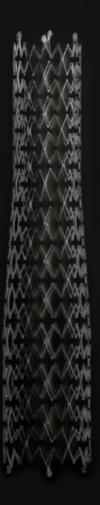
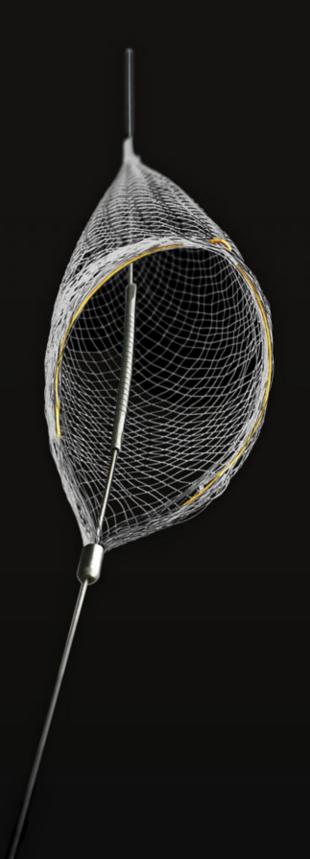
# **Carotid Innovations**

designed for confidence.



From initial lesion crossing to stent deployment and filter recovery.



## Predictable Deployment. Visible Results.

Protégé™ RX Carotid Stent System

#### Designed for Confidence in Carotid Artery Stenting

### **Predictable Deployment**

- Proprietary EX.P.R.T.™ Release Technology essentially eliminates premature deployment or jumping
- · No stent shortening
- Unique anatomically designed tapered stent for better fit in the carotid bifurcation
- 0.014" rapid exchange catheter with 6F low crossing profile and flexible atraumatic tip
- Radiopaque marker on catheter clearly indicates tapered location for precise positioning







- Tantalum GPS™ Markers enhance visibility for precise positioning and result confirmation
- · Cell design produces expansion force that resists compression while providing excellent wall apposition
- · Straight and tapered options for customized fit in carotid vessels

#### Filter the Risk™.

SpiderFX<sup>™</sup> Embolic Protection Device

#### **Guidewire of Choice**

 Works with any 0.014" or 0.018" guidewire to cross the most challenging carotid lesions

## **Extensive Portfolio**

 Treat vessels from 2 mm to 7 mm with a variety of sizes; available to use for Carotid, Coronary and Peripheral interventions



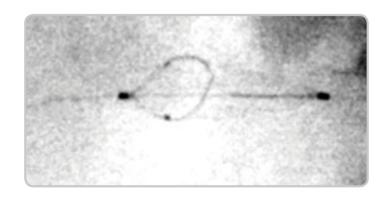
### **Excellent Stability**

- Controlled filter positioning throughout the intervention and during device exchanges
  - Braided nitinol design provides full-wall apposition
  - Capture wire designed to rotate and move longitudinally independent of the filter
  - Heparin coated filter provides up to 60 minutes patency



## **Enhanced Visibility**

 Clearly visible radiopaque markers and direct mouth indicator enable quick and controlled positioning of the filter throughout the intervention



#### Protégé™ RX Carotid Stent System

STRAIGHT				STRAIGHT			
Product Number	Diameter (mm)	Stent Length (mm)	Lumen Size (mm)	Product Number	Diameter (mm)	Stent Length (mm)	Lumen Size (mm)
SEPX-6-20-135	6	20	4.5-5.5	SEPX-9-40-135	9	40	7.5-8.5
SEPX-7-20-135	7	20	5.5-6.5	SEPX-10-40-135	10	40	8.5-9.5
SEPX-8-20-135	8	20	6.5-7.5	SEPX-6-60-135	6	60	4.5-5.5
SEPX-9-20-135	9	20	7.5-8.5	SEPX-7-60-135	7	60	5.5-6.5
SEPX-10-20-135	10	20	8.5-9.5	SEPX-8-60-135	8	60	6.5-7.5
SEPX-6-30-135	6	30	4.5-5.5	SEPX-9-60-135	9	60	7.5-8.5
SEPX-7-30-135	7	30	5.5-6.5	SEPX-10-60-135	10	60	8.5-9.5
SEPX-8-30-135	8	30	6.5-7.5	TAPERED			
SEPX-9-30-135	9	30	7.5-8.5				
SEPX-10-30-135	10	30	8.5-9.5	SEPX-8-6-30-135	8/6	30	(6.5-7.5)-(4.5-5.5)
SEPX-6-40-135	6	40	4.5-5.5	SEPX-8-6-40-135	8/6	40	(6.5-7.5)-(4.5-5.5)
SEPX-7-40-135	7	40	5.5-6.5	SEPX-10-7-30-135	10/7	30	(8.5-9.5)-(5.5-6.5)
SEPX-8-40-135	8	40	6.5-7.5	SEPX-10-7-40-135	10/7	40	(8.5-9.5)-(5.5-6.5)

