BARBED LUER LOCK ADAPTOR

INTENDED USE: Connection point for catheter and male luer for peritoneal dialysis catheter.

WARNINGS:

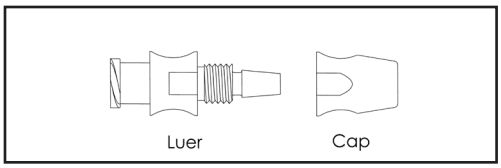
- Federal Law (USA) restricts the device to sale by or on the order of a physician.
- This device is for Single Use Only.
- Do not resterilize by any method.
- Re-use may lead to infection or illness/injury.
- The manufacturer shall not be liable for any damages caused by re-use or resterilization of the device.
- Contents sterile and non-pyrogenic in unopened, undamaged package. STERILIZED BY ETHYLENE OXIDE
- Do not use if package is opened or damaged.
- Do not use if any sign of product damage is visible.

STERILE	EO

Caution: When cutting catheter to desired length, assure lumen is cut square and that remaining catheter lumen is not damaged.

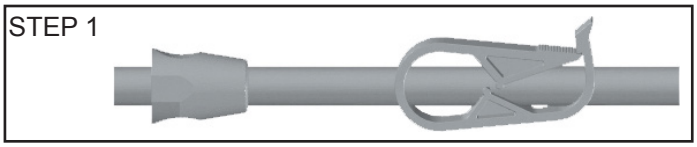
INSTRUCTIONS FOR USE:

1. Remove the luer and cap from the package. Note: Both components must be utilized to complete installation.

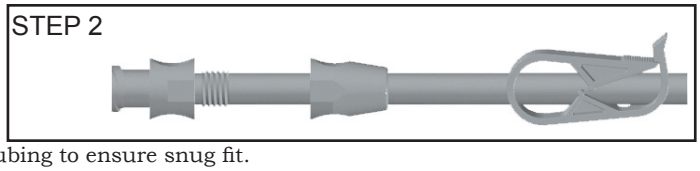


2. Assemble the provided luer to the catheter as follows:

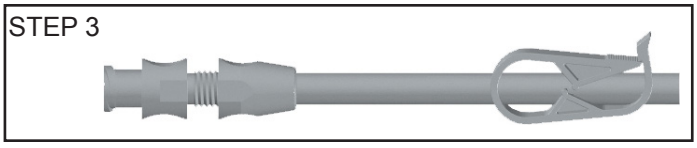
- Slide cap over the catheter tubing such that the taper faces the catheter exit site. (Step 1)



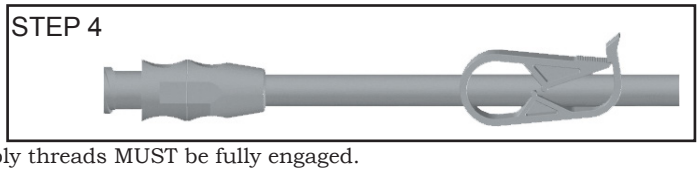
- Push barbed end of the luer into the catheter tubing. Position completely over barbed end of fitting making sure the tubing is FULLY seated. (Step 2)



- Gently tug the tubing to ensure snug fit.
- Slide the cap toward the threads on the luer, and rotate it to engage the threads. (Step 3)



- Continue to thread the cap by hand onto the luer until secure. (Step 4)



Caution: Assembly threads MUST be fully engaged.

3. Grasping the luer in one hand, and the tubing in the other, gently tug on the joint to test the security of the connector. If the luer pulls out of tubing, repeat the connection procedure. A connection failure may be due to one, or a combination of the following:

    - The luer is not fully inserted into the catheter tubing.
    - The catheter tubing is damaged, preventing a secure connection. If the failure is due to damaged tubing, then the catheter may need to be removed and replaced. CAUTION: Be sure to pull on the external tubing and the connector only and not on the catheter In Situ.

4. Attach end cap.

5. Use a sterile 10cc luer lock syringe to aspirate any air introduced during the connection procedure.

6. Continue with catheter Instructions for Use.

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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SYMBOL TABLE

5.1.1	Manufacturer*
5.3.4	Keep Dry*
5.4.2	Do Not Re-use*
5.6.3	Non-pyrogenic*
5.3.2	Keep Away from Sunlight*
5.2.3 STERILE EO	Sterilized Using Ethylene Oxide*
5.2.8	Do Not Use if Package is Damaged*
5.1.4	Use-by Date*
5.2.6	Do Not Resterilize*
5.1.5 LOT	Batch/Lot Number*
5.1.6 REF	Catalogue Number*
5.1.2 EC REP	Authorized Representative in the European Community*
5.4.4	Caution, consult Accompanying Documents*
Rx Only	Prescription Use Only***
MR	MR Conditional - 3 Tesla ****

***This symbol is in accordance with ISO 15223-1.**

***** FDA guidance Use of Symbols in Labeling.**

******This Symbol is in accordance with ASTM F 2503-13**

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