PHILIPS

Arterial and Venous

Peripheral Vascular

See clearly. Treat optimally.

Only Philips offers the powerful combination of advanced imaging and specialized treatment options to accurately assess inside the vessel, successfully select the right treatment algorithm and optimize outcomes for your patients.

See clearly | Visualize the best path forward with IVUS

To determine the right treatment path for your patient, it is critical that you first accurately assess the disease state within the vessel. That's where the powerful combination of angiography and IVUS comes into play. Angiography provides a roadmap and flow characteristics of the vessel. IVUS brings you inside the vessel for deeper insights into vessel sizing and plaque morphology. Together, IVUS and angiography give you the advanced visualization you need to deliver exceptional patient care.



Accurately assess critical lesion characteristics with the four pillars of arterial IVUS

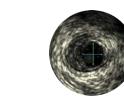
Only Philips provides a complete and simple plug-and-play digital array IVUS portfolio that includes both an 0.014" and 0.018" (RX and OTW) platform.

Vessel size

Guides device sizing to ensure precise wall apposition, drug delivery and placement



Vessel diameter



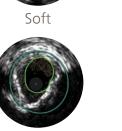


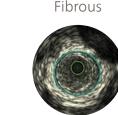
Lumen diameter



Plaque morphology

Understand plaque type and severity to help guide proper device selection



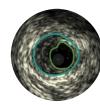


Thrombus

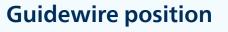
Plaque geometry

Visualize plaque burden location for precise treatment

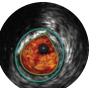


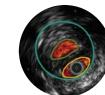


Eccentric



Confirm true lumen or sub-intimal guidewire location





True lumen



Sub-intimal



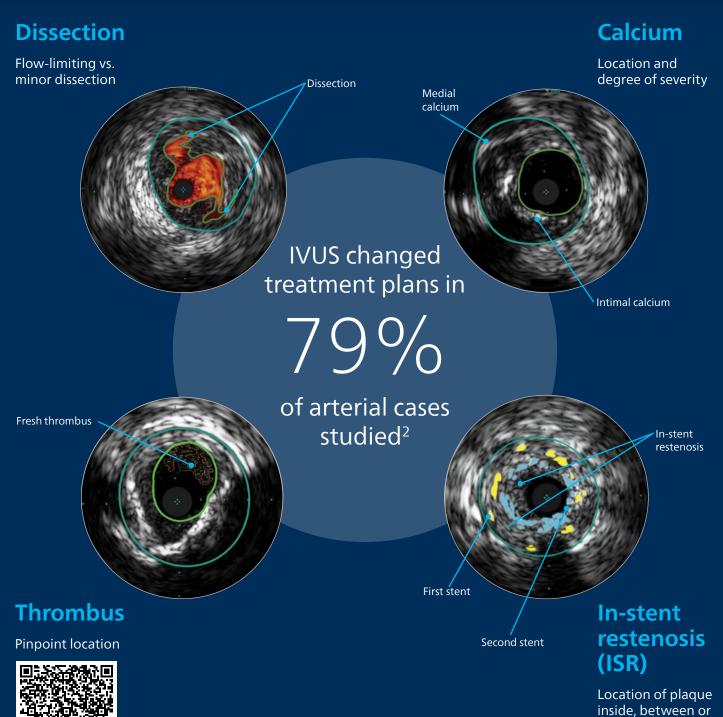


Calcific





"Angiography provides information on luminal characteristics of peripheral arteries, but severely underestimates the extent of atherosclerosis in patients with PAD, even in 'normal appearing' vessels."¹



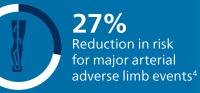
Angiogram alone is not enough

Is this a dissection, calcium, thrombus or stenosis?

Scan here to watch a video that shows how IVUS can help you see clearly and treat optimally. outside the stent

Demonstrated real-world results to improve patient outcomes

First ever global consensus guidance on IVUS³





28% Risk reduction for repeat venous intervention, nospitalization or death⁵

Venous

Accurately assess critical lesion characteristics with the four pillars of venous IVUS

Only Philips provides a simple plug-and-play digital array IVUS with the market leading 0.035"* IVUS platform.