135 cm catheter length / 0.078" stent crossing profile / 6F compatible 0.014" quidewire compatible

#### SpiderFX<sup>™</sup> Embolic Protection Device

•							
	COMPONENT		CAPTL	JRE WIRE	DELIVERY END	RECOVERY END	GUIDE CATHETER/ SHEATH
Product Number	Filter Size (mm)	Lumen Size (mm)	Wire Length OTW/RX (cm)	Wire Diameter (inch/mm)	Crossing Profile (F)	Diameter (F)	Minimum ID (inch)
SPD2-030-190	3.0	2.0-3.0	190	0.014/0.36	3.2	4.2	0.066
SPD2-030-320	3.0	2.0-3.0	320/190	0.014/0.36	3.2	4.2	0.066
SPD2-040-190	4.0	3.1-4.0	190	0.014/0.36	3.2	4.2	0.066
SPD2-040-320	4.0	3.1-4.0	320/190	0.014/0.36	3.2	4.2	0.066
SPD2-050-190	5.0	4.1-5.0	190	0.014/0.36	3.2	4.2	0.066
SPD2-050-320	5.0	4.1-5.0	320/190	0.014/0.36	3.2	4.2	0.066
SPD2-060-190	6.0	4.5-6.0	190	0.014/0.36	3.2	4.2	0.066
SPD2-060-320	6.0	4.5-6.0	320/190	0.014/0.36	3.2	4.2	0.066
SPD2-070-190	7.0	5.5-7.0	190	0.014/0.36	3.2	4.2	0.066
SPD2-070-320	7.0	5.5-7.0	320/190	0.014/0.36	3.2	4.2	0.066

#### Protégé Rx Indications:

Peripheral

The stent is indicated for use in occlusions, lesions at high risk for abrupt closure or threatened closure following percutaneous transluminal angioplasty (PTA); or lesions believed to be at high risk for restenosis following PTA in the common iliac, external iliac, or subclavian arteries. Stenting is intended to improve and maintain artery luminal diameter.

The stent is indicated for treatment of stenosis of the common carotid artery (CCA), internal carotid artery (ICA), and carotid bifurcation.

#### **SpiderFX Indications:**

The SpiderFX Embolic Protection Device provides distal embolization protection during general vascular use, including peripheral, coronary, and carotid interventions.

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

NEUROVASCULAR I PERIPHERAL VASCULAR Access - Balloons - Carotid - Embolic Coils - Embolic Protection - Liquid Embolics - Plaque Excision - Procedural Support - Retrieval Devices - Stents ev3 Neurovascular 9775 Toledo Way Irvine, CA 92618 USA ev3 SAS France ev3 S.r.l Italy PH +39 0267 977 61 FX +39 0266 711 637 ev3 Europe ev3 Corporate ev3 GmbH Germany, Austria International Headquarters **PH** +33 (0) 156 88 31 10 **FX** +33 (0) 156 88 31 11 **World Headquarters** PH +49 228 528 830 Peripheral Vascular 3033 Campus Drive Plymouth, MN 55441 106-108 rue La Boétie FX +49 228 528 8360 **PH** +1 949 837 3700 **FX** +1 949 837 2044 75008 Paris **ev3 B.V. Benelux PH** +31 (0) 433 659 223 **FX** +31 (0) 433 650 283 **ev3 Nordic AB PH** +46 859 000 950 **FX** +46 859 000 959 ev3 Ltd. United Kingdom PH +33 156 88 59 10 FX +33 156 88 59 11 PH +44 1279 659 900 FX +44 1279 654 900 UŚA **PH** +1 763 398 7000 **FX** +1 763 398 7001 ev3 International Distribution Centre ev3 Technologies Iberica, S.L. Spain PH +34 91 656 7154 Europalaan 25 6199 AB Maastricht-Airport **ev3 Sp z o.o. Poland PH** +48 32 747 01 44 **FX** +48 32 747 01 45 The Netherlands **PH** +31 (0) 433 659 220 **FX** +31 (0) 43 364 6395 FX +34 91 656 7214 www.ev3.net

