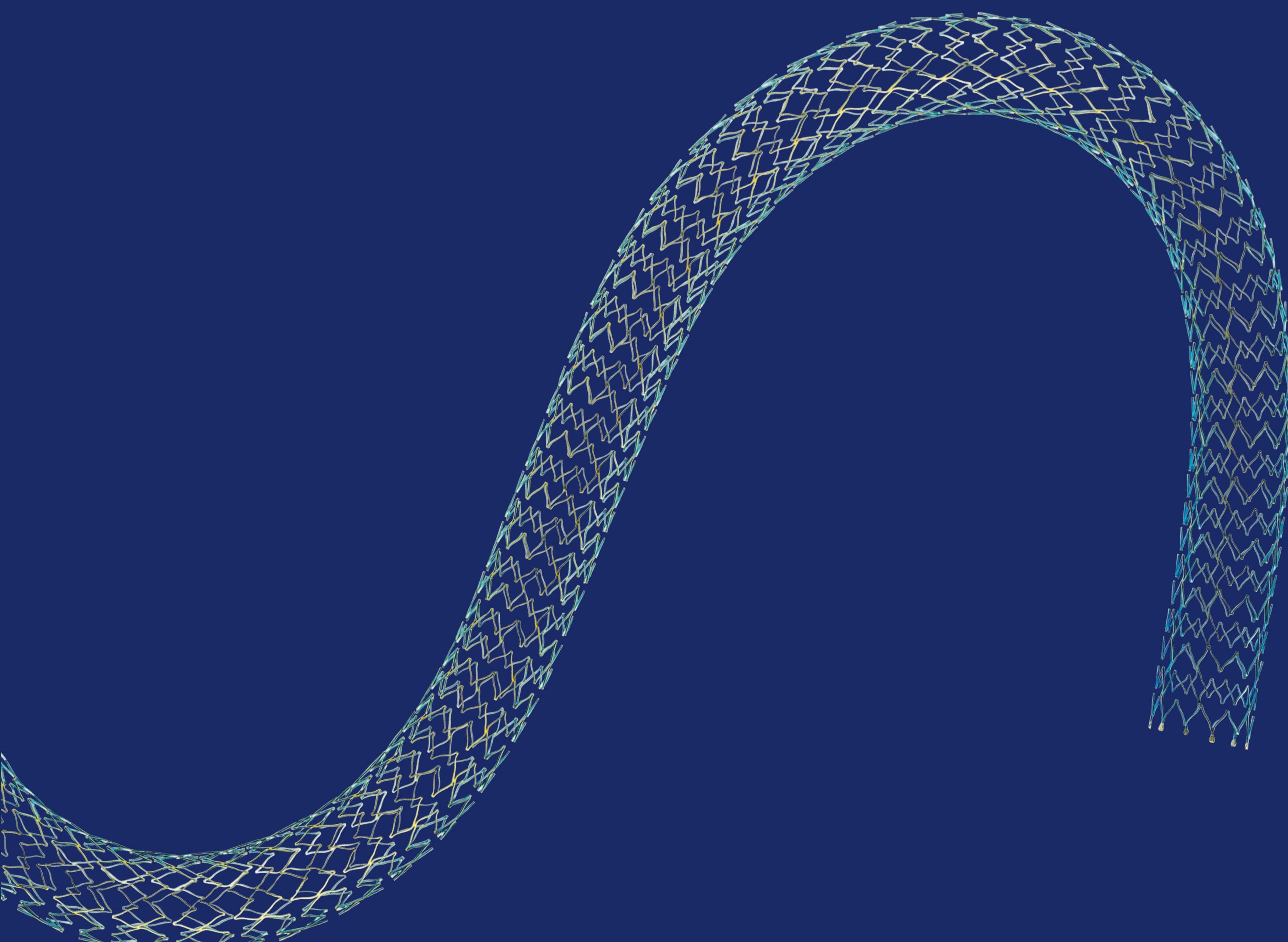
Technical data /
ordering info

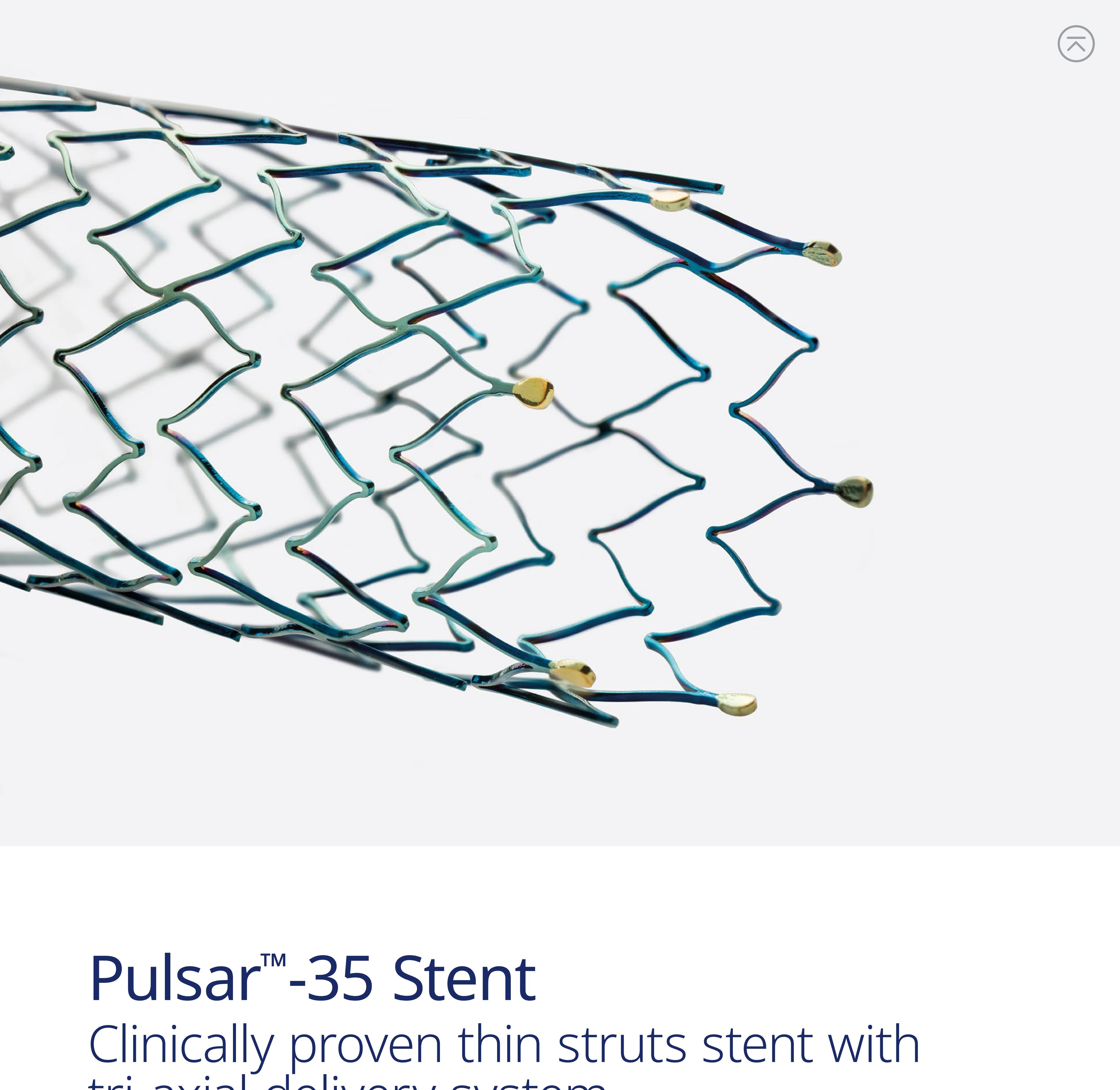
Teleflex™

Empowering the future of healthcare

