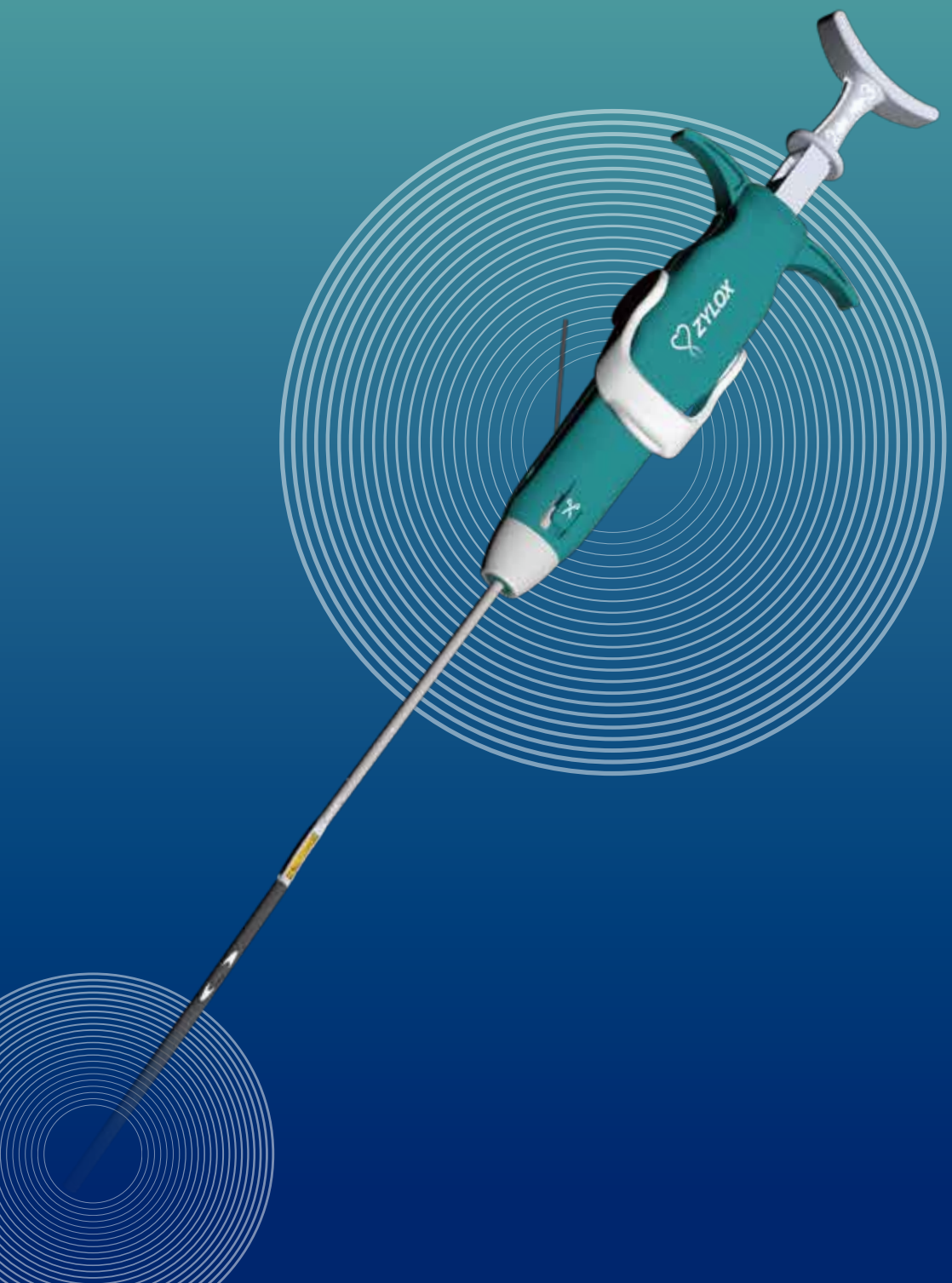
ZYLOX TONBRIDGE

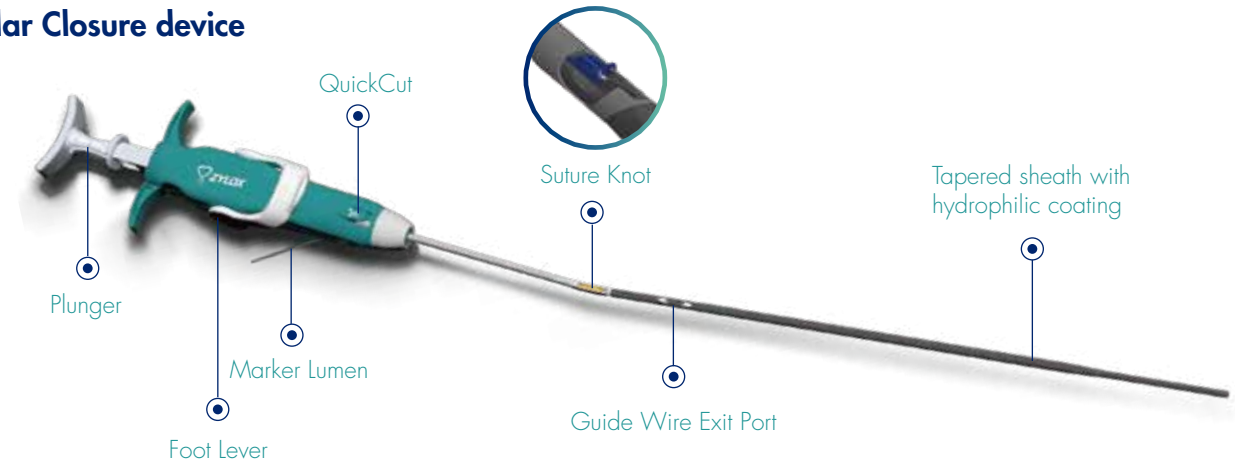
ZYLOX Unicorn™

Vascular Closure System

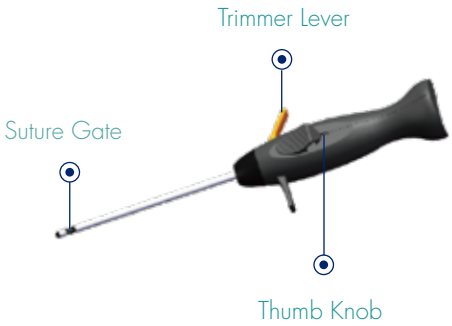


ZYLOX Unicorn

Vascular Closure device



Vascular Closure Suture Trimmer



Clinically Validated

Primary Endpoint

Femoral Artery Access Site Vascular Complication at 30 days	ZYLOX Unicorn N = 114	ProGlide N = 115	p-value
[95% Confidence Interval]	1.75% [0.00%, 4.16%]	0.87% [0.00%, 2.57%]	P>0.05

A prospective, randomized, concurrently-controlled, and non-inferiority clinical trial has been carried out in 4 research centers in China. The ZYLOX Unicorn Vascular Closure System is non-inferior to the Perclose ProGlide Suture-Mediated Closure System on the safety and effectiveness of femoral artery closure up to 22F sheath size.

Ordering Information

The ZYLOX Unicorn™ Vascular Closure System includes:
(1) Vascular Closure device
(2) Vascular Closure Suture Trimmer

Specification	Compatible guidewire	Sheath sizes
SV1401	0.038 inch (or smaller)	5-22F

Indications: The Vascular Closure System is indicated for the percutaneous delivery of suture for closing the common femoral artery access site of patients who have undergone diagnostic or interventional catheterization procedures using 5F to 22F sheaths. For sheath sizes greater than 8F, at least two devices and the pre-close technique are required.

The ZYLOX Unicorn™ Vascular Closure System is not available in markets requiring FDA clearance and CE certification.

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