

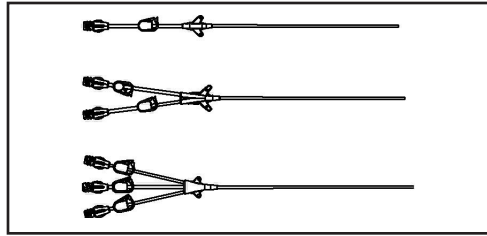
medCOMP®

Pro-PICC® CT PERIPHERALLY
INSERTED CENTRAL VEIN ACCESS
CATHETER

INSTRUCTIONS FOR USE

DESCRIPTION:

- A family of peripherally inserted central catheters made from specially formulated biocompatible medical grade materials. Catheters are packaged in a tray with accessories necessary for a percutaneous microintroducer introduction (Modified Seldinger or Seldinger technique).



INDICATIONS FOR USE:

- The Pro-PICC® CT catheter is indicated for short or long term peripheral access to the central venous system for intravenous therapy and power injection of contrast media, and allows for central venous pressure monitoring. For blood sampling, infusion, or therapies, use a 4F or larger catheter. The maximum recommended infusion rate varies by catheter French size and is printed on the catheter.

IMPORTANT INFORMATION PERTAINING TO POWER INJECTION:

- Contrast media should be warmed to body temperature prior to power injection. **Warning:** Failure to warm contrast to body temperature prior to power injection may result in catheter failure.
- Vigorously flush the Pro-PICC® CT catheter using a 10cc or larger syringe and sterile normal saline prior to and immediately following the completion of power injection studies. This will ensure the patency of the catheter and prevent damage to the catheter. Resistance to flushing may indicate partial or complete catheter occlusion. **Do not** proceed with power injection study until occlusion has been cleared. **Warning:** Failure to ensure patency of the catheter prior to power injection studies may result in catheter failure.
- Do not** exceed the maximum flow rate printed on the catheter. **Warning:** Power injector machine pressure limiting feature may not prevent over pressurization of an occluded catheter. **Warning:** Exceeding the maximum indicated flow rate may result in catheter failure and/or catheter tip displacement.

- Warning:** Pro-PICC® CT catheter indication of power injection of contrast media implies the catheter's ability to withstand the procedure, but does not imply appropriateness of the procedure for a particular patient. A suitably trained clinician is responsible for evaluating the health status of a patient as it pertains to a power injection procedure.

CONTRAINDICATIONS:

- The presence of device related infection, bacteremia, or septicemia is known or suspected.
- The patient's body size is insufficient to accommodate the size of the implanted device.
- The patient is known or is suspected to be allergic to materials contained in the device.
- There has been past irradiation of prospective insertion site.
- There have been previous episodes of venous thrombosis or vascular surgical procedures at the prospective placement site.
- There are local tissue factors that may prevent proper device stabilization and/or access.

POSSIBLE COMPLICATIONS:

- Air Embolus
- Bleeding
- Brachial Plexus Injury
- Cardiac Arrhythmia
- Cardiac Tamponade
- Catheter Erosion Through the Skin
- Catheter Embolism
- Catheter Occlusion
- Catheter Related Sepsis
- Endocarditis
- Exit Site Infection
- Exit Site Necrosis
- Exsanguination
- Fibrin Sheath Formation
- Hematoma
- Intolerance Reaction to Implanted Device
- Laceration of the Vessels or Viscus
- Myocardial Erosion
- Perforation of Vessels or Viscus
- Pleuritis
- Spontaneous Catheter Tip Malposition or Retraction
- Thromboembolism
- Venous Thrombosis
- Ventricular Thrombosis
- Vessel Erosion
- Risks normally associated with local or general anesthesia, surgery, and post-operative recovery
- Before attempting the insertion, ensure that you are familiar with the potential complications and their emergency treatment should any of them occur.


WARNINGS:


- In the rare event that a hub or connector separates from any component during insertion or use, take all necessary steps and precautions to prevent blood loss or air embolism and remove the catheter.

- Do not advance the guidewire or catheter if unusual resistance is encountered.

