



ARROW

3000 Bernville Road Reading, PA 19605 1-800-523-8446 8 a.m. - 8 p.m. EST

H-09800-105A (8/96)



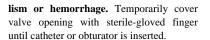


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Arrow International, Inc. recommends that the user be acquainted with the reference literature. *If you have any questions or would like additional reference information, please contact Arrow International, Inc.





Sheath Removal Procedure:

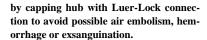
- 1. Precaution: Place the patient in a supine position.
- Remove dressing, if applicable. Precaution: To avoid cutting of the sheath, do not use scissors to remove the dressing.
- If applicable, remove sutures from sheath.
 Precaution: Be careful not to cut the sheath.
- 4. Withdraw device from sheath. Cover hemostasis valve with sterile-gloved finger. Warning: Hemostasis valve must be occluded at all times to minimize the risk of air embolism or hemorrhage.
- 5. Warning: Exposure of the central vein to atmospheric pressure may result in entry of air into the central venous system. Remove sheath slowly, pulling it parallel to the skin. As sheath exits the site, apply pressure with a dressing impermeable to air, e.g. vaseline gauze. Because the residual sheath track remains an air entry point until completely sealed, the occlusive dressing should remain in place for at least 24-72 hours dependent upon the amount of time the sheath was indwelling ^{12,16,17,19}.
- Upon removal of the sheath, inspect it to make sure that the entire length has been withdrawn.
- Verify that the sheath was intact upon removal.
- 8. Document removal procedure.

References:

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- Arditis J, Giala M, Anagnostidou A. Accidental puncture of the right lymphatic duct during pulmonary artery catheterization. Acta Anaesthesiol Scand. 1988;32:67-68.

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- Kashuk JL, Penn I. Air embolism after central venous catheterization. Surg Gynecol Obstet. September 1984;159:249-252.
- Kondo K, O'Reily LP, Chiota J. Air embolism associated with an introducer for pulmonary arterial catheters. *Anesth Analg.* 1984;63:871-872.





22. Hold catheter in place and reposition catheter contamination shield so that distal hub is approximately five inches (12.7 cm) from hemostasis valve/side port assembly (refer to Fig. 4).

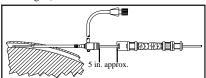


Fig. 4

23. Hold proximal hub of catheter contamination shield in place. Disengage distal hub from inner feed tube by pulling forward. Advance distal hub forward toward hemostasis valve/side port assembly. Hold assembly in place (refer to Fig. 5).

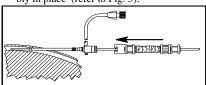


Fig. 5

24. Press distal hub of catheter contamination shield over assembly cap. Twist to lock (refer to Fig. 6).

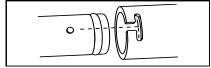


Fig. 6

- Orient slot in hub with locking pin on assembly cap.
- Slide hub forward over cap and twist.
- 25. While maintaining catheter position, twist the upper half of the distal hub in clockwise direction to lock catheter in place. Reposition proximal end of catheter shield as

desired. Twist upper and lower halves in opposite directions to lock in place. Test the adapter by gently tugging on the catheter to ensure a secure grip on the catheter (refer to Fig. 7). **Precaution: Do not reposition proximal hub once locked in final position.**

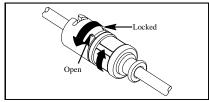


Fig. 7

- 26. Use suture tab to secure sheath and/or anchor with a purse string suture around the sheath suture ring. Precaution: Do not suture directly to the outside diameter of the sheath to avoid cutting or damaging the sheath or impeding flow.
- 27. Verify that the hemostasis valve/side port assembly to sheath connection is tightened securely. The connection may be further secured with tape to avoid patient tampering. Warning: Routinely examine connection to avoid inadvertent disconnection and possible air embolism, hemorrhage or exsanguination^{10,13}.
- 28. Dress puncture site per hospital protocol.

 Precaution: Maintain the insertion site with regular, meticulous redressing using aseptic technique.
- 29. Record the insertion procedure on the patient's chart.

Catheter Removal Procedure:

- 1. Precaution: Place the patient in a supine position.
- Remove dressing, if applicable. Precaution: To avoid cutting of the sheath, do not use scissors to remove the dressing.
- 3. Withdraw catheter from sheath. Warning: Hemostasis valve must be occluded at all times to minimize the risk of air embo-



Alternate technique:

Introducer needle may be used in the standard manner as alternative to catheter/needle assembly.

13. Because of the potential for inadvertent arterial placement, verify venous access via a wave form obtained by a calibrated pressure transducer (refer to Fig. 3).



Fig. 3

If a pressure transducer is not available, check for pulsatile flow. Pulsatile flow is usually an indicator of inadvertent arterial puncture.