Pulsar™-35

Self-Expanding
Stent System

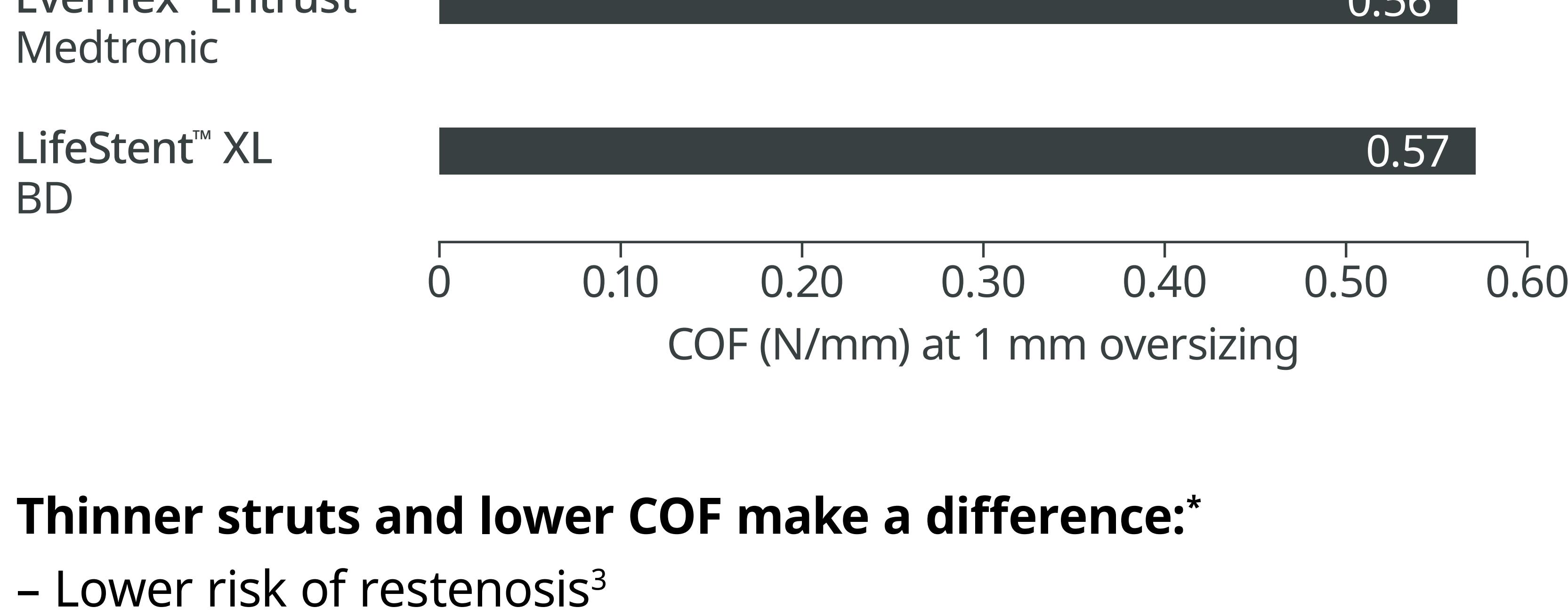




Pulsar™-35 Stent

Clinically proven thin