- Do not insert or withdraw the guidewire forcibly from any component. The wire may break or unravel. If the guidewire becomes damaged, the introducer needle or sheath/dilator and guidewire must be removed together.

- Federal Law (USA) restricts this device to sale by or on the order of a physician.

- This catheter is for Single Use Only. 


- Do not resterilize the catheter or accessories 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 this catheter or accessories.

- Contents sterile and non-pyrogenic in unopened, undamaged package. STERILIZED BY ETHYLENE OXIDE

STERILE EO

- Do not use catheter or accessories if package is opened or damaged. 

- Do not use catheter or accessories if any sign of product damage is visible.

CATHETER PRECAUTIONS:

- Small syringes will generate excessive pressure and may damage the catheter. Ten (10)cc or larger syringes are recommended.

- Do not use sharp instruments near the extension lines or catheter lumen.

- Do not use scissors to remove dressing.

- Catheter will be damaged if clamps other than what is provided with this kit are used.

- Clamping of the tubing repeatedly in the same location will weaken tubing. Avoid clamping near the luer(s) and hub of the catheter.

- Examine catheter lumen and extension(s) before and after each infusion for damage.

- To prevent accidents, assure the security of all caps and connections prior to and between uses.

- Use only Luer Lock (threaded) Connectors with this catheter.

- Repeated overtightening of luer lock connections, syringes, and caps will reduce connector life and could lead to potential connector failure.

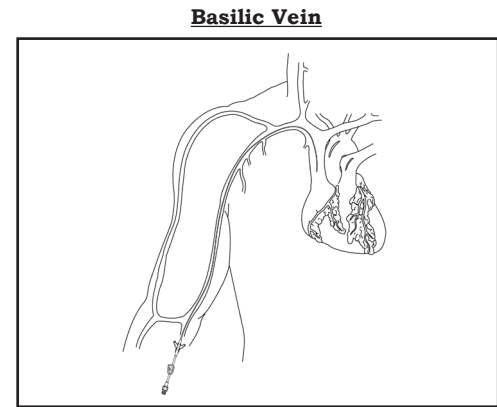
- Confirm catheter tip position by x-ray prior to use. Monitor tip placement routinely per institution policy.

- The catheter allows for blood draws, power injection of contrast media, intravenous therapy, and infusion of medications into the central venous system. Refer to standards of practice and institutional policies for compatible infusion agents for central venous access.

- Follow all contraindications, warnings, precautions, and instructions for all infusates including contrast media as specified by their manufacture.

INSERTION SITES:

- The basilic, median cubital, or cephalic vein may be catheterized. The basilic vein above antecubital fossa is the preferred site.



DIRECTIONS FOR MODIFIED SELDINGER INSERTION

- Read instructions carefully before using this device. The catheter should be inserted, manipulated, and removed by a qualified, licensed physician or other qualified health care professional under the direction of a physician.

- The medical techniques and procedures described in these instructions for use do not represent all medically acceptable protocols, nor are they intended as a substitute for the physician's experience and judgment in treating any specific patient.

- Use standard hospital protocols when applicable.

PRIOR TO PLACEMENT:

Identify Insertion Site and Vein, taking into account the following variables:

- patient diagnosis
- age and size of patient
- unusual anatomical variables
- type and purpose of IV therapy
- anticipated dwell time of catheter

- Apply tourniquet to arm above anticipated insertion site.
- Select vein based on assessment.
- Release tourniquet.

PREPARE CATHETER:

- Preflush catheter, sideport adaptor, and needleless access port(s).

- Attach saline filled syringe to luer of sideport adaptor and flush adaptor and catheter. Clamp sideport extension and remove syringe. If using multiple lumen catheter, attach needleless access port to remaining extension (s). Attach saline filled syringe to the needleless access port and completely flush catheter lumen. Remove syringe from access port prior to clamping extension.

Caution: Never close clamp on catheter stylet; stylet and catheter damage may result.

Caution: The needleless access port should not be used with needles, blunt cannula, or other non-luer connectors, or luer connectors with visible defects. If needle access is attempted, the needleless access port must be replaced immediately. Do not exceed 100 actuations.

