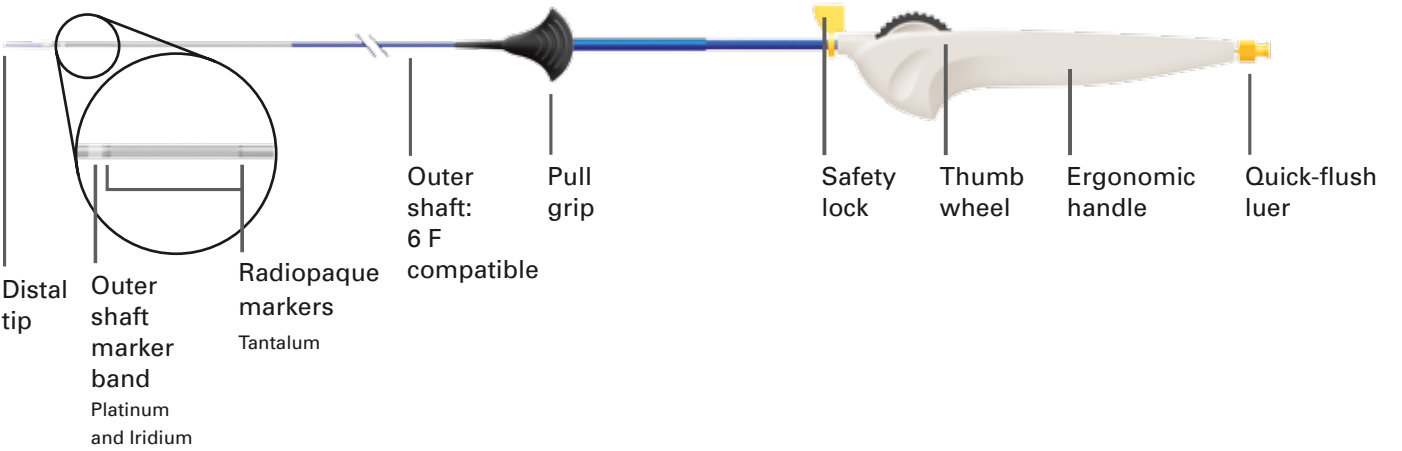


Epic™ Vascular Self-Expanding Stent System Product Codes

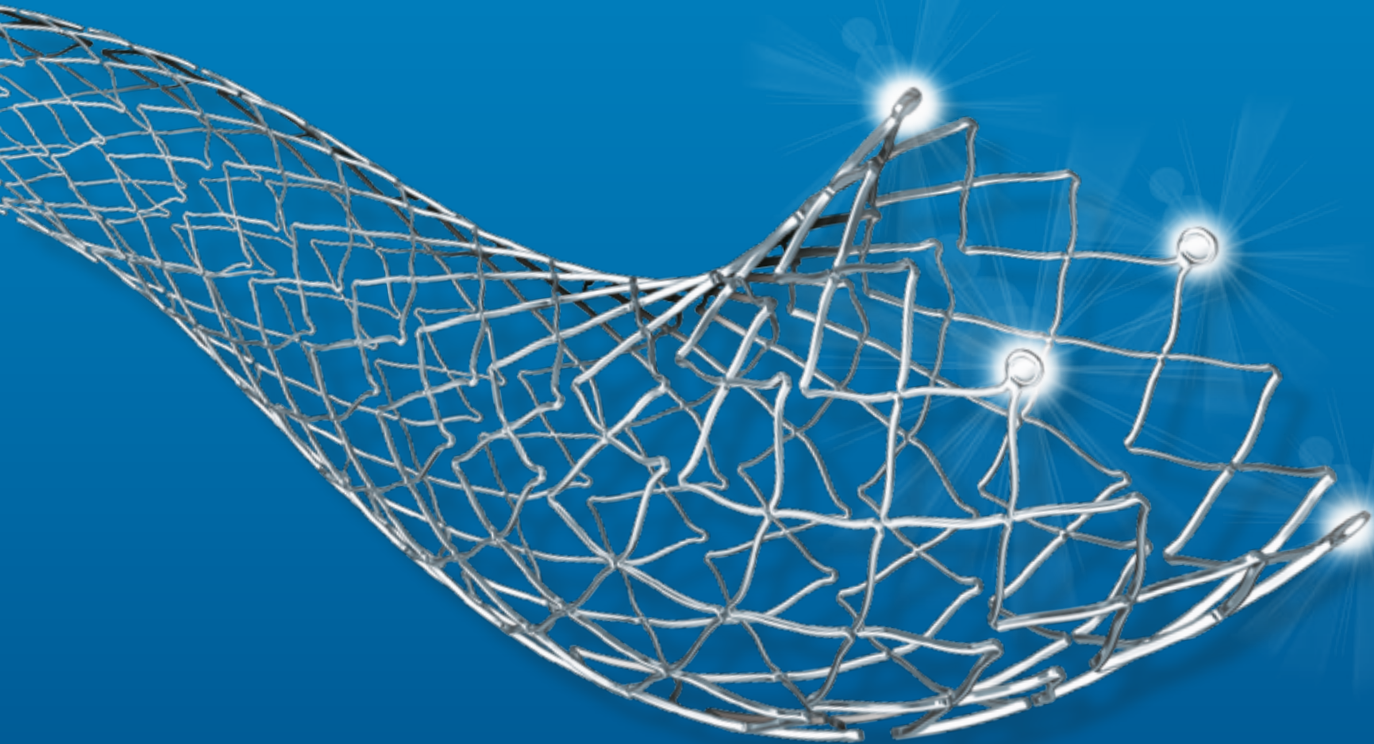
UPN Order Code		Diameter (mm)	Length (mm)	Sheath Compatibility (F/mm)
75 cm shaft	120 cm shaft			
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H74939054053070	H74939054053020	5	30	6/2.0
H74939054054070	H74939054054020	5	40	6/2.0
H74939054055070	H74939054055020	5	50	6/2.0
H74939054056070	H74939054056020	5	60	6/2.0
H74939054057070	H74939054057020	5	70	6/2.0
H74939054058070	H74939054058020	5	80	6/2.0
H74939054051070	H74939054051020	5	100	6/2.0
H74939054051270	H74939054051220	5	120	6/2.0
H74939054062070	H74939054062020	6	20	6/2.0
H74939054063070	H74939054063020	6	30	6/2.0
H74939054064070	H74939054064020	6	40	6/2.0
H74939054065070	H74939054065020	6	50	6/2.0
H74939054066070	H74939054066020	6	60	6/2.0
H74939054067070	H74939054067020	6	70	6/2.0
H74939054068070	H74939054068020	6	80	6/2.0
H74939054061070	H74939054061020	6	100	6/2.0
H74939054061270	H74939054061220	6	120	6/2.0
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H74939054075070	H74939054075020	7	50	6/2.0
H74939054076070	H74939054076020	7	60	6/2.0
H74939054077070	H74939054077020	7	70	6/2.0
H74939054078070	H74939054078020	7	80	6/2.0
H74939054071070	H74939054071020	7	100	6/2.0