struts stent with tri-axial delivery system.

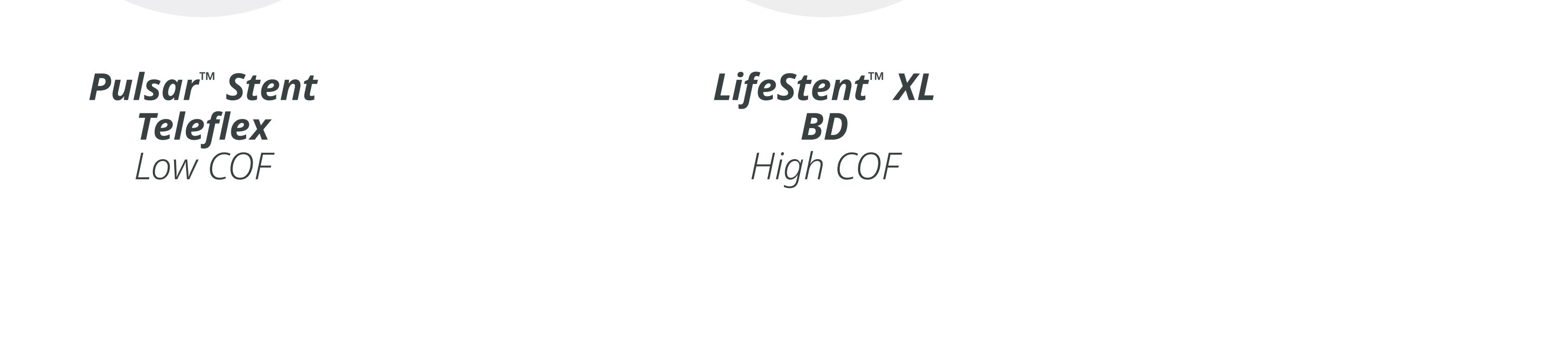
140 μm thin struts – thinner than the leading brands¹



