



SOLARIS[®]
SELF-EXPANDING



Innovation for life



MISSION

- Improve patients' quality of life
- Provide effective and Innovative solutions for healthcare professionals
- Make our customers feel unique and have outstanding customer service



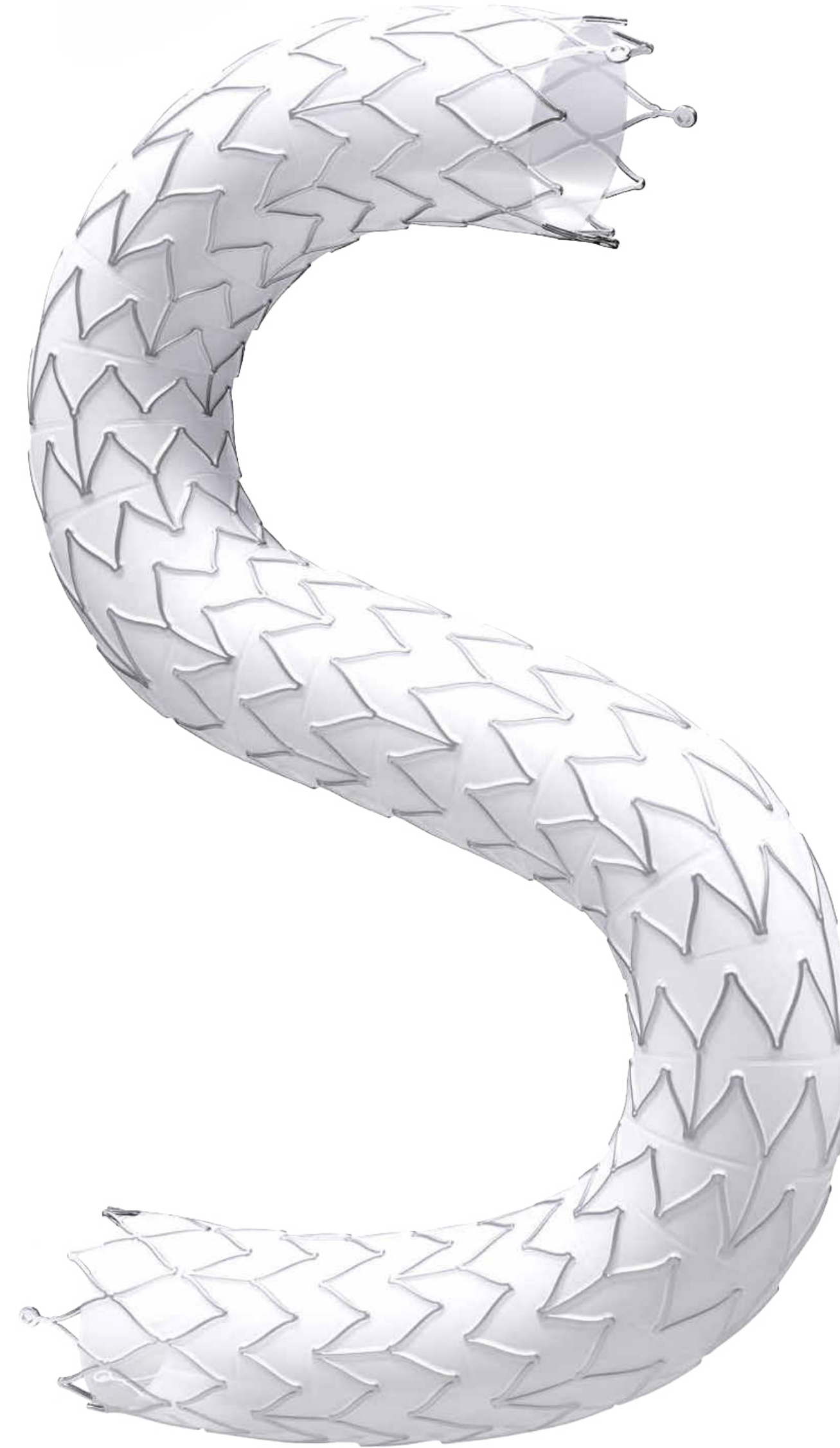
PRESENT IN MORE
THAN 45 COUNTRIES

SCITECH Medical is a minimally invasive medical device company that was founded over 18 years ago and is currently present in more than 45 countries. Through state-of-the-art technology and the use of the highest quality materials, tested and proven by the most rigorous international standards and clinical trials, SCITECH manufactures products that empower healthcare professionals to save or improve the quality of life of their patients.