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H74939054085070	H74939054085020	8	50	6/2.0
H74939054086070	H74939054086020	8	60	6/2.0
H74939054087070	H74939054087020	8	70	6/2.0
H74939054088070	H74939054088020	8	80	6/2.0
H74939054081070	H74939054081020	8	100	6/2.0
H74939054081270	H74939054081220	8	120	6/2.0
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H74939054095070	H74939054095020	9	50	6/2.0
H74939054096070	H74939054096020	9	60	6/2.0
H74939054097070	H74939054097020	9	70	6/2.0
H74939054098070	H74939054098020	9	80	6/2.0
H74939054091070	H74939054091020	9	100	6/2.0
H74939054102070	H74939054102020	10	20	6/2.0
H74939054103070	H74939054103020	10	30	6/2.0
H74939054104070	H74939054104020	10	40	6/2.0
H74939054105070	H74939054105020	10	50	6/2.0
H74939054106070	H74939054106020	10	60	6/2.0
H74939054107070	H74939054107020	10	70	6/2.0
H74939054108070	H74939054108020	10	80	6/2.0
H74939054101070	H74939054101020	10	100	6/2.0
H74939054123070	H74939054123020	12	30	6/2.0
H74939054124070	H74939054124020	12	40	6/2.0
H74939054125070	H74939054125020	12	50	6/2.0
H74939054126070	H74939054126020	12	60	6/2.0
H74939054143070	H74939054143020	14	30	6/2.0
H74939054144070	H74939054144020	14	40	6/2.0
H74939054145070	H74939054145020	14	50	6/2.0
H74939054146070	H74939054146020	14	60	6/2.0