INSERTION:

- Strict aseptic technique must be used during insertion, maintenance, and catheter removal procedures. Provide a sterile operative field. Use sterile drapes, instruments, and accessories. Perform surgical scrub. Wear gown, cap, gloves, and mask.

- Apply tourniquet to arm above anticipated insertion site to distend the vein.

- Insert the introducer needle with attached syringe into the target vein. Aspirate to insure proper placement. Release tourniquet.

- Remove the syringe and place thumb over the end of the needle to prevent blood loss or air embolism. Draw the flexible end of marked .018" guidewire back into advancer so that only the end of the guidewire is visible. Insert the advancer's distal end into the needle hub. Advance guidewire with forward motion into and past the needle hub into the target vein.

Caution: Guidewire is hydrophilic. Hydrate prior to use.

Caution: The length of the wire inserted is determined by the size of the patient. Monitor patient for arrhythmia throughout this procedure. The patient should be placed on a cardiac monitor during this procedure. Cardiac arrhythmias may result if guidewire is allowed to pass into the right atrium. The guidewire should be held securely during this procedure.

Note: For alternate insertion method, see Directions for Seldinger Insertion Section.

- Remove needle, leaving guidewire in the target vein. Thread sheath/dilator over the proximal end of the guidewire into target vein. Advance the guidewire until it reaches the caval atrial junction. Once the guidewire is in place, measure the depth of the guidewire by reading the markings on the wire. Remove the guidewire leaving the sheath and dilator

in the vein.

Caution: DO NOT bend the sheath/dilator during insertion as bending will cause the sheath to prematurely tear. Hold sheath/dilator close to the tip (approximately 3cm from tip) when initially inserting through the skin surface. To progress the sheath/dilator towards the vein, regrab the sheath/, dilator a few centimeters (approximately 5cm) above the original grasp location and push down on the sheath/dilator. Repeat procedure until sheath/ dilator is fully inserted.

Caution: Never leave sheath in place as an indwelling catheter. Damage to the vein will occur.

- Loosen locking collar of sideport and withdraw stylet back beyond the point where the catheter is to be trimmed by at least ¼ inch (1cm). Cut catheter to length determined by marked guidewire.

Caution: Never attempt to cut stylet.

Caution: Always withdraw stylet back beyond the tip of the catheter prior to insertion.

- Once proper catheter length and stylet position has been achieved, tighten locking collar to keep stylet in place.

- Remove dilator from sheath.

- Insert distal tip of catheter into and through the sheath until catheter tip is correctly positioned in the target vein.

- Remove the tearaway sheath by slowly pulling it out of the vessel while simultaneously splitting the sheath by grasping the tabs and pulling them apart (a slight twisting motion may be helpful).

Caution: Do not pull apart the portion of the sheath that remains in the vessel. To avoid vessel damage, pull back the sheath as far as possible and tear the sheath only a few centimeters at a time.

- Make any adjustments to catheter under fluoroscopy. The distal tip should be positioned at the level of the caval atrial junction.

Caution: Do not clamp the lumen portion of the catheter. Clamp only the extension(s). Do not use the serrated forceps, use only the in-line clamp(s) provided.

- Loosen locking collar of sideport. Remove the stylet by applying gentle pressure with one hand above the insertion site while grasping the stylet with the other hand and slowly pulling back with a constant motion. Remove sideport adaptor and replace with needleless access port. Attach saline filled syringe to needleless access port, aspirate lumen and then irrigate with saline. Remove syringe prior to clamping extension.

Caution: If difficulty and/or bunching of the catheter lumen are experienced while removing the stylet, additional

flushing of the catheter may be helpful. The catheter may need to be repositioned to allow for removal of the stylet.

Caution: Do not attempt to reinsert stylet once it has been withdrawn.

Caution: Never leave stylet in place after catheter insertion; injury may occur. Remove both stylet and sideport adaptor after insertion.

- Attach syringe(s) to extension(s) and open clamp(s). Blood should aspirate easily. If excessive resistance to blood aspiration is experienced, the catheter may need to be repositioned to obtain adequate flow.

- Once adequate aspiration has been achieved, lumen(s) should be irrigated with saline filled syringe(s). Clamp(s) should be open for this procedure.