Thinner struts and lower COF make a difference:^{*}

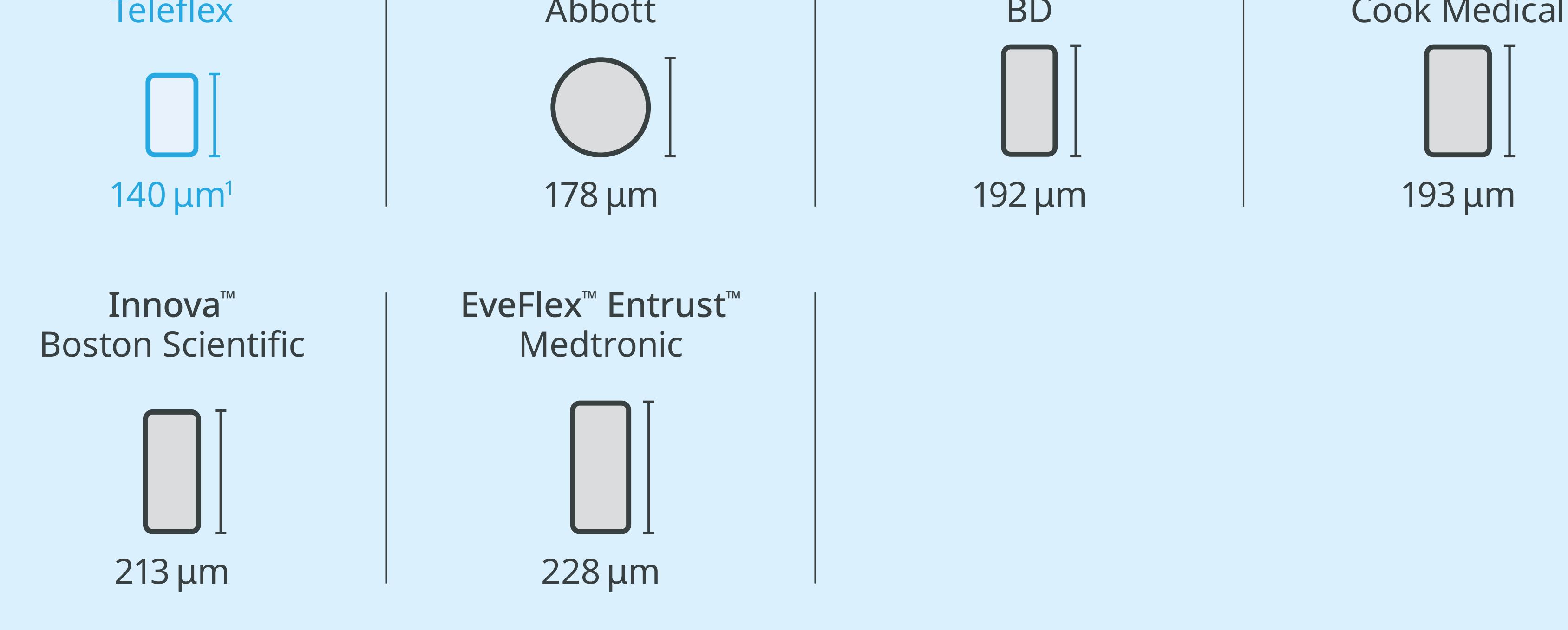
- Lower risk of restenosis³
- Reduced vessel injury and inflammation³
- Faster endothelialization^{4,5}

Vessel response on SE stent 1 mm oversizing showing neointimal hyperplasia at 90 days^{6*}



^{*}As demonstrated in pre-clinical studies

Stent strut thickness in perspective¹





Clinically proven

Long term safety and efficacy (24-month data)