Epic™ Vascular Self-Expanding Stent System



Product Specs	
Stent Diameters	5 to 14 mm
Stent Lengths	20 to 120 mm
Catheter Lengths	75, 120 cm
GuideWire Compatibility	0.035" (0.90 mm) OTW
Catheter Nominal Outer Diameter	0.079" (2.03 mm) to 0.077" (1.97 mm) (distal to proximal)

EPIC™ Vascular Self-Expanding Stent System



All cited trademarks are the property of their respective owners. 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 labeling supplied with each device. Information for the use only in countries with applicable health authority product registrations.

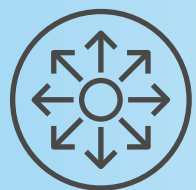
Illustrations are property of Boston Scientific Corporation. Illustrations are not necessarily to scale.
PI-274705-AB FEB2015 Printed in Germany by medicalvision.

Designed to provide **Exactly the Right Balance**



EXCEPTIONAL FLEXIBILITY

Hybrid Architecture Design with open- and closed-cell geometry is engineered to provide flexibility and deployment uniformity



BALANCED RADIAL FORCE

Macro and Micro Struts are engineered to work in tandem for balanced force – even in tortuous vasculature



EXCELLENT DEPLOYMENT ACCURACY

Intuitive Delivery System and Radiopaque Markers facilitate accurate deployment



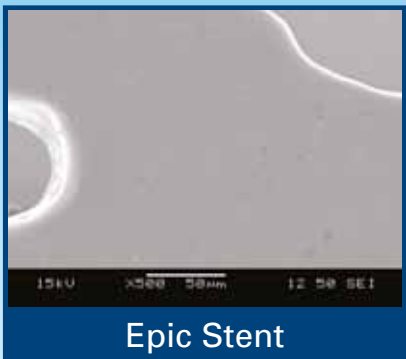
FRACTURE RESISTANCE

Top-Grade Nitinol with Meticulous Surface Finishing and Polishing contribute to superior fracture resistance

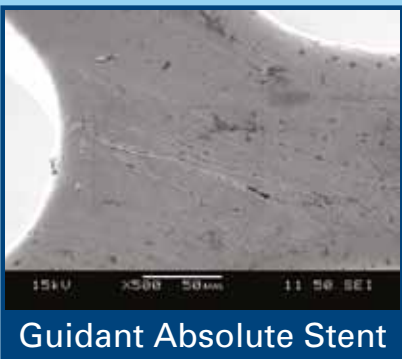
Designed to deliver **Superior Performance**

Meticulous Surface Finishing

Scanning Electron Microscopy x 500 magnification



Epic Stent



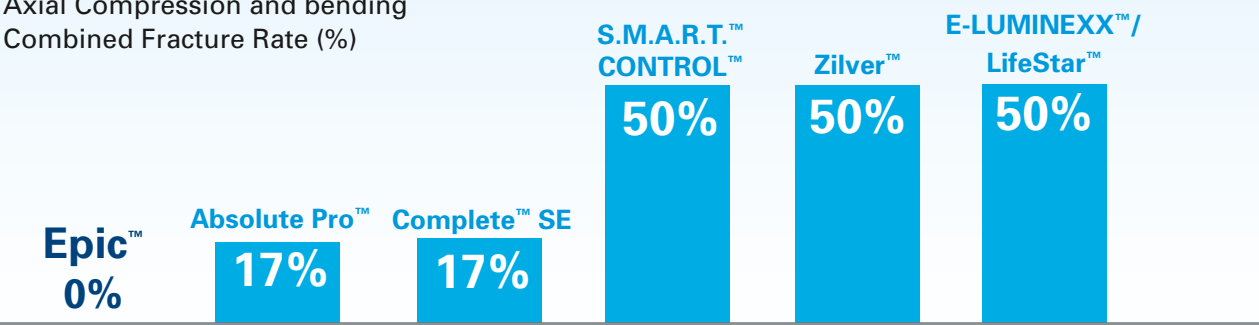
Guidant Absolute Stent

Produced by Boston Scientific. Data on file.