Caution: Small syringes will generate excessive pressure and may damage the catheter. Ten (10)cc or larger syringes are recommended.

- Remove the syringe(s) and close extension clamp(s). Avoid air embolism by keeping catheter tubing clamped at all times when not in use and by aspirating then irrigating the catheter with saline prior to each use. With each change in tubing connections, purge air from the catheter and all connecting tubing and caps.

- Confirm and document proper tip placement with fluoroscopy prior to use. The distal tip should be positioned at the level of the caval atrial junction.

Caution: Failure to verify catheter placement may result in serious trauma or fatal complications.

Note: If there is no blood return, verify catheter position before use.

CATHETER SECUREMENT AND WOUND DRESSING:

- The insertion site and external portion of the catheter should always be covered with a protective dressing.

- Cover the exit site with an occlusive dressing according to the facility policy.

- Record catheter length, catheter lot number, and tip position on patient's chart.

DIRECTIONS FOR SELDINGER INSERTION

- Follow directions for Modified Seldinger Insertion, up to step #5.

- Remove needle, leaving guidewire in the targeted vein. Advance the guidewire until it reaches the caval atrial junction. Once the guidewire is in place, measure the depth of the guidewire by reading the markings on the wire.

- Cut catheter to length determined by marked guidewire.

- Insert proximal end of wire into distal tip of catheter lumen. Feed catheter lumen into the vessel following the guidewire. Advance catheter lumen along the guidewire until the distal tip is correctly positioned in the target vein. The distal tip should be positioned at the level of the caval atrial junction.

Caution: A skin nick may be required to feed the catheter smoothly into the vessel.

- Make any adjustments to catheter under fluoroscopy. The distal tip should be positioned at the level of the caval atrial junction.

Caution: Do not clamp the lumen portion of the catheter. Clamp only the extension(s). Do not use serrated forceps; use only in-line clamps provided.

- Remove the wire from the catheter. Remove by applying gentle pressure with one hand above the insertion site while grasping the 130cm wire with the other hand and pulling slowly back with a constant motion.

- Follow Directions for Modified Seldinger Insertion, from step #14 on.

POWER INJECTION PROCEDURE

- Remove the injection/needleless cap from the Pro-PICC® CT catheter.
- Using a 10cc or larger syringe(s), aspirate catheter lumen(s) to assure patency and remove locking solution. Discard syringe(s).
- Attach a 10cc or larger syringe filled with sterile normal saline and vigorously flush the catheter with the full 10cc of sterile normal saline. **Warning:** Failure to ensure patency of the catheter prior to power injection studies may result in catheter failure.
- Detach syringe.
- Attach the power injection device to the Pro-PICC® CT catheter per manufacturer's recommendations.

Warning: Always use connector tubing between power injector syringe and catheter. Do not attempt to connect power injector syringe directly to the catheter. Damage may result.

- Complete power injection study taking care not to exceed the flow rate limits. **Warning:** Exceeding the maximum indicated flow rate may result in catheter failure and/or catheter tip displacement.

- Disconnect the power injection device.

- Flush the Pro-PICC® CT catheter with 10cc of sterile normal saline, using a 10cc or larger syringe. For multi-lumen catheters, flush all lumens after power injection.

- Replace the injection/needleless cap on the Pro-PICC® CT catheter.

INFUSION

- Before infusion begins, all connections should be examined carefully.

- Frequent visual inspection should be conducted to detect leaks to prevent blood loss or air embolism.

- If a leak is found, the catheter should be clamped immediately and replaced.

- Necessary remedial action must be taken prior to the continuation of the treatment.

Note: Excessive blood loss may lead to patient shock.

CATHETER MAINTENANCE

- Dressing Changes** - A dressing should cover the insertion site at all times. The dressing should be changed per institutional policy or any time the dressing becomes soiled, wet, or non-occlusive.

Note: During all dressing changes, the external length of the catheter should be measured to determine if catheter migration has occurred. Periodically confirm catheter placement and tip location by imaging method.

- Flushing and Locking** - Flush and lock catheter according to your institutional policy.

- The catheter should be flushed with normal saline prior to drug administration to remove locking solution.

