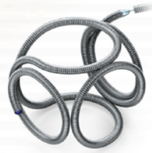
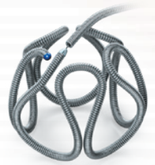
Target™ Detachable Coils family product offering

All deliverable through Excelsior™ SL-10™ and XT-17™ Microcatheters



360 shape



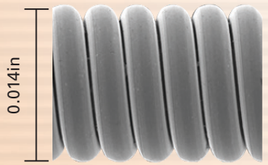
3D shape

Helical shape

Secondary OD (mm)	Target XL Standard	Target XL Soft	Standard	Soft	Ultra	Nano	Standard	Target XL Soft	Ultra	Nano
	Coil length (cm)									
24	50									
22	50									
20	50	50								
18	50	50								
16	50	50								
15			40							
14	50	50	30	30						
13			30	30						
12	45	45	30	30						
11			30	30						
10	40	40	30	30			30	40		
9	30	30	30, 20	30, 20				30		
8	30	30	30, 20	30, 20			25	30		
7	20	20	30, 20, 15	30, 20, 15			15	20		
6	20	20	30, 20, 15	30, 20, 15, 10			15	20		
5		15, 10	20, 15	20, 15, 10	15, 10		10	15		
4.5				12	10				8	
4		12, 8	15, 10, 8	15, 10, 8, 6	15, 10, 8, 6		8		15, 8, 6	
3.5				10	8	6			6	
3		9, 6	8, 6	10, 8, 6	10, 8, 6, 4	6, 4	6		10, 8, 6, 4	8, 6, 4
2.5					4	4			6, 4, 3	6, 4, 3
2		6, 3		4	6, 4, 3	4, 3			8, 6, 4, 3, 2, 1	6, 4, 3, 2
1.5						4, 3, 2				4, 3, 2, 1
1						3, 2				3, 2, 1

Size and softness

Target XL™ Coils



Target XL Standard

Target XL Soft

10-Size Coils

Target™
DETACHABLE COILS

Target™ Detachable Coil

See package insert for complete indications, contraindications, warnings and instructions for use.

INTENDED USE / INDICATIONS FOR USE

Target Detachable Coils are intended to endovascularly obstruct or occlude blood flow in vascular abnormalities of the neurovascular and peripheral vessels.

Target Detachable Coils are indicated for endovascular embolization of:

- Intracranial aneurysms
- Other neurovascular abnormalities such as arteriovenous malformations and arteriovenous fistulae
- Arterial and venous embolizations in the peripheral vasculature

THIS DOCUMENT IS INTENDED SOLELY FOR THE USE OF HEALTHCARE PROFESSIONALS.

A physician must always rely on his or her own professional clinical judgment when deciding whether to use a particular product when treating a particular patient. Stryker does not dispense medical advice and recommends that physicians be trained in the use of any particular product before using it in a procedure. The information presented is intended to demonstrate the breadth of Stryker product offerings. A physician must always refer to the package insert, product label and/or instructions for use before using any Stryker product. Products may not be available in all markets because product availability is subject to the regulatory and/or medical practices in individual markets. Please contact your Stryker representative if you have questions about the availability of Stryker products in your area. The Stryker products listed above are CE marked according to the Medical Device Directive 93/42/EEC.

AUS **Australian Sponsor Address**
Stryker Australia Pty Ltd
8 Herbert Street
St Leonards, NSW 2065
Australia

Stryker Neurovascular
47900 Bayside Parkway
Fremont, CA 94538
strykerneurovascular.com

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