



**A REVOLUTIONARY STENT SYSTEM
TO TREAT COMPLEX CORONARY LESIONS**

BIODEGRADABLE DRUG ELUTION

X² LIMUS[®]

**Sirolimus-eluting cobalt chromium
coronary stent technology.**



Unmatched

deliverability and
XLIMUS performance.



XLIMUS®

The most innovative Sirolimus-eluting cobalt chromium coronary stent technology available.



Leading the future of drug-eluting coronary stenting.

- XLIMUS is the ultimate coronary DES stent system to treat complex coronary artery disease by reaching and crossing the most challenging lesions.
- XLIMUS offers a stable, flexible stent delivery system featuring a flexible tip technology allowing navigating the most tortuous coronary anatomies.
- XLIMUS is a next generation thin stent strut Sirolimus DES, using the No. 1 drug which demonstrated long-term patient safety and optimal clinical efficacy, in more than 10.0 Million Patients.

MACE 0%

0.0029"

optimized thin stent strut design

Perfectly

homogenous vessel wall scaffolding

L-605

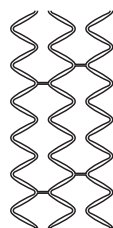
Cobalt Chromium

The outstanding
XLIMUS 6-8-10
stent cell design.

Ensures even vessel wall coverage.
Any different artery lesion diameter ranging
from 2.25 up to 5.00mm is stented evenly.

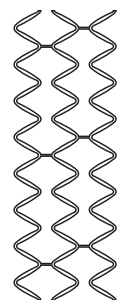
XLIMUS®

6 cells



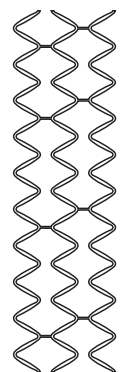
A1 2.25-2.50

8 cells



A2 2.75-3.50

10 cells



A3 4.00-5.00

Homogenous deployment distribution: Surface A1 = A2 = A3



No stent flaring.
No tissue prolapse.
Clinically effective.

Homogenous, clinically effective drug delivery optimizes the anti-proliferative protection of the stented lesion segment.

No stent strut flaring. No open gaps.

The technically high standard of 6-8-10 intermediate and closed-cell stent architecture covers all vessel diameters evenly. XLIMUS quality ensures the best possible intracoronary stenting stability and minimizes stenting trauma and restenosis. Extraordinary homogenous vessel wall scaffolding.

XLIMUS assists the cardiologist

with an optimal, unsurpassed tracking performance. It has an innovative Hydrophilic-coated shaft and an extra-low tip profile to access the most tortuous lesions. The ultra-low lesion crossing profile measures only 0.90 mm. The novel XLIMUS Sirolimus-eluting coronary stent system protects the stented lesion segment through extraordinary homogeneous vessel wall scaffolding which minimizes the risk of tissue prolapse.



XLIMUS drug-coating technology.

A novel drug-eluting, cobalt chromium coronary stent system, which provides clinically effective antiproliferative, drug delivery to the coronary artery lesion to prevent restenosis, followed by a rapid functional endothelial healing.

● Biodegradable Polymer ● Sirolimus Drug Elution



Following the nature. Stent flexibility by design.

Pulse Synchronous Stent Dynamics respond to coronary artery movement, with every heart beat. Natural stent flexion minimizes friction and shear stress to avoid vessel wall trauma. For a lifetime patient safety!



Controlled biodegradable Sirolimus drug release for rapid functional endothelial healing.

The highly biocompatible Poly (lactic acid) drug containing release matrix degrades smoothly and provides an optimal release kinetic profile. Within 30 days, about 70% of the anti-proliferative drug is distributed into the surrounding arterial tissue of the stent struts, ensuring a highly effective inhibition of smooth muscle cell migration and proliferation. Pharmacokinetic study result confirm sustained anti-proliferative drug efficacy up to 120 days.

Thin stent struts minimize foreign body metal volume.

XLIMUS reduces the inflammatory signal potential for prevention of late restenosis.

Cypher	140 μm
Taxus Liberte	97 μm
Endeavor	91 μm
Xience V	81 μm
XLIMUS*	71 μm

Source: Peter Smits, MD, from the COMPARE trial presentation at TCT 2009. *XLIMUS low metal volume stenting. Caution! Thinner struts, much below 71 μm may influence stent stability.

