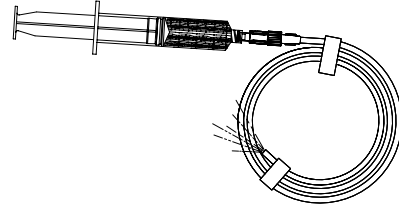


INSTRUCTIONS FOR PREFLUSHING THE CATHETER AND STYLET:

1. Flush the catheter with heparinized saline solution or sterile normal saline. **Note:** The catheter may be trimmed if a shorter length is required.
2. Attach a syringe with heparinized saline solution or sterile normal saline to the Luer Lock fitting of the flush through stylet hub.
3. Inject enough solution to wet the stylet surface entirely.
4. Remove the stylet from its holder and insert it into the PICC. If the catheter has been trimmed, only advance the stylet to the distal end of the catheter. **Note:** If the surface of the stylet becomes dry after removal from the holder, wet with additional heparinized saline or sterile normal saline.
5. The catheter stylet assembly can now be introduced. (See device Instructions For Use for further information.)

WARNING: Do not cut stylet.
WARNING: Hydrate stylet prior to use.



INTENDED USE:

The Flushable Guidewire Stylet is a coiled guidewire used to stiffen the catheter during insertion to aid the advancement of the lumen into the vasculature.

POTENTIAL COMPLICATIONS:

Cardiac Events, Embolisms, Infections, Perforation or trauma to vessel, Bleeding

WARNING AND PRECAUTIONS:

- Caution when using this device. Be aware of sharps.
- Do not use if components are damaged, deformed or missing.
- Do not proceed if resistance is felt or interaction between components is failing.
- Do not insert or withdraw the guidewire stylet forcibly from any components. The guidewire may break or unravel. If the guidewire becomes damaged, the introducer needle or sheath/dilator and guidewire must be removed together.
- The guidewire stylets are a single use device. Do not re-use or attempt to re-sterilize.

paper size 6.5" x 4"

CAUTIONS:

- Federal Law (USA) restricts this device to the sale and use by or on the order of a physician.
- Disposable: for single patient use only. Re-use may lead to infection or illness/injury. Do not resterilize.
- Do not use if package is damaged.
- Ethylene oxide (EtO) sterilized; non-pyrogenic.
- Carefully read all instructions prior to use. Observe all warnings and cautions.



STERILE EO

INSTRUCTIONS FOR USE:

- Examine the device after it is removed from the patient to ensure no foreign material remains inside the patient.
- Discard biohazard according to facility protocol.

ADDITIONAL STATEMENT:

Medcomp® does not recommend a particular technique for the use of this device. The physician should evaluate the appropriateness according to individual patient conditions and his or her medical training and experience.

WARRANTY

Medcomp® WARRANTS THAT THIS PRODUCT WAS MANUFACTURED ACCORDING TO APPLICABLE STANDARDS AND SPECIFICATIONS. PATIENT CONDITION, CLINICAL TREATMENT, AND PRODUCT MAINTENANCE MAY AFFECT THE PERFORMANCE OF THIS PRODUCT. USE OF THIS PRODUCT SHOULD BE IN ACCORDANCE WITH THE INSTRUCTIONS PROVIDED AND AS DIRECTED BY THE PRESCRIBING PHYSICIAN.

Because of continuing product improvement, prices, specifications, and model availability are subject to change without notice. Medcomp® reserves the right to modify its products or contents without notice.

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5.1.9 	5.4.2 	5.2.6
5.2.3 STERILE EO	Sterilized Using Ethylene Oxide	

* This symbol is in accordance with ISO 15223-1.
 *** FDA guidance Use of Symbols in Labeling.

paper size 6.5" x 4"