For further information visit the website: scitechmed.com



The **SOLARIS** is a flexible, self-expanding endograft, comprised of a thin, multi-direction, durable electrospinning PTFE membrane encapsulating a Nitinol stent structure.



The device has been **engineered to effectively cover and instantaneously seal off** diseased tissue with a high multidirectional resistance membrane, providing an endoluminal bypass option for **physicians faced with complex lesions**. Its design provides high flexibility without compromising the requirement length, balanced radial force and low shortening rate. **The pull-back hydrophilic delivery system** provides superior navigability, and its anti-jumping system guarantees accurate deployment during the procedure.

HIGH FLEXIBILITY

PRECISION

BALANCED RADIAL STRENGTH

MINIMAL SHORTENING

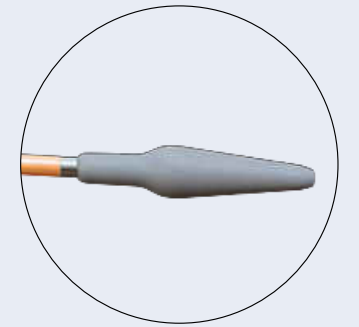


MULTIDIRECTIONAL
RESISTANCE STRENGTH WITH
INSTANTANEOUS SEALING

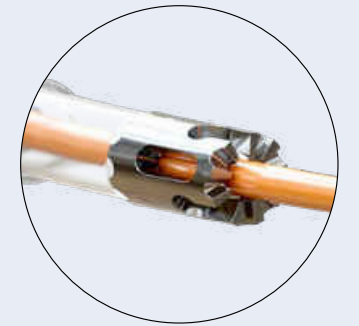
PULLBACK
DELIVERY
SYSTEM



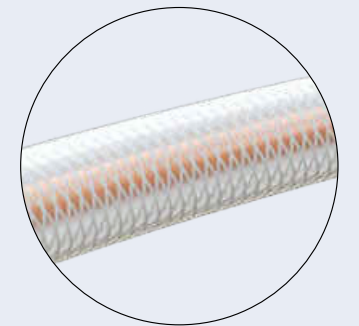
ATRAUMATIC
FLEXIBLE TIP



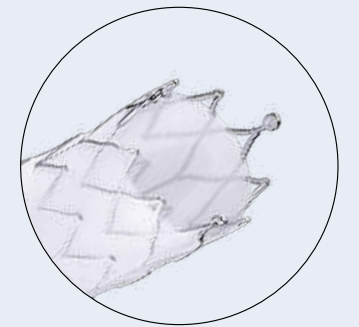
ANTI-JUMP
FEATURE



HYDROPHILIC
COATING



3 TANTALUM
MARKER BANDS
(DISTAL/PROXIMAL)





High Flexibility

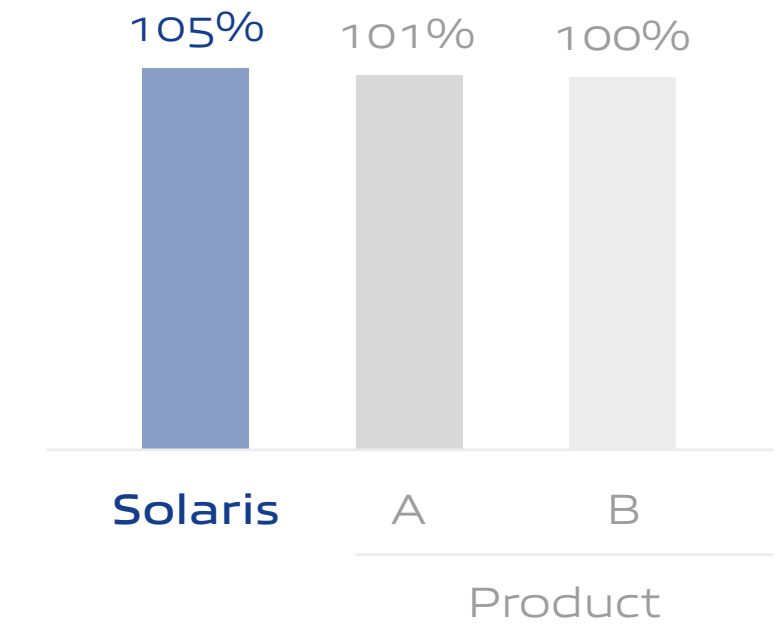
The **Solaris Endograft** is made from a laser cut Nitinol alloy. Due to its design, which maintains a distance between one cell and another, the product has high flexibility.