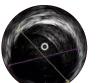
Vessel size

Guides device sizing to ensure precise wall apposition and stent placement

Thrombus type

Determine chronicity and

burden of clot⁶



Vessel

Acute

thrombus



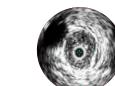
Vessel and lumen

Subacute

thrombus



Stent wall apposition



Chronic thrombus with fibrosis

Compression or stenosis

Determine degree of compression or stenosis

Post-thrombotic

lesion detection

*Venous use may not be available in select international markets



Stenosis

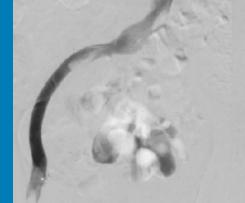


Severe stenosis with wall thickening





scaring



Venography alone is not enough

"IVUS applications in central venous pathologies are related to assessment and management of venous stenotic disease, thrombo-occlusive disease, IVC filter placement and retrieval."7

May-Thurner syndrome

Assess the degree of compression

Left commor iliac vein

Right commor iliac vein

Common iliac artery

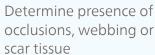
Post phlebitic stenosis

Common

femoral vein

Post thrombotic lesion detection

Assess and grade lesion severity





Compression

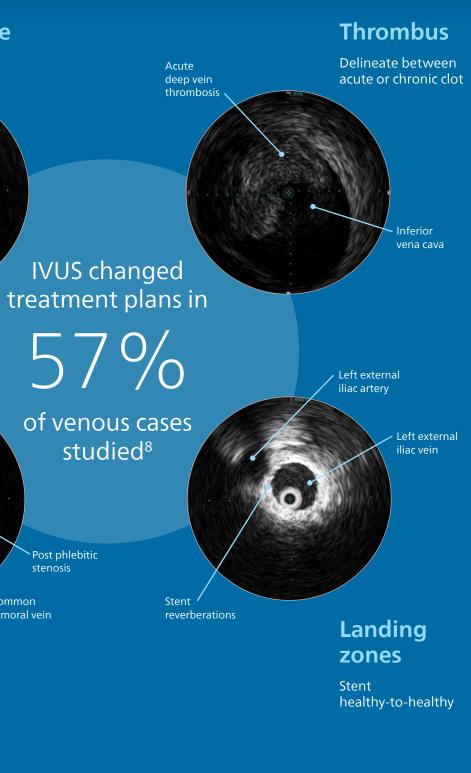
Webbing



Post thrombotic fibrosis occlusion







Treat optimally | Successfully select the right treatment algorithm

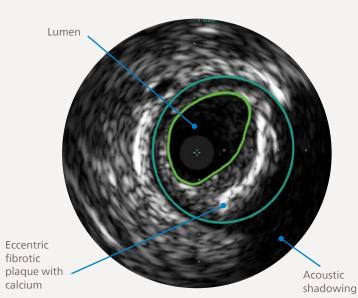
The Philips portfolio of therapeutic devices offers specialized treatment options for your arterial or venous interventions.

Successfully select Support Catheters **Quick-Cross, Quick-Cross Extreme and** Arterial **Quick-Cross Select Cross your** #1 selling support catheter toughest lesions **IVUS-guided Re-entry Catheter Pioneer Plus** Arterial The only IVUS-guided re-entry catheter Laser Atherectomy Catheters **Turbo-Elite and Turbo-Power** Arterial Strong performance in a variety of challenging lesion types above and below the knee⁹, and indicated for ISR* Mechanical Atherectomy System Phoenix Arterial Prepare and treat Front-cutting mechanical atherectomy for treating multiple lesions mixed morphologies with low risk of embolization¹⁰ including CTOs, ISR, calcium, neo-intimal hyperplasia, mixed Thrombectomy System morphologies and Arterial QuickClear thrombus Intuitive and powerful fresh thrombus aspiration Venous supports faster procedure times PTA Scoring Balloon Catheter AngioSculpt Arterial Reduces risk of flow-limiting dissections, including calcified lesions1 Drug-Coated Angioplasty Balloon Catheter **Stellarex Definitive treatment** for arterial lesions Durable treatment effect with a low-drug dose in common to complex patients¹² **Optimize outcomes Dissection Repair Solution Tack Endovascular System** Arterial Minimal metal dissection repair device for optimized PTA above- and below-the-knee Take patient care a step further IVC Filter Removal Laser Sheath CavaClear Venous The first and only FDA-cleared device designed for advanced IVC filter removal

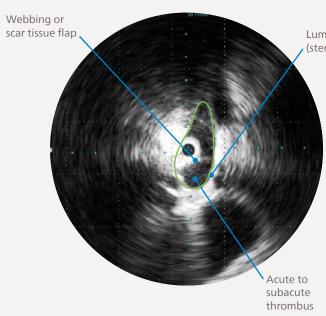
Full view treatment planning

The integration of visualization and interventional technologies allows physicians to see the complete picture and create a more informed and individualized procedural approach.

Arterial treatment plan example



Venous treatment plan example



Optimize outcomes for your patients with IVUS

No dissections | Reduce residual stenosis | Stent fully deployed | Treated entire lesion

See clearly.

Vessel size: 5.5 mm diameter Plaque morphology: Fibrotic plaque with intimal and medial calcium Plaque geometry: Eccentric lesion Guidewire position: True lumen

Treat optimally.

