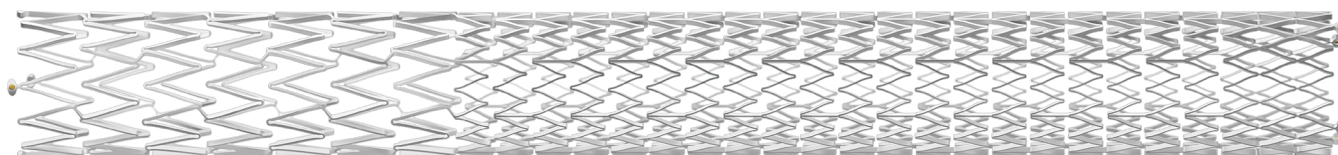


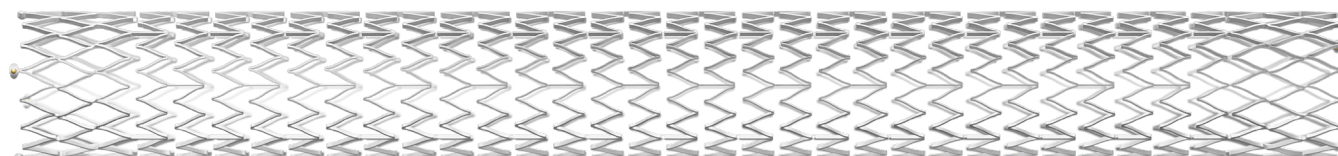
Product specifications

Engineered for greater crush resistance and flexibility, Philips Duo Venous Stent System is built to address the multiple anatomical challenges of the deep venous system and includes two stents – Duo Hybrid and Duo Extend. Available in a wide range of sizes, the option to combine Duo Hybrid and Duo Extend enables versatile treatment solutions based on disease length and severity, without the need to mix and match stent brands.¹



Duo Hybrid

	Lengths (mm)					
Diam (mm)	60	80	100	120	140	160
12	9 Fr	9 Fr	9 Fr	9 Fr	9 Fr	9 Fr
14	9 Fr	9 Fr	9 Fr	9 Fr	9 Fr	9 Fr
16	10 Fr	10 Fr	10 Fr	10 Fr	10 Fr	10 Fr
18	10 Fr	10 Fr	10 Fr	10 Fr	10 Fr	10 Fr

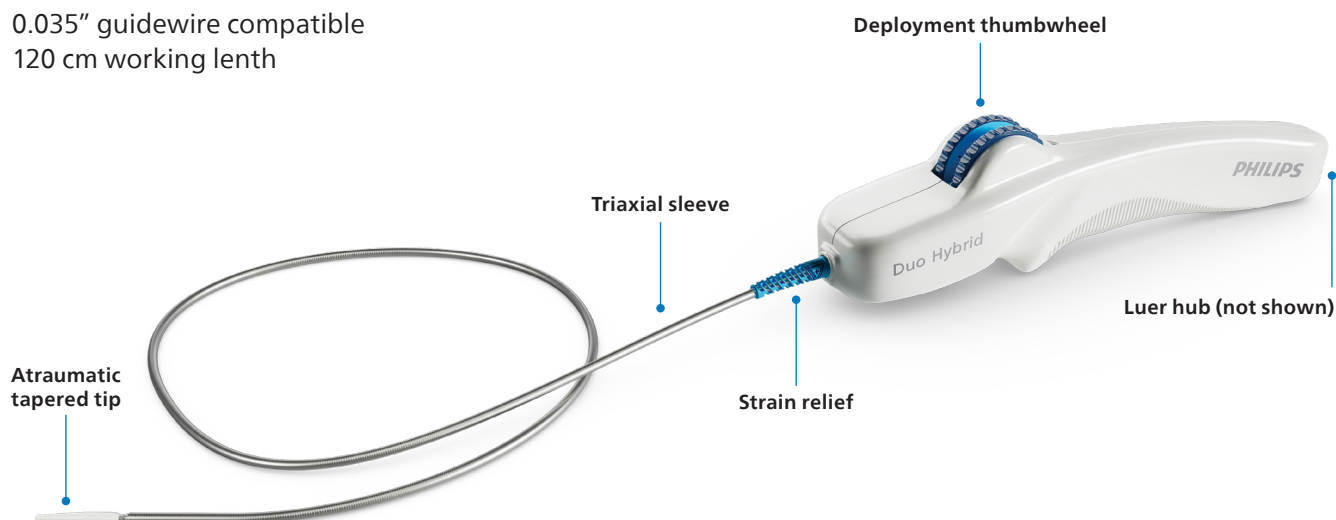


Duo Extend

	Lengths (mm)					
Diam (mm)	40	60	80	100	120	140
12	9 Fr	9 Fr	9 Fr	9 Fr	9 Fr	9 Fr
14	9 Fr	9 Fr	9 Fr	9 Fr	9 Fr	9 Fr
16	10 Fr	10 Fr	10 Fr	10 Fr	10 Fr	10 Fr

Triaxial, OTW delivery catheter

0.035" guidewire compatible
120 cm working length



1. Data on file: D062749

Brief statement

Philips Duo Venous Stent System

Intended use:

The Duo Venous Stent System is intended for use in the iliofemoral veins for the treatment of symptomatic venous outflow obstruction. The Duo Hybrid Stent is intended to be used in the iliac vein at the confluence of the inferior vena cava only. The Duo Extend Stent is intended for use in the common iliac and common femoral vein.

Contraindications for use:

The Duo Venous Stent System is contraindicated for the following: 1. Patients with a known hypersensitivity to nickel-titanium alloy (Nitinol). 2. Patients unable to receive standard medication used for interventional procedures including anticoagulants, contrast agents and antiplatelet therapy. 3. Patients who are judged to have a lesion that prevents complete inflation of a balloon dilation catheter or proper placement of the stent or the stent delivery system. 4. Tortuous vascular anatomy significant enough to prevent safe introduction and passage of the device to its intended location. 5. Duo Hybrid jugular or contralateral vascular access.

Prior to implanting the Duo Venous Stent System, please review the instructions for use for a complete listing of indications, contraindications, warnings, precautions, potential adverse events and directions for use.

Caution:

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



©2024 Koninklijke Philips N.V. All rights reserved.
Approved for external distribution.
D062728-01 022024

Philips
3721 Valley Centre Drive, Suite 500
San Diego, CA 92130 USA
www.philips.com/IGTdevices