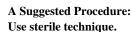
- 14. Insert desired tip of spring-wire guide through the introducer needle or catheter into vein. If the "J" tip is used, prepare for insertion by sliding the plastic tube over the "J" to straighten. Advance the spring-wire guide in the routine fashion to the desired depth. Advancement of "J" tip may require a gentle rotating motion. Warning: Do not cut spring-wire guide to alter length. Do not withdraw spring-wire guide against needle bevel to avoid possible severing or damaging of spring-wire guide.
- 15. Hold spring-wire guide in place and remove introducer needle or catheter. Precaution: Maintain firm grip on spring-wire guide at all times.
- 16. Enlarge cutaneous puncture site with cutting edge of scalpel positioned away from the spring-wire guide. Precaution: Do not cut guide wire.
- 17. Thread tapered tip of dilator/sheath/valve assembly over spring-wire guide. Grasping near skin, advance assembly with slight twisting motion to a depth sufficient to enter vessel. Dilator may be partially withdrawn to facilitate advancement of sheath through tortuous vessel. Precaution: Do not with-

draw dilator until the sheath is well within the vessel to prevent damage to sheath tip.

- 18. Advance sheath/valve assembly off dilator into vessel, again grasping near skin and using slight twisting motion.
- 19. To check for proper sheath placement within the vessel, remove side port end cap and attach syringe for aspiration. Hold sheath/valve assembly in place and withdraw spring-wire guide and dilator sufficiently to allow venous blood flow to be aspirated into side port. Precaution: Maintain firm grip on spring-wire guide at all times.
- 20. Holding sheath/valve assembly in place, remove guide wire and dilator as a unit. Place sterile-gloved finger over hemostasis valve. Warning: To avoid possible vessel wall perforation do not leave vessel dilator in place as an indwelling catheter. Warning: Although the incidence of spring-wire guide failure is extremely low, practitioner should be aware of the potential for breakage if undue force is applied to the wire. Flush and connect side port to appropriate line as necessary.
- 21. Feed catheter through sheath/valve assembly into vessel. Advance catheter to desired position. Warning: Hemostasis valve must be occluded at all times to minimize the risk of air embolism or hemorrhage. If catheter introduction is delayed, temporarily cover valve opening with sterilegloved finger until obturator is inserted. Use Arrow obturator, either included with this product or sold separately, as dummy catheter with hemostasis valve/ side port assembly and sheath. This will ensure that leakage does not occur and seal is protected from contamination¹⁵. Precaution: Do not disconnect hemostasis valve/side port assembly from sheath hub without occluding lumen with sterile-gloved finger followed







- Precaution: Place patient in slight Trendelenburg position as tolerated to reduce the risk of air embolism. If femoral approach is used, place patient in supine position.
- 2. Prep area of anticipated venipuncture.
- 3. Drape puncture site as required.
- 4. Perform skin wheal using desired needle. In kits where provided, a SharpsAway[®] disposal cup is used for the disposal of needles. Push needles into foam after use. Discard entire cup at completion of procedure. Precaution: Do not reuse needles after they have been placed into the disposal cup. Particulate matter may adhere to needle tip.
- 5. Prepare flow-directed catheter according to manufacturer's instructions. Wet balloon with flush solution to facilitate passage through valve of catheter contamination shield. Precaution: Do not inflate balloon prior to insertion through catheter contamination shield to minimize the risk of balloon damage.
- Ensure that double TwistLockTM of catheter contamination shield is fully opened (refer to Fig. 1).

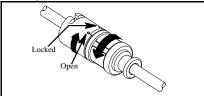


Fig. 1

Insert tip of desired catheter through proximal end of catheter contamination shield. Advance catheter through tubing and hub at other end (refer to Fig. 2).

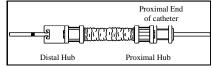


Fig. 2

- 7. Slide entire catheter contamination shield to proximal end of catheter.
- 8. If flow directed catheter is used, inflate and deflate balloon with syringe to ensure integrity. Precaution: Do not exceed balloon catheter manufacturer's recommended volume. Place catheter and catheter contamination shield on sterile field awaiting final sheath placement.
- 9. Secure hemostasis valve/side port assembly to sheath. Connection must be tight. Precaution: If sheath and dilator are preassembled, dilator must be removed prior to connecting assembly to sheath. Warning: Hemostasis valve/side port assembly to sheath connection must be tightened securely and routinely examined to avoid disconnection and possible air embolism, hemorrhage or exsanguination^{10,13}.
- 10. Insert entire length of dilator through hemostasis valve into sheath pressing hub of dilator firmly into hub of hemostasis valve/side port assembly. Place assembly on sterile field awaiting final sheath placement.
- 11. In kits where provided, use a 22 Ga. needle and syringe to locate central vein.
- 12. Insert introducer catheter/needle assembly with attached syringe into vein beside locater needle and aspirate. Remove locater needle. Withdraw needle and attached syringe from introducer catheter. If no free flow of venous blood is observed after needle is removed, attach syringe to the catheter and aspirate until good venous blood flow is established. Precaution: The color of the blood aspirated is not always a reliable indicator of venous access¹¹. Do not reinsert needle into introducer catheter.