Excellent Radial Force

Bench tests showed excellent radial force for 1mm Stent compression and it is comparable to competitor values.



RADIAL FORCE FOR
1MM COMPRESSION



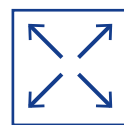
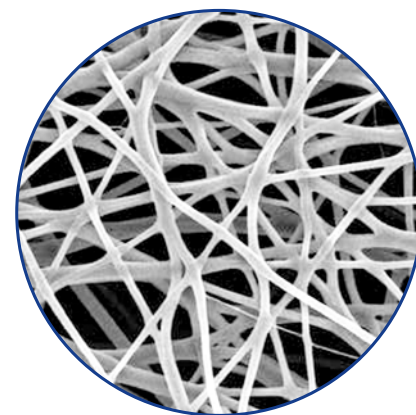
RADIAL FORCE

High strength and elasticity electrospinning PTFE Membrane

The Solaris Endograft is encapsulated both externally and internally by a thin PTFE membrane, produced by electrospinning process. Proven in bench tests, the Solaris Endograft membrane has high resilience and elasticity.

ELECTROSPINNING MEMBRANE

SOLARIS
(Electrospinning PTFE)



Solaris Covering
Multidirecional Strength

UNIDIRECTIONAL
(ePTFE)



Should be compensated by
a multi ePTFE layers

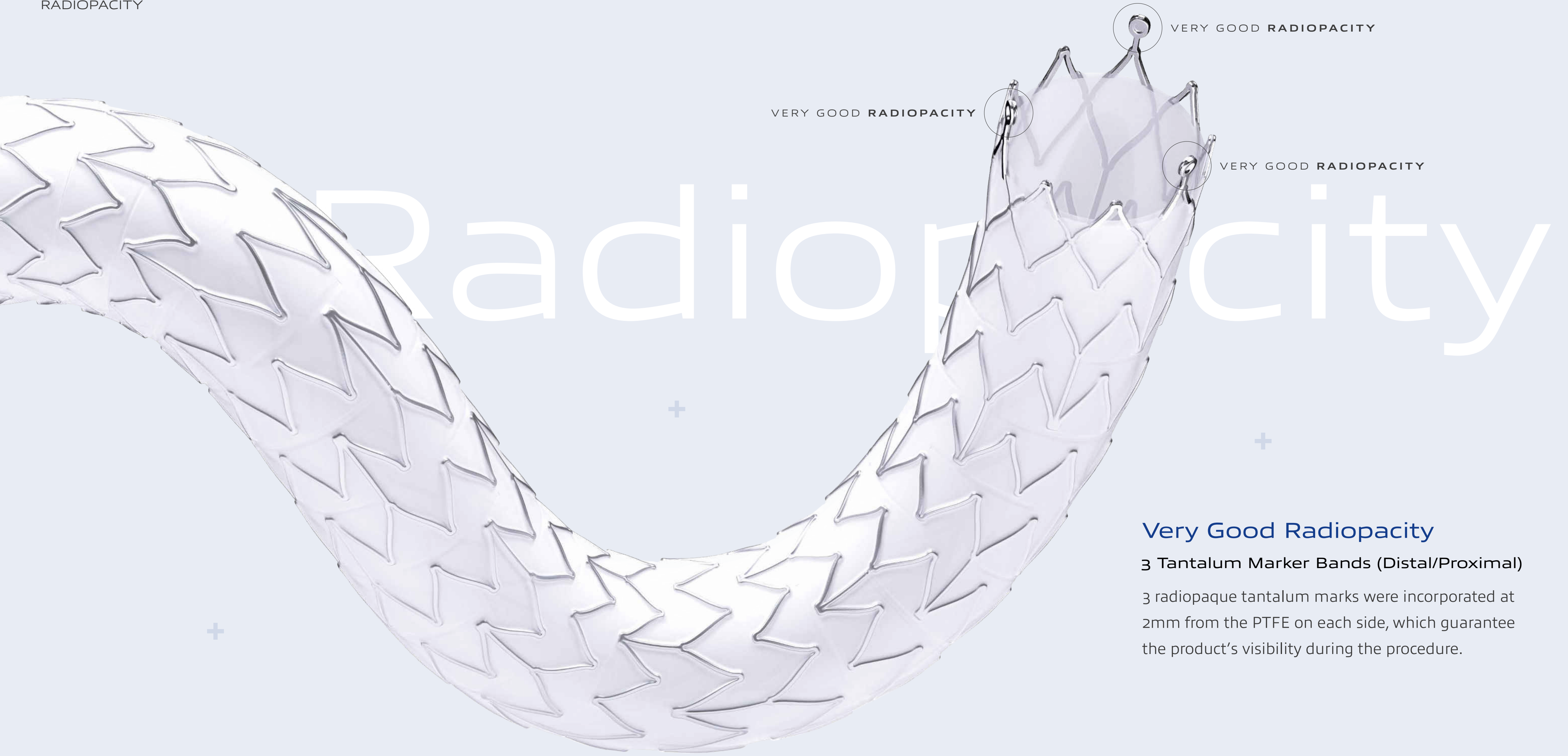
SOLARIS



X



MULTI EPTFE LAYER



VERY GOOD RADIOCAPACITY

VERY GOOD RADIOCAPACITY

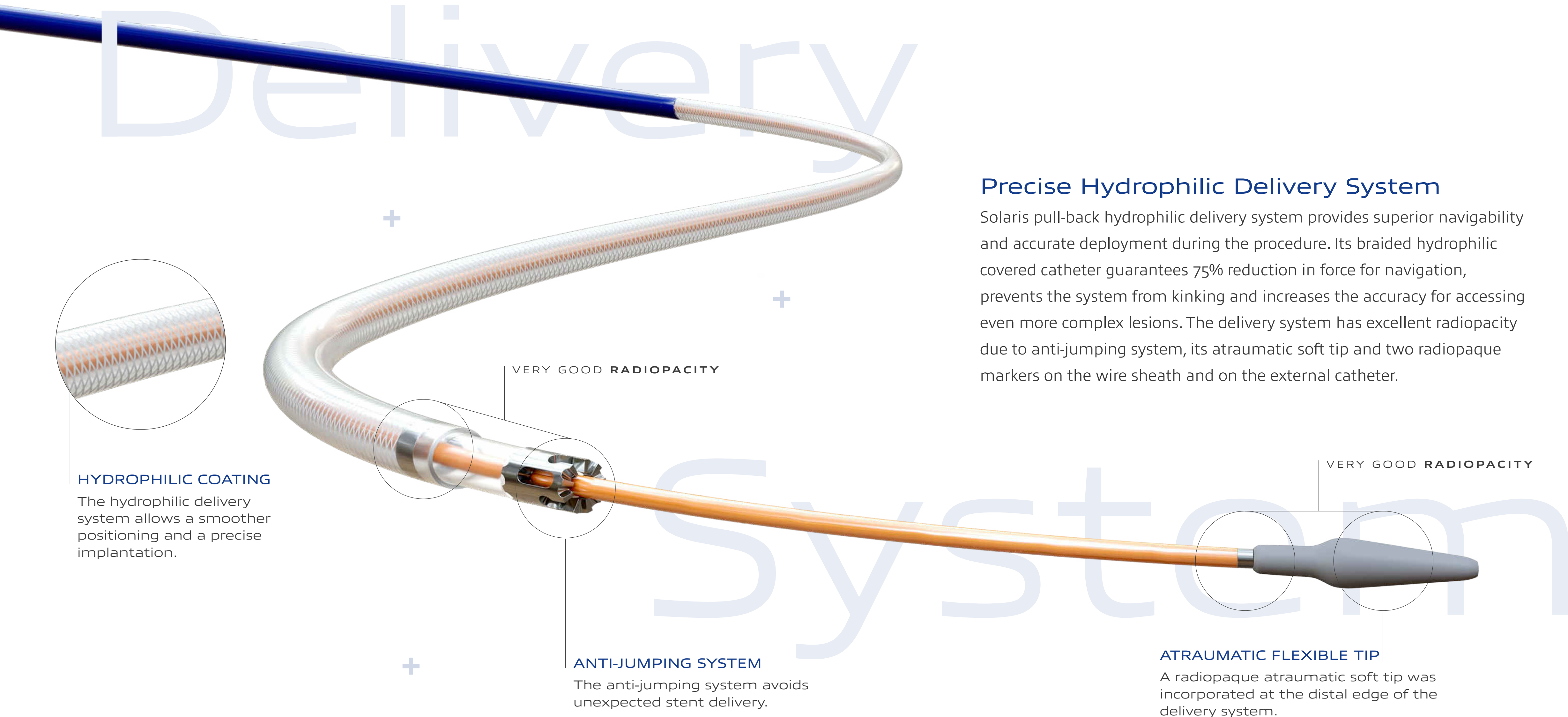
VERY GOOD RADIOCAPACITY

Radiopacity

Very Good Radiopacity

3 Tantalum Marker Bands (Distal/Proximal)

3 radiopaque tantalum marks were incorporated at 2mm from the PTFE on each side, which guarantee the product's visibility during the procedure.



HYDROPHILIC COATING

The hydrophilic delivery system allows a smoother positioning and a precise implantation.

VERY GOOD RADIOPACITY

ANTI-JUMPING SYSTEM

The anti-jumping system avoids unexpected stent delivery.

