Protégé™ EverFlex™ Self-Expanding Peripheral Stent System ORDERING INFORMATION

Product Number	Product Number	Stent	Stent	Lumon	Sheath	Recommended	Crossing
Catheter Length 80 cm	Catheter Length 120 cm	Diameter (mm)	Length (mm)	Lumen Size (mm)	Size (Fr)	Guidewire (in)	Profile (in)
PRP35-05-020-080	PRP35-05-020-120	5	20	3.5 - 4.5	6	0.035	0.079
PRP35-05-030-080	PRP35-05-030-120	5	30	3.5 - 4.5	6	0.035	0.079
PRP35-05-040-080	PRP35-05-040-120	5	40	3.5 - 4.5	6	0.035	0.079
PRP35-05-060-080	PRP35-05-060-120	5	60	3.5 - 4.5	6	0.035	0.079
PRP35-05-080-080	PRP35-05-080-120	5	80	3.5 - 4.5	6	0.035	0.079
PRP35-05-100-080	PRP35-05-100-120	5	100	3.5 - 4.5	6	0.035	0.079
PRP35-05-120-080	PRP35-05-120-120	5	120	3.5 - 4.5	6	0.035	0.079
PRP35-05-150-080	PRP35-05-150-120	5	150	3.5 - 4.5	6	0.035	0.079
PRP35-06-020-080	PRP35-06-020-120	6	20	4.5 - 5.5	6	0.035	0.079
PRP35-06-030-080	PRP35-06-030-120	6	30	4.5 - 5.5	6	0.035	0.079
PRP35-06-040-080	PRP35-06-040-120	6	40	4.5 - 5.5	6	0.035	0.079
PRP35-06-060-080	PRP35-06-060-120	6	60	4.5 - 5.5	6	0.035	0.079
PRP35-06-080-080	PRP35-06-080-120	6	80	4.5 - 5.5	6	0.035	0.079
PRP35-06-100-080	PRP35-06-100-120	6	100	4.5 - 5.5	6	0.035	0.079
PRP35-06-120-080	PRP35-06-120-120	6	120	4.5 - 5.5	6	0.035	0.079
PRP35-06-150-080	PRP35-06-150-120	6	150	4.5 - 5.5	6	0.035	0.079
-	PRP35DR-06-200-120	6	200	4.5 - 5.5	6	0.035	0.079
PRP35-07-020-080	PRP35-07-020-120	7	20	5.5 - 6.5	6	0.035	0.079
PRP35-07-030-080	PRP35-07-030-120	7	30	5.5 - 6.5	6	0.035	0.079
PRP35-07-040-080	PRP35-07-040-120	7	40	5.5 - 6.5	6	0.035	0.079
PRP35-07-060-080	PRP35-07-060-120	7	60	5.5 - 6.5	6	0.035	0.079
PRP35-07-080-080	PRP35-07-080-120	7	80	5.5 - 6.5	6	0.035	0.079
PRP35-07-100-080	PRP35-07-100-120	7	100	5.5 - 6.5	6	0.035	0.079
PRP35-07-120-080	PRP35-07-120-120	7	120	5.5 - 6.5	6	0.035	0.079
PRP35-07-150-080	PRP35-07-150-120	7	150	5.5 - 6.5	6	0.035	0.079
-	PRP35DR-07-200-120	7	200	5.5 - 6.5	6	0.035	0.079
PRP35-08-020-080	PRP35-08-020-120	8	20	6.5 - 7.5	6	0.035	0.079
PRP35-08-030-080	PRP35-08-030-120	8	30	6.5 - 7.5	6	0.035	0.079
PRP35-08-040-080	PRP35-08-040-120	8	40	6.5 - 7.5	6	0.035	0.079
PRP35-08-060-080	PRP35-08-060-120	8	60	6.5 - 7.5	6	0.035	0.079
PRP35-08-080-080	PRP35-08-080-120	8	80	6.5 - 7.5	6	0.035	0.079
PRP35-08-100-080	PRP35-08-100-120	8	100	6.5 - 7.5	6	0.035	0.079
PRP35-08-120-080	PRP35-08-120-120	8	120	6.5 - 7.5	6	0.035	0.079
PRP35-08-150-080	PRP35-08-150-120	8	150	6.5 - 7.5	6	0.035	0.079
-	PRP35DR-08-200-120	8	200	6.5 - 7.5	6	0.035	0.079

INDICATIONS: The Protégé EverFlex Stent is indicated for use in common iliac, external iliac, superficial femoral, proximal popliteal, and subclavian arteries. Protégé EverFlex is a trademark of ev3 Inc.

Indications, contraindications, warnings and instructions for use can be found in the product labeling supplied with each device.