Ouick-Cross Catheter: Confidently cross challenging morphologies **Phoenix Deflecting Atherectomy System:** Front facing to cut, capture and clear mixed morphologies, including calcium Deflecting capabilities for larger luminal gain **Stellarex DCB:** Proven performance in calcium¹³ Tack Endovascular System: Optimizing PTA with precision dissection repair

Lumen (stenosed)

See clearly.

Vessel size: 10-13 mm **Thrombus type:** Acute to subacute thrombus Percentage compression or stenosis: 50-80% Other considerations: Occlusion webbing and collaterals

Treat optimally.

QuickClear Thrombectomy System: All-in-one, simple, single use aspiration pump and catheter Appropriate balloon venoplasty: Treat narrowed vessel by stretching it with air pressure Appropriate venous stent: Use stent to keep vein open

With vast IVUS experience and a consultative approach, Philips offers clinical support at every critical step—from integrating visualization and interventional technologies to focusing on solutions to help you provide individualized patient treatment plans.



Important safety information

Stellarex .035 Drug-Coated Balloon

The Stellarex 0.035" OTW Drug-coated Angioplasty Balloon is indicated for percutaneous transluminal angioplasty (PTA), after appropriate vessel preparation, of de novo, restenotic, or in-stent restenotic lesions up to 180 mm in length in native superficial femoral or popliteal arteries with reference vessel diameters of 4-6 mm. The Stellarex™ 0.035" OTW Drug-coated Angioplasty Balloon is contraindicated for use in:

- Patients with known hypersensitivity to paclitaxel or structurally re-lated compounds.
- Patients who cannot receive recommended antiplatelet and/or anticoagulation therapy.
- Women who are breastfeeding, pregnant or are intending to become pregnant or men intending to father children.
- Coronary arteries, renal arteries, and supra-aortic/cerebrovascular arteries
- Patients judged to have a lesion that prevents complete inflation of an angioplasty balloon or proper placement of the delivery system

Possible adverse effects associated with the balloon dilation procedure include, but are not limited to: Abrupt vessel closure; Allergic reaction to contrast medium, antiplatelet therapy, or catheter system components (drug, excipients, and materials); Amputation/ Loss of limb; Arrhythmias; Arterial aneurysm; Thrombosis; Arteriovenous fistula (AVF); Bleeding; Death; Embolism/Device embolism; Fever; Hematoma; Hemorrhage; Hypertension/Hypotension; Infection or pain at insertion site; Inflammation; Ischemia or infarction of tissue/organ; Occlusion; Pain or tenderness; Peripheral edema; Pseudoaneurysm; Renal insufficiency or failure; Restenosis; Sepsis or systemic infection; Shock; Stroke/Cerebrovascular accident; Vessel dissection, perforation, rupture, spasm, or recoil; Vessel trauma which requires surgical repair; Balloon rupture; Detachment of a component of the balloon and/or catheter system; Failure of the balloon to perform as intended; Failure to cross the lesion.

Additional complications which may be associated with the addition of paclitaxel to the balloon include, but may not be limited to the following: Allergic/immunologic reaction to paclitaxel; Alopecia; Anemia; Gastrointestinal symptoms (diarrhea,

nausea, pain, vomiting); Hematologic dyscrasia (including neutropenia, leucopenia, thrombocytopenia); Hepatic enzyme changes; Histologic changes in vessel wall including inflammation, cellular damage, or necrosis; Myalgia/Arthralgia; Myelosuppression; Peripheral neuropathy.

Caution: Federal law restricts this device to sale by or on the order of a physician.

Tack Endovascular System

The Tack Endovascular System (6F, 3.5-6.0 mm US/ 2.5-6.0 mm EU and 4.0-8.0 mm) is intended for use in the superficial femoral and proximal popliteal arteries ranging in diameter from 3.5 mm to 6.0 mm US/ 2.5 to 6.0 mm EU and 4.0 mm to 8.0 mm for the repair of post percutaneous transluminal balloon angioplasty (PTA) dissection(s).

The Tack Endovascular System (4F, 1.5-4.5 mm) is intended for use in mid/distal popliteal, tibial, and peroneal arteries, ranging in diameter from 1.5 mm to 4.5 mm, for the treatment of post percutaneous transluminal balloon angioplasty (PTA) dissection(s).

The Tack Endovascular System is contraindicated for the following:

- 1. Patients with residual stenosis in the treated segment equal to or greater than 30% after PTA.
- 2. Tortuous vascular anatomy significant enough to prevent safe introduction and passage of the device.
- 3. Patients with a known hypersensitivity to nickel titanium alloy (Nitinol).
- 4. Patients unable to receive standard medication used for interventional procedures such as anticoagulants, contrast agents and antiplatelet therapy.

Prior to using the Tack Endovascular System, please review the instructions for use for a complete listing of indications, contraindications, warnings, precautions, potential adverse events, and directions for use.

Tack Endovascular System is CE Mark authorized under EC Directive 93/42/EEC. Caution: Federal law restricts this device to sale by or on the order of a physician.

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