VERY GOOD RADIOPACITY

ATRAUMATIC FLEXIBLE TIP

A radiopaque atraumatic soft tip was incorporated at the distal edge of the delivery system.

Precise Hydrophilic Delivery System

Solaris pull-back hydrophilic delivery system provides superior navigability and accurate deployment during the procedure. Its braided hydrophilic covered catheter guarantees 75% reduction in force for navigation, prevents the system from kinking and increases the accuracy for accessing even more complex lesions. The delivery system has excellent radiopacity due to anti-jumping system, its atraumatic soft tip and two radiopaque markers on the wire sheath and on the external catheter.

SOLARIS
NEW FRENCH SIZE

8F Frenchs



New French size
8F for 5-8mm until
80mm of length.

Solaris Stent Graft implant in the treatment of a failed basilica loop arteriovenous fistula due to swing point stenosis.

Dr. Leonardo Harduin

BACKGROUND:

A 29-year-old female patient with SAH, SLE and CRF on hemodialysis for 8 months through a basilica loop arteriovenous fistula in the left arm. She started about 30 days ago with pain during hemodialysis sessions, increased venous resistance and increased bleeding time after hemodialysis sessions. She was admitted to the emergency room with disappearance of the fremitus in the AVF, hardening and local hyperemia, pain and puncture of the AVF with the release

