

**Abbott**

2 p.d Kraujagyslių uždarymo po kateterizacijos sistema

Perclose™ ProStyle™ Suture-Mediated Closure and Repair (SMCR) System Tender Specification

Kraujagyslių uždaryma su silo ir mazgo pagalba.

INDICATIONS FOR USE

The Perclose™ ProStyle™ SMCR System is indicated for the percutaneous delivery of suture for closing the common femoral artery and vein access site of patients who have undergone diagnostic or interventional catheterization procedures.

- For access sites in the common femoral artery using 5F to 21F sheaths. For sheath sizes greater than 8F, at least two devices and the pre-close technique are required.
- For access sites in the common femoral vein using 5F to 24F sheaths. For sheath sizes greater than 8F, at least one device and the pre-close technique is required.

Skirta 5 – 24 F diametro punkcijos vietai užsiti.

COMPONENTS	MATERIAL SPECIFICATIONS	DIMENSION SPECIFICATIONS
Perclose™ ProStyle™ Device		
Sistem sudaro polipropileno silas ir 2 adatos.		
Suture	Monofilament Polypropylene	Diameter 0.0088 ± 0.0010” Length 20.0” minimum
Needles	300 Series Stainless Steel	Diameter 0.021 ± 0.0015”
Sheath	Polyether Block Amide	Diameter 0.113”; Length 8.8 ± 0.5”
Handle	Polycarbonate	N/A
Suture Cutter	400 Series Stainless Steel	N/A
Marker Lumen	Pebax	N/A
Needle Guide / Foot	PPA	N/A
Lever / Plunger	ABS	N/A
Perclose™ ProStyle™ Suture Trimmer		
Handle	ABS	N/A
Sheath / Trimmer	304 Stainless Steel	N/A
Tip	PAA IXEF	N/A
Perclose™ Snared Knot Pusher		
Body	Nylon 66/Carbon Fiber	N/A
Snare	304V Stainless Steel Wire	N/A
Snare Tab	Polycarbonate	N/A

TECHNICAL SPECIFICATIONS

- Method of Sterilization: Ethylene Oxide
- Shelf Life: 2 years
- Storage: Store in a cool dry location

CAUTION: This product is intended for use by or under the direction of a physician. Prior to use, reference the Instructions for Use, inside the product carton (when available) or at www.vascular.eifu.abbott or at medical.abbott/manuals for more detailed information on Indications, Contraindications, Warnings, Precautions and Adverse Events.

Information contained herein for DISTRIBUTION outside the U.S. only. Check the regulatory status of the device in areas where CE marking is not the regulation in force.

Data of File at Abbott

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