- After drug administration, each lumen should be flushed again with normal saline and then locked to maintain patency.

Injection Caps - Injection cap(s) or needleless access port(s) should be changed per institutional policy. If using the supplied needleless access port(s), do not exceed 100 actuations.

CATHETER PERFORMANCE

- Occluded/Partially Occluded Catheter- If resistance is encountered to aspirating or flushing, the lumen may be partially or completely occluded.

Warning: Do not flush against resistance.

- If the lumen will neither aspirate nor flush, and it has been determined that the catheter is occluded with blood, follow institutional de clotting procedure.

Infection:

Caution: Due to risk of exposure to HIV or other blood borne pathogens, health care professionals should always use Universal Blood and Body Fluid Precautions in the care of all patients.

- Sterile technique should always be strictly adhered to.

- Clinically recognized infection should be treated promptly per institutional policy.

CATHETER REMOVAL

Warning: Only a clinician familiar with the appropriate techniques should attempt the following procedure.

Caution: Always review facility protocol, potential complications and their treatment, warnings, and precautions prior to catheter removal.

- Wash hands, gather equipment.

- Remove old dressing/securement device and inspect insertion site for redness, tenderness, and drainage.

- Grasp catheter near insertion site and using a slow steady motion remove catheter from vein. **Caution:** Do not apply direct pressure to the insertion site or along the course of the vein.

Warning: Forceful removal can result in vein or catheter rupture.

- Apply pressure on site a few minutes to promote hemostasis as needed. Apply a sterile occlusive dressing per institutional policy.

Note: Inspect catheter and measure length. It must be equal to baseline measurement taken when the catheter was inserted.

TROUBLESHOOTING:

- Resistance is felt during removal – STOP!!! NEVER pull against resistance

a. Release any pressure along catheter path

b. Venous spasm may cause resistance; waiting a few minutes may allow the patient and the vein to relax

c. Warm packs placed proximal to the insertion site may help relax the vein walls and resolve the venospasm

d. Reposition the limb and try again after 20 minutes

Continued resistance: STOP, clean, secure the catheter, re-dress, and notify the physician.

PRO-PICC®				
Catheter Size	Gravity Flow		Full Length Priming Volume	
3F X 55CM SINGLE PRO-PICC® CT	6.12ml/min		0.46cc	
4F X 55CM SINGLE PRO-PICC® CT	20.60ml/min		0.70cc	
5F X 55CM DUAL PRO-PICC® CT	9.96 ml/min		0.60cc	
5F X 60CM SINGLE PRO-PICC® CT	28.40 ml/min		0.79cc	
6F X 60CM DUAL PRO-PICC® CT	11.75 ml/min		0.67cc	
	17Ga	19Ga	Purple Luer	Yellow & White Luer
6F X 60CM TRIPLE PRO-PICC® CT	15.43 ml/min	5.9 ml/min	0.69cc	0.39cc

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 in accordance with all relevant regulatory requirements.

Medcomp® and Pro-PICC® are trademarks of Medical Components, Inc. registered in the United States.

SYMBOL TABLE

^{5.1.1}	Manufacturer *
^{5.1.4}	Keep Dry *
^{5.4.2}	Do Not Re-use *
^{5.6.3}	Non-pyrogenic *
^{5.3.2}	Keep Away from Sunlight *
^{5.2.3}	Sterilized Using Ethylene Oxide *
^{5.2.8}	Do Not Use if Package is Damaged *
^{5.1.4}	Use-by Date *
^{5.2.1}	Do Not Resterilize *
^{5.1.5}	Batch/Lot Number *
^{5.1.8}	Catalogue Number *
	Prescription Use Only ***
^{5.1.2}	Authorized Representative in the European Community*
^{5.4.4}	Caution, consult Accompanying Documents *

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

*** FDA guidance Use of Symbols in Labeling.

Medical Components, Inc.
1499 Delp Drive
Harleysville, PA 19438 U.S.A.
Tel:215-256-4201
Fax:215-256-1787
www.medcompnet.com

MPS Medical Product Service GmbH
Borgasse 20
35619 Braunfels
Germany