of multiple clots, compatible with access thrombosis. Color Doppler ultrasound showed thrombosis of the entire basilic vein from the anastomosis with the brachial artery to the outflow in the axillary vein.

METHODS:

Access through dissection of the arteriovenous fistula. Thrombectomy with Fogarty 4F catheter. Diagnostic phlebography showed 90% stenosis at the swing point. Recanalization of the stenotic segment and 9F introducer implant. Pre-dilation of the

stenosed segment with a 7x30mm high pressure balloon. Preoperative phlebography with severe residual stenosis. Solaris 9x60mm Stent Graft implant and accommodation with a 9x30mm high pressure balloon.

RESULTS:

Control phlebography with good results and without residual stenosis or folds. Fremitus at the end of the 4+/4+ procedure. Hemodialysis performed by the AVF immediately after the procedure. Control color Doppler ultrasound (30 days) demonstrating the patency of the

stent and AVF flow volume of 1099 ml/min. Control phlebography at 6 months showing patent Stent Graft without stenosis.

CONCLUSION::

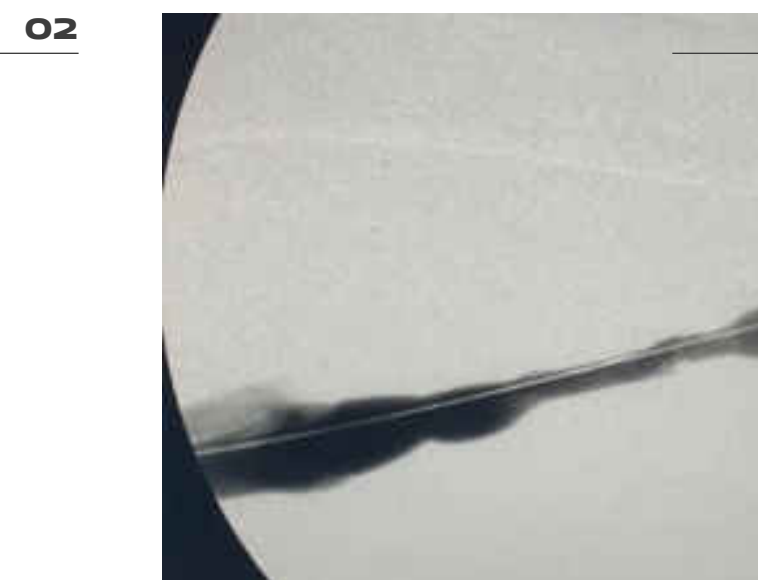
The use of the Solaris Stent Graft in the treatment of swing point lesions of the transposed basilic vein in failure was safe and effective, leading to an improvement in the quality of dialysis and maintenance of the access patency.