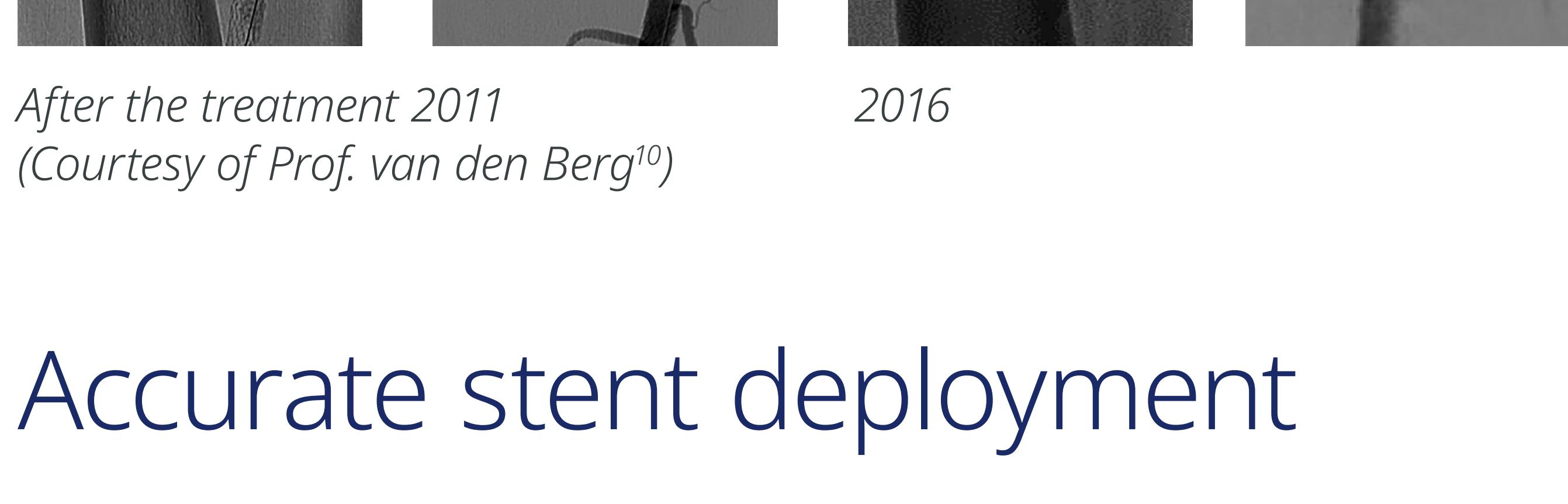
Clinically proven even in calcified lesions (4EVER), total occlusions (TASC D) and in all-comers registry (BIOFLEX PEACE).[¤]

	A.L.L ^{††}	12 MONTHS		24 MONTHS	
		PP [†]	FTLR ^{**}	PP [†]	FTLR ^{**}
ALL-COMERS BIOFLEX PEACE ⁷ (stent only)	8.2 cm	84.7 %	89.3 %	78.4 %	89.3 %
4F INTERVENTIONS 4EVER ⁸	7.1 cm	81.4 %	89.3 %	72.3 %	82.7 %
LONG & OCCLUDED TASC D ⁹	24.5 cm	77.0 %	86.0 %	-	-

[¤]Clinical outcomes of Pulsar™-18 stent can be used to illustrate clinical outcomes of Pulsar™-35 stent due to identical stent platforms