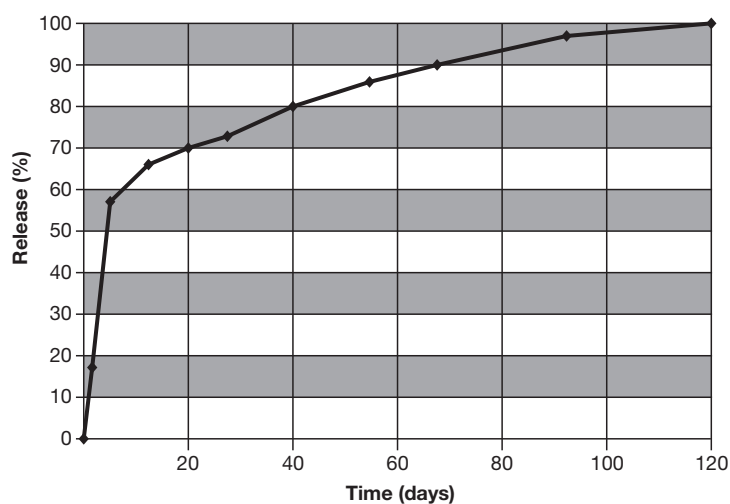
XLIMUS®



Life deserves the best.

XLIMUS controlled biodegradable Sirolimus drug release.

XLIMUS ensures a controlled drug release after stent implantation.



The widest range of 72 stent sizes.

Stent Length (mm)	Stent Diameter (mm)							
	2.25 mm	2.50 mm	2.75 mm	3.00 mm	3.50 mm	4.00 mm	4.50 mm	5.00 mm
8 mm	XL 2.25-8*	XL 2.50-8*	XL 2.75-8*	XL 3.00-8*	XL 3.50-8*	XL 4.00-8*	XL 4.50-8*	XL 5.00-8*
12 mm	XL 2.25-12	XL 2.50-12	XL 2.75-12	XL 3.00-12	XL 3.50-12	XL 4.00-12	XL 4.50-12*	XL 5.00-12*
16 mm	XL 2.25-16	XL 2.50-16	XL 2.75-16	XL 3.00-16	XL 3.50-16	XL 4.00-16	XL 4.50-16*	XL 5.00-16*
20 mm	XL 2.25-20	XL 2.50-20	XL 2.75-20	XL 3.00-20	XL 3.50-20	XL 4.00-20	XL 4.50-20*	XL 5.00-20*
24 mm	XL 2.25-24	XL 2.50-24	XL 2.75-24	XL 3.00-24	XL 3.50-24	XL 4.00-24	XL 4.50-24*	XL 5.00-24*
28 mm	XL 2.25-28	XL 2.50-28	XL 2.75-28	XL 3.00-28	XL 3.50-28	XL 4.00-28	XL 4.50-28*	XL 5.00-28*
32 mm	XL 2.25-32	XL 2.50-32	XL 2.75-32	XL 3.00-32	XL 3.50-32	XL 4.00-32	XL 4.50-32*	XL 5.00-32*
36 mm	XL 2.25-36	XL 2.50-36	XL 2.75-36	XL 3.00-36	XL 3.50-36	XL 4.00-36	XL 4.50-36*	XL 5.00-36*
40 mm	XL 2.25-40	XL 2.50-40	XL 2.75-40	XL 3.00-40	XL 3.50-40	XL 4.00-40	XL 4.50-40*	XL 5.00-40*

*upon request, no lead time available

Technical Data	
Material	Cobalt Chromium Alloy L-605
Total strut thickness	73µm (71µm Alloy + 2µm coating layer)
Coating layer	2µm
Device lengths	Stent length = balloon length = markers distance
Metal to artery ratio	14% average
Nominal pressure	8 ATM
Rated burst pressure	16 ATM except diameters 4.5 / 5.0 and diameter 4.0 with length higher than 20mm (14 ATM)
Average foreshortening	< 1%
Guiding catheter compatibility	5F (0.058" ID) except diameters 4.5 and 5.0 → 6F (0.071")
Guidewire compatibility	0.014" maximum recommended
Sirolimus (Rapamycin) drug-coating	1.25µg/mm ² stent surface



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