90% stenosis at Swing point.



Angioplasty with a 7x30mm high pressure balloon.



Preoperative phlebography with severe residual stenosis.



9x60mm Solaris Stent Graft implant.



Control phlebography without residual stenosis and with satisfactory result after post ballooning with a 9x30mm high pressure balloon.



Control phlebography at 6 months via femoral access demonstrating the patent Solaris stent graft without fractures.

Treatment of Subocclusive Atherosclerotic Lesion of the Left Subclavian Artery with Covered Stent

Prof. Dr. Paulo Eduardo Ocke Reis¹

CASE REPORT:

A 67-year-old female patient complaining of left upper limb (LUL) claudication reported limitation of simple activities such as brushing hair or washing hands. She presented worsening of pain and numbness complaint of the LUL despite the clinical treatment. Por "On physical examination, the absence of left brachial, radial and ulnar pulses was noticed. The angiotomography images (Figures I and II) confirmed the subocclusion of the left subclavian artery and a high degree of calcification of the lesion in the proximal segment of the artery.

TREATMENT:

After confirming the diagnosis, we

indicated an endovascular procedure with puncture access in the right femoral artery and revascularization of the left subclavian artery with a covered Solaris® stent. Demonstration before and after delivery of the covered stent (Figures III, IV, V). Six-month follow-up with angiotomography (figure VI).

CONCLUSION:

The Solaris® covered stent showed excellent navigation, delivery and precision in a sub-occluded and calcified artery. The immediate radiological and clinical result was satisfactory in the brachial, radial and ulnar arteries, with mediumterm broad pulses observed during patient follow-up.

FIGURE I

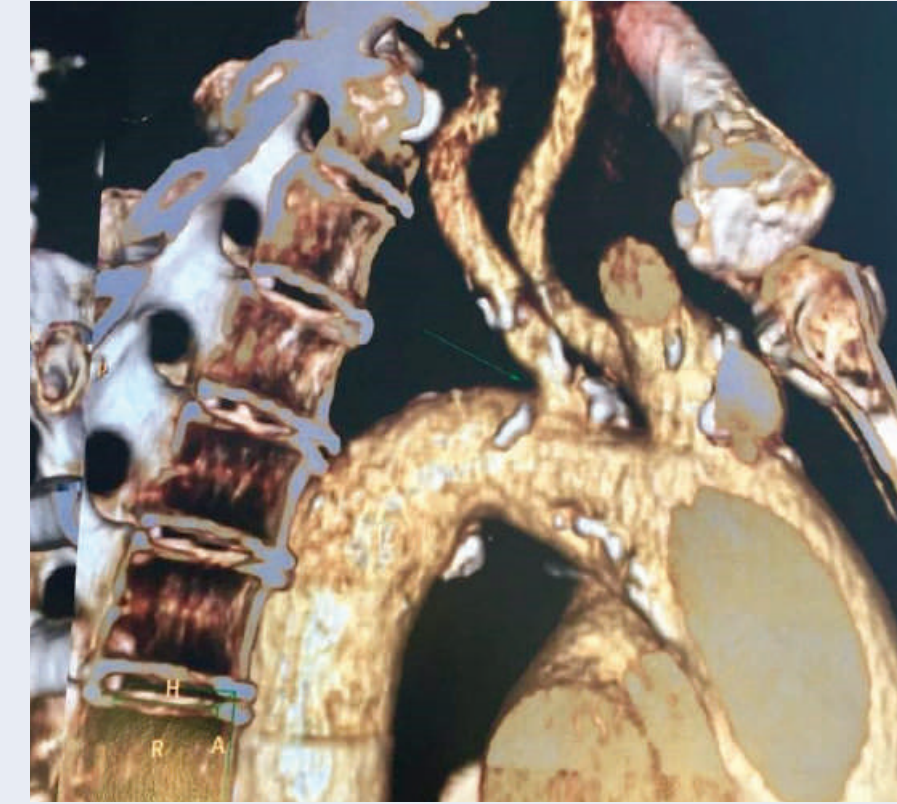


FIGURE II



Figures I and II - APre-procedure angiotomography of the left subclavian artery confirming subocclusion and calcification of the artery.