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



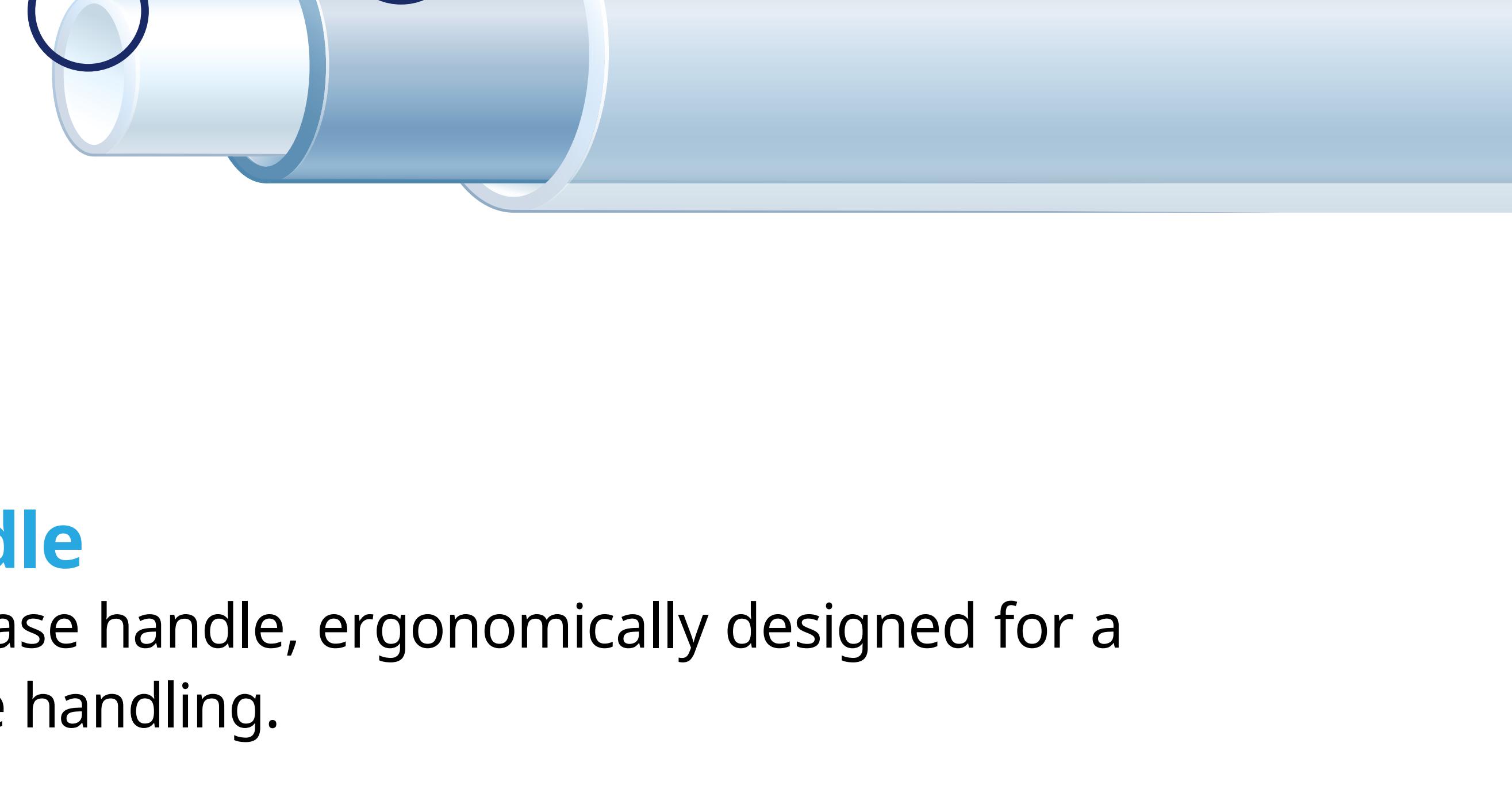
After the treatment 2011
(Courtesy of Prof. van den Berg¹⁰)