Comments and comparisons with competitor stent are made of date by BSC based on information provided from data on file.

Lower Combined Fracture Rate¹

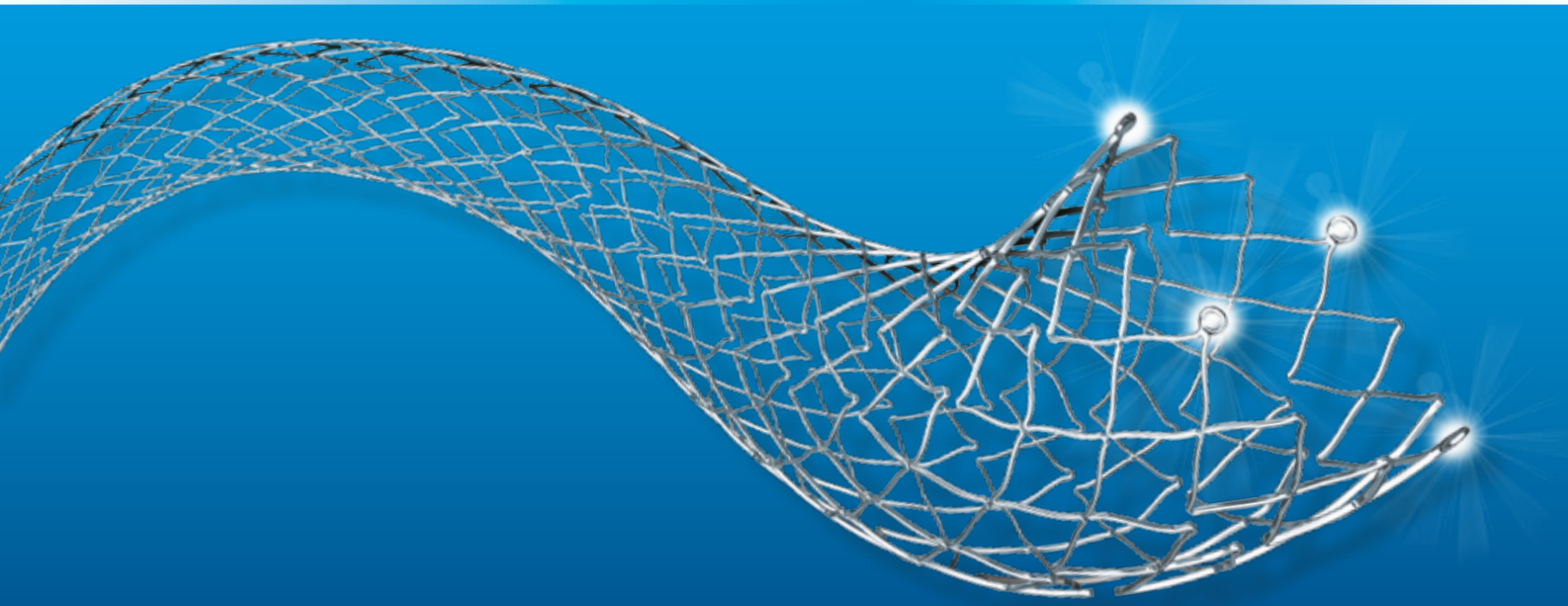
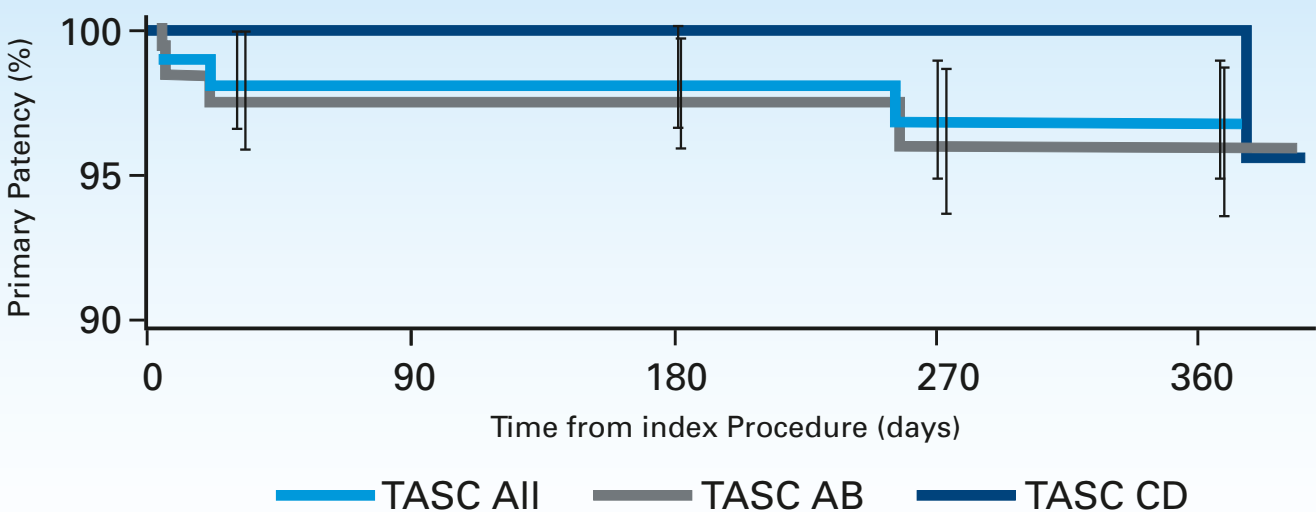
Axial Compression and bending Combined Fracture Rate (%)