FIGURE III



FIGURE IV



Figures III and IV - After the right femoral access, preoperative arteriography confirms a high degree of stenosis and an irregular calcified ostial atherosclerotic plaque.

FIGURE V

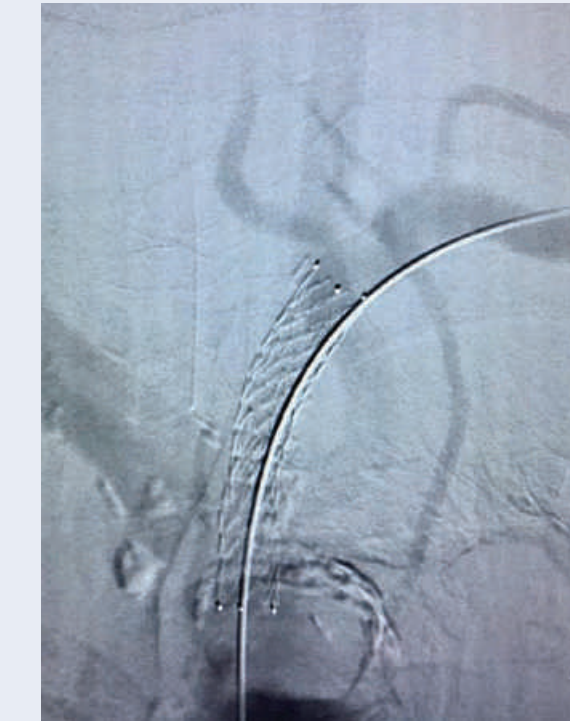
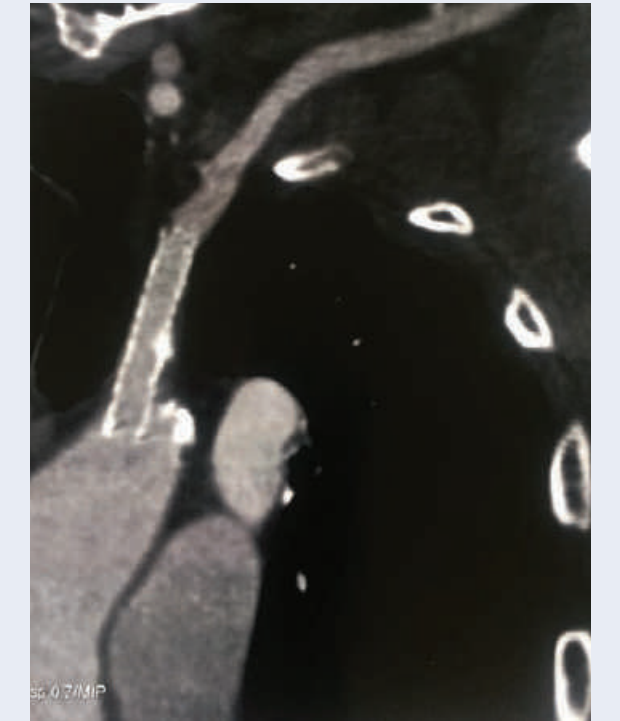


FIGURE VI



Figures V and VI - Final preoperative control and angiotomography after six months with excellent results. Lesion treatment and delivery of Solaris® 8x40mm safely and efficiently.

SOLARIS
ORDER INFORMATION



Diameter	Length							
	40mm		60mm		80mm		100mm	
	Ref	∅	Ref	∅	Ref	∅	Ref	∅
5	112429	8F	112430	8F	112431	8F	112432	8F
6	112434	8F	112435	8F	112436	8F	111715	9F
7	112439	8F	112440	8F	112441	8F	111720	9F
8	112444	8F	112445	8F	112446	8F	111725	9F
9	111727	9F	111728	9F	111729	9F	111730	9F

DELIVERY SYSTEM (LENGTH): 130CM

LATAM/ASIA