2016

Accurate stent deployment

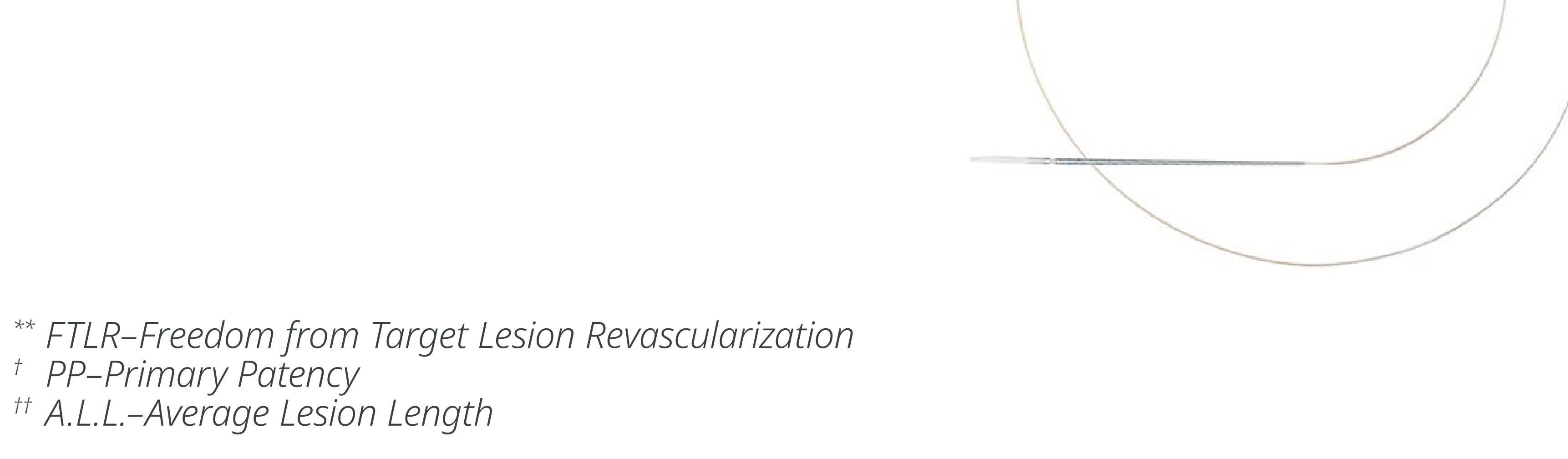
Tri-axial delivery system

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



Easy release handle

One-handed stent release handle, ergonomically designed for a comfortable and stable handling.

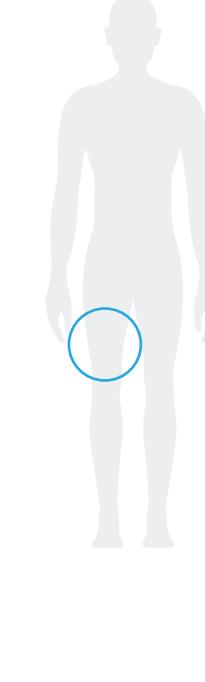


^{**} FTLR-Freedom from Target Lesion Revascularization

[†] PP-Primary Patency

^{††} A.L.L-Average Lesion Length





Pulsar™-35 Stent

Indicated for use in patients with atherosclerotic disease of the femoral and proximal popliteal arteries, in particular for the treatment of insufficient results after percutaneous transluminal angioplasty (PTA).*

Technical data

STENT	
Catheter type	OTW
Recommended guide wire	0.035"
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 5.0–7.0 mm; L: 30–200 mm
Proximal shaft	6F, hydrophobic coating
Usable length	90 and 135 cm

Ordering Information

STENT Ø	CATHETER LENGTH 90 CM; STENT LENGTH									
	20 mm	40 mm	60 mm	80 mm	100 mm	120 mm	150 mm	170 mm	200 mm	
5.0 mm	379878	379879	379880	379881	379917	379918	379919	379920	379921	
6F	6.0 mm	379883	379884	379885	379886	379922	379923	379924	379925	379926
	7.0 mm	379888	379889	379890	379891	379927	379928	379929	379930	379931

STENT Ø	CATHETER LENGTH 135 CM; STENT LENGTH									
	20 mm	40 mm	60 mm	80 mm	100 mm	120 mm	150 mm	170 mm	200 mm	
5.0 mm	379898	379899	379900	379901	379937	379938	379939	379940	379941	
6F	6.0 mm	379903	379904	379905	379906	379942	379943	379944	379945	379946
	7.0 mm	379908	379909	379910	379911	379947	379948	379949	379950	379951

* Indication as per IFU.

References:

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- 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 Data on file.

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.

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Teleflex Headquarters International, Ireland · Teleflex Medical Europe Ltd. · IDA Business & Technology Park

Dublin Road · Athlone · Co Westmeath · Tel. +353 (0)9 06 46 08 00 · Fax +353 (0)14 37 07 73 · orders.intl@teleflex.com

United Kingdom Tel. +44 (0)14 94 53 27 61 · info.uk@teleflex.com

South Africa Tel. +27 (0)11 807 4887 · assist.africa@teleflex.com

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