Warnings and Precautions:*

- 1. Warning: Practitioners must be aware of complications associated with percutaneous sheath introduction including vessel wall perforation 18 , pleural and mediastinal injuries 1,14 , air embolism 5,10,13,15 , sheath embolism, thoracic duct laceration², bacteremia. septicemia. thrombosis³, inadvertent arterial puncture⁶, nerve damage, hematoma formation, hemorrhage⁴, and dysrhythmias.
- 2. Warning: Do not apply excessive force in removing guide wire, dilator or sheath. If withdrawal cannot be easily accomplished, a chest x-ray should be obtained and further consultation requested.
- 3. Warning: The practitioner must be aware of potential air embolism problems associated with leaving open needles, sheaths, or catheters in venous puncture sites or as a consequence of inadvertent disconnects. To lessen the risk of disconnects, only securely tightened Luer-Lock connections should be used with this device. Follow hospital protocol for all sheath and side port maintenance to guard against air embolism.
- 4. Warning: The connection between the hemostasis valve/side port assembly to sheath must be tightened securely and examined routinely to avoid disconnection and possible air embolism, hemorrhage or exsanguination^{10,13}.
- 5. Warning: Hemostasis valve must be occluded at all times to minimize the risk of air embolism or hemorrhage. If catheter introduction is delayed, or catheter is removed, temporarily cover valve opening with sterile-gloved finger until catheter or obturator is inserted. Use Arrow obturator, either included with this product or sold separately, as dummy catheter with hemostasis valve/side port assembly and sheath. This will ensure that leakage does not occur and inner seal is protected from contamination¹⁵.

- 6. Warning: Care should be exercised in passing spring-wire guide. Use of excessive length of the guide wire into the right heart can cause dysrhythmias, right bundle branch block⁹, and vessel wall, atrial or ventricular perforation.
- 7. Warning: Due to the risk of exposure to HIV (Human Immunodeficiency Virus) or other blood borne pathogens, health care workers should routinely use universal blood and body-fluid precautions in the care of all patients.
- Precaution: Do not suture directly to the outside diameter of the sheath to avoid cutting or damaging the sheath or impeding sheath flow.
- Precaution: Indwelling sheaths should be routinely inspected for desired flow rate, security of dressing, correct position and for proper Luer-Lock connection.
- Precaution: Maintain the insertion site with regular meticulous redressing using aseptic technique.
- 11. Precaution: Alcohol and acetone can weaken the structure of polyurethane material. Therefore, care should be taken when instilling drugs containing alcohol or when using high concentration of alcohol or acetone when performing routine insertion site care and maintenance. Alcohol should not be utilized to declot polyurethane sheaths.
- 12. Precaution: Do not disconnect hemostasis valve/side port assembly from sheath hub without occluding lumen with sterilegloved finger followed by capping hub with Luer-Lock connection to prevent possible air embolism, hemorrhage or exsanguination.
- 13. Precaution: Do not inflate balloon of flow-directed catheter prior to insertion through catheter contamination shield to minimize the risk of balloon damage.

Carefully read all warnings and precautions throughout procedure instructions.





Percutaneous Sheath Introducer Product

Safety and Efficacy Considerations:

The product is designed for single use only. Do not resterilize or reuse. Do not alter the sheath or any other kit/set component during insertion, use, or removal.

Procedure must be performed by trained personnel well versed in anatomical landmarks, safe technique and potential complications.

Two-Piece Percutaneous Sheath Introducer Warning

Two-piece percutaneous sheath introducers (PSIs) are used in the clinical setting because they have advantages over a one-piece or integral introducer in certain clinical situations. The two-piece introducers consist of a detachable hemostasis valve/side port assembly that attaches to a sheath via a Luer-Lock connection. Advantages to this type of PSI include the following:

- (1) the ability to remove the hemostasis valve/side port to facilitate a greater flow rate directly through the sheath during fluid resuscitation.
- (2) the ability to test the integrity of the balloon on a balloon catheter after it has been put through the hemostasis valve.
- (3) the ability to replace the hemostasis valve with a new, sterile assembly of the same size or one that is larger or smaller, depending on the need for different sized catheters or wires to be placed through the sheath.

Although there are advantages to the twopiece design, there is also an inherent risk that is associated with its use. If the connection is not secured and maintained properly, or excess tension is placed on the connection, the two parts can separate and place the patient at risk for AIR EMBOLISM, HEMORRHAGE or EXSANGUINATION. Cases of two-piece PSI separation have been documented in the literature^{8,10}.

To reduce the potential for serious complications associated with two-piece disconnects the following recommendations should be observed:

- discontinue the use of the percutaneous sheath introducer as soon as possible after the cardiac or central venous catheter has been removed^{7,13}.
- PSIs should be maintained in closely monitored environments (e.g. critical care units, operating room, recovery room), not on general nursing units¹⁵.
- securely tighten and routinely examine the hemostasis valve/side port assembly to sheath connection^{10,13}.
- do not use two-piece introducers on patients who are agitated, disoriented or uncooperative⁷.
- avoid placing traction on the side port lumen or intravenous tubing attached to the side port.
- tape the side port lumen securely to the patient's skin⁷.
- use of a one-piece PSI (available from Arrow International, Inc.) is also recommended for further risk reduction⁷.

Indications for Use:

The Arrow Percutaneous Sheath Introducer permits venous access and catheter introduction to the central circulation.

Contraindications:

None known.

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