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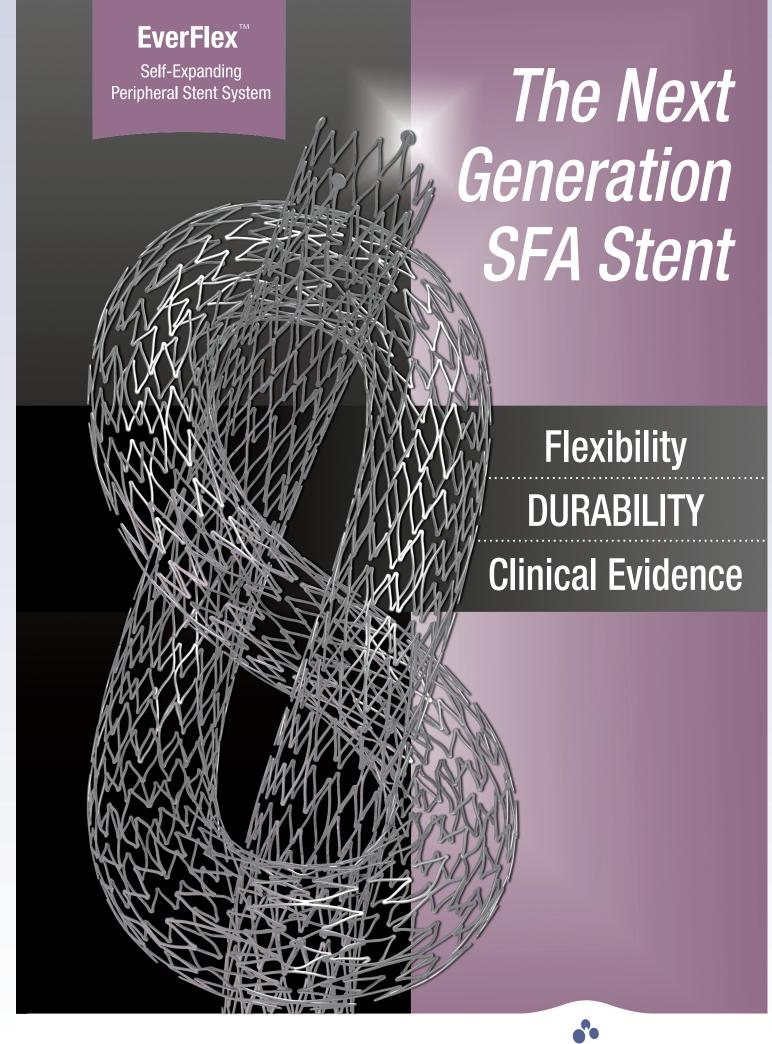
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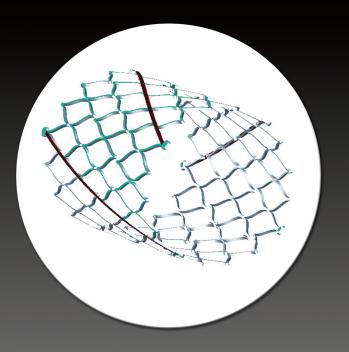


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True 2nd Generation stent design

EverFlex provides superb strength and excellent threedimensional flexibility for challenging SFA stenting.

- Spiral connections significantly improve flexibility and vessel conformability without sacrificing radial force.
- Minimal shortening during deployment



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durability



Superior DURABILITY

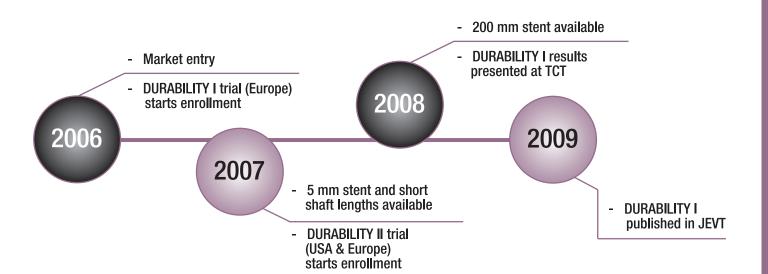
More wave peaks between each interconnection distribute stress across multiple struts, greatly enhancing fatigue performance, durability and uniformity of deployment.

Stent manufacturing and finishing:

Superb fatigue performance through

- Laser Cutting
- Microblasting
- Electropolishing
- Surface passivation
- Riveted markers

EverFlex: The Story of Innovation and Evidence



clinical evidence



Nitinol Stent Implantation in Long Superficial Femoral Artery Lesions:

12 MONTH RESULTS OF THE DURABILITY I STUDY

Bosiers M, et al: Journal of Endovascular Therapy 2009;16(3):261-269

Results (n=151 patients)				
Lesion Length:	96.4 mm			
Primary Patency:	72.2%			
Freedom from TLR:	79.1%			
Rutherford Class:	2.84 baseline to 0.60 at 1 year (significant)			
ABI Improvement:	0.6 baseline to 0.9 at 1 year (significant)			

141 out of 151 patients received only 1 single EverFlex stent.