



Edwards

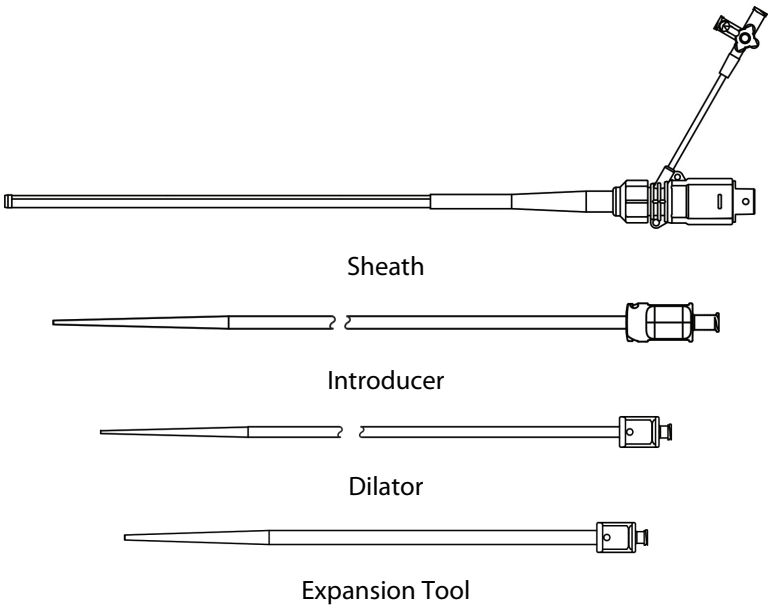
Edwards eSheath+ Introducer Set

Instructions for Use

This product should be used by physicians trained and experienced in interventional techniques.

1.0 Device Description

The Edwards eSheath+ introducer set contains one introducer, one dilator, one expansion tool and a sheath, as well as a loader packaged with an Edwards delivery system. The working lengths of the introducer, dilator, and sheath contain a hydrophilic coating.



Model	Sheath I.D.	Sheath Working Length	Compatible THV	Introducer O.D.	Introducer Working Length	Dilator O.D.	Dilator Working Length
914ESP	14 F	36 cm	20 mm 23 mm 26 mm	14 F	57 cm	16 F	38 cm
916ESP	16 F	36 cm	29 mm	16 F	57 cm	18 F	38 cm

2.0 Indications

The Edwards eSheath+ introducer set is indicated for the introduction and removal of devices used with the Edwards transcatheter heart valves.

Edwards, Edwards Lifesciences, the stylized E logo, Edwards eSheath, Edwards eSheath+, eSheath, and eSheath+ are trademarks of Edwards Lifesciences Corporation. All other trademarks are the property of their respective owners.

3.0 Contraindications

There are no known contraindications.

4.0 Warnings

The devices are designed, intended, and distributed for single use only.

Do not resterilize or reuse the devices. There is no data to support the sterility, nonpyrogenicity, and functionality of the devices after reprocessing.

The Edwards eSheath+ introducer set must be used with a compatible 0.035" (0.89 mm) guidewire to prevent vessel injury.

5.0 Precautions

- Caution should be used in vessels that have diameters less than 5.5 mm or 6 mm as it may preclude safe placement of the 14F and 16F Edwards eSheath+ introducer set respectively.
- Use caution in tortuous or calcified vessels that would prevent safe entry of the introducer set.
- Do not use the Edwards eSheath+ introducer set if the packaging sterile barriers and any components have been opened or damaged.
- When inserting, manipulating or withdrawing a device through the sheath, always maintain sheath position.
- When puncturing, suturing or incising the tissue near the sheath, use caution to avoid damage to the sheath.
- Expansion tool does not contain a hydrophilic coating. Do not use as a dilator.

6.0 Potential Adverse Events

Complications associated with standard catheterization and use of angiography include, but are not limited to, injury including perforation or dissection of vessels, thrombosis and/or plaque dislodgement which may result in emboli formation, distal vessel obstruction, hemorrhage, infection, and/or death.