THE GRAPHIC FOR FRACTURES ABOVE REFERS TO BENCH TESTING.

Is proven by **Clinical Results**

ORION is the **ONLY** Clinical Trial to demonstrate **similar patency across all TASC classifications**



Fractured stents are **2.8x more likely** to cause restenosis or reocclusion compared to non-fractured stents at 12 months²

¹Combined fracture rate includes any fracture whether associated with ≤6% axial compression or ≥57 mm radius bending deformation during fatigue testing. Axial compression and bending fatigue testing was performed at body temperature and at stent use diameter. N = 6. Bench testing performed by Boston Scientific. Data on file. Bench test results may not necessarily be indicative of clinical performance.

²D. Scheinert, S. Scheinert, et al. Prevalence and clinical impact of stent fractures after femoropopliteal stenting. *J Am Coll Cardiol*, 45 (2005), pp. 312–315.

Absolute Pro™ Vascular Self-Expanding Stent System is a trademark of Abbott Vascular
Complete™ SE Vascular Self-Expanding Stent System is a trademark of Medtronic
S.M.A.R.T.™ CONTROL™ Self-Expanding Nitinol Stent is a trademark of Cordis Corporation, a Johnson & Johnson Company
Zilver™ Vascular Self-Expanding Stent is a trademark of Cook Inc.
E-LUMINEXX™ Nitinol Transhepatic Biliary Endoprosthesis and Delivery System is a trademark of C. R. Bard, Inc.
LifeStar™ Biliary Stent System is a trademark of C. R. Bard, Inc.

89.9%
(n=139 lesions)

Freedom from TLR at 3 years with 0 Amputations

94.4%
(n=126 lesions)

Primary patency³ at 12 months

95.3%
(n=106 subjects)

Rutherford Becker classification improved by ≥1 at 12 months

100%
(n=166 lesions)

Technical success⁴

Note: All 12 month clinical data is from the *Journal of Endovascular Therapy*: April 2014, Vol. 21, No. 2, pp. 213-222. All 3 year clinical data is on file at Boston Scientific Corporation.

³Primary patency (per lesion) is defined as DUS SVR ≤2.5 with no target lesion revascularization, bypass of the target lesion, or amputation.

⁴Technical success (per lesion) is defined as ≤30% stenosis post-procedure.