7.0 Directions for Use

1. Inspect the length of the introducer, dilator, expansion tool and sheath for surface defects and damage prior to clinical use.
2. Flush the introducer and dilator using heparinized saline through the guidewire lumen.
3. Hydrate the length of the introducer, dilator, and sheath with heparinized saline to activate the hydrophilic coating. Wet the surface of the expansion tool.
4. Flush the sheath using heparinized saline through the flush port; close the flush port.
5. Use the expansion tool to pre-expand the partially expandable portion of the sheath prior to procedural use.

Note: After pre-expanding the sheath, inspect the length of the expandable portion for damage prior to use.

6. After removing the expansion tool, flush the sheath a second time using heparinized saline through the flush port; close the flush port.
7. Insert the introducer completely into the sheath and turn clockwise to lock the introducer hub to the sheath hub.
8. Using standard catheterization techniques, gain access to the vessel and dilate as necessary with the dilator to accommodate the sheath.
9. Orient the sheath appropriately and maintain orientation for the duration of the procedure. Insert the sheath assembly using standard technique while following its progression on fluoroscopy.

Note: The proximal tapered end of the sheath working length is larger in diameter.

10. If possible, suture the sheath into place.
11. Remove the introducer from the sheath by turning counterclockwise to unlock the introducer hub from the sheath.
12. Insert the device into the sheath (reference device specific instructions for use).
 - a. When using the loader, flush the loader with heparinized saline and insert the device into the loader.
 - b. Insert the loader into the sheath until the loader stops. Advance device through the loader and into the sheath.
 - c. Retract the loader once the device is passed through the sheath. If additional working length is needed unscrew the cap from the loader and peel the loader from the shaft of the device.
13. After the completion of the procedure, remove the device from the sheath.

Note: The sheath should be intermittently flushed with heparinized saline per standard technique.

14. Remove the suture (if necessary) and remove the sheath entirely without torquing. Do not reinsert the sheath.

8.0 How Supplied

The Edwards eSheath+ introducer set is supplied pouched and sterilized by ethylene oxide.








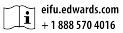











9.0 Storage



















The Edwards eSheath+ introducer set should be stored in a cool, dry place.


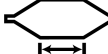









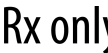




10.0 Device Disposal

Used sheath sets may be handled and disposed of in the same manner as hospital waste and biohazardous materials. There are no special risks related to the disposal of these devices.

Symbol Legend

	English
	Reorder Number
	Model Number
	Usable length
	Do not re-use
	Lot Number
	Caution
	Consult instructions for use
	Consult instructions for use on the website
	Do not use if package is damaged and consult instructions for use
	Exterior diameter
	Inner diameter
	Store in a cool, dry place
	Keep dry
	Keep away from sunlight
	Unique Device Identifier
	Temperature limit
	Sterile
	Sterilized using ethylene oxide
	Sterilized using irradiation

	English
	Do not re-sterilize
	eSheath compatibility
	eSheath compatibility
	Single sterile barrier system
	Single sterile barrier system with protective packaging inside
	Quantity
	Use-by date
	Serial Number
	Manufacturer
	Date of manufacture
	Authorized representative in the European Community/European Union
	Guidewire compatibility
	Nominal Pressure
	Rated burst pressure
	Recommended guidewire length
	Minimum sheath size
	Catheter shaft size
	Importer

	English
	Balloon diameter
	Balloon working length
	For use with size 20 mm Edwards transcatheter heart valve
	For use with size 23 mm Edwards transcatheter heart valve
	For use with size 26 mm Edwards transcatheter heart valve
	For use with size 29 mm Edwards transcatheter heart valve
	MR Conditional
	Contents
	Non-pyrogenic
	Medical device
	Contains biological material of animal origin
	Caution: Federal (USA) law restricts this device to sale by or on the order of a physician.
	Time & Temperature Sensitive
	Contains hazardous substances
	Size
	Work Order

Note: Not all symbols may be included in the labeling of this product.



Edwards

2023-06
10058027001 A

© Copyright 2023, Edwards Lifesciences LLC
All rights reserved.

Manufacturer 
Edwards Lifesciences LLC
One Edwards Way
Irvine, CA 92614 USA

Telephone +1.949.250.2500
+1.800.424.3278
FAX +1.949.250.2525

Web IFU