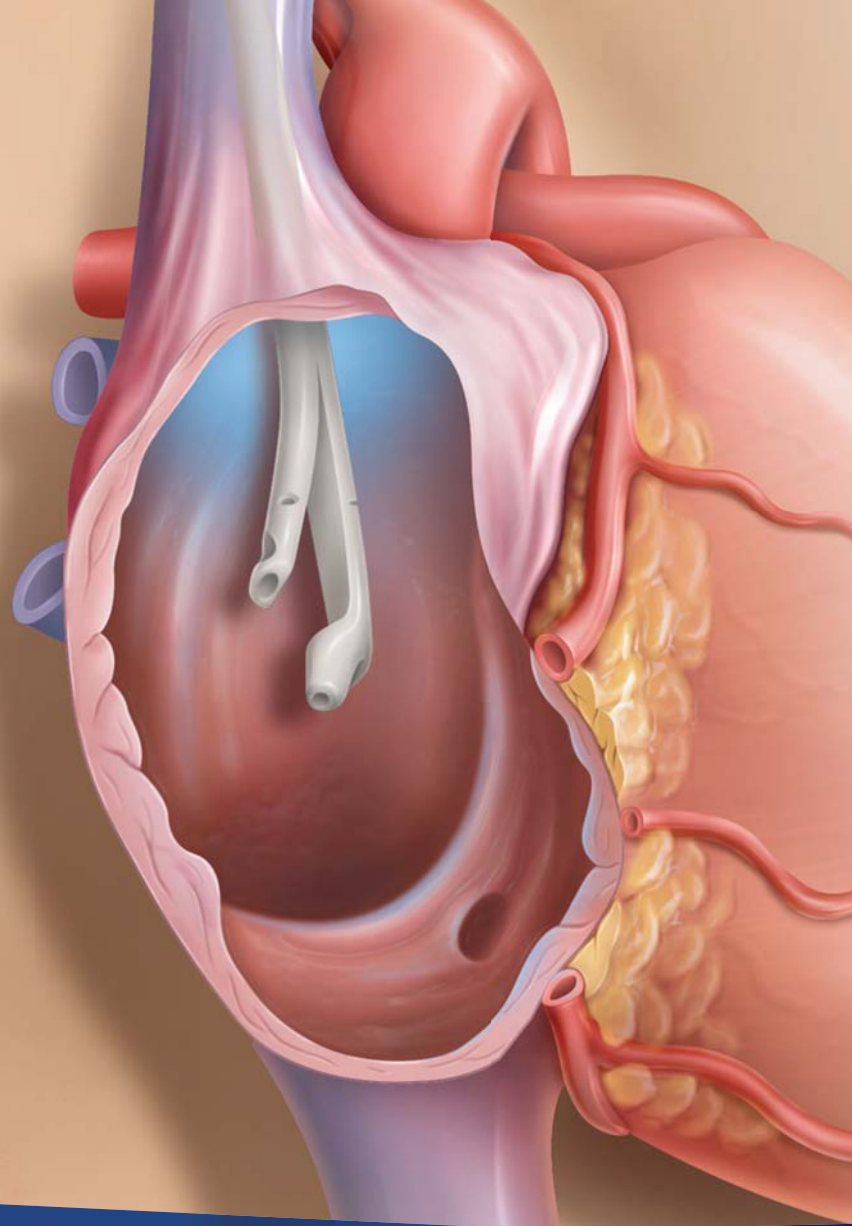


Step Tip Placement Split Tip Performance

Equistream™
Long-Term Hemodialysis Catheter

Equistream™ XK
Long-Term Hemodialysis Catheter



The split-tip technology of the 14.5F Equistream™ and 16F Equistream™ XK Long-Term Hemodialysis Catheters provides optimized flow for efficient dialysis and ease of access for right atrium placement.



Efficient Dialysis

- Nested tip design enables right atrium placement of both tips for optimal flow per KDOQI guidelines¹
- Recirculation <2% on average in forward and reverse when tested in-vitro²
- High flow rates of 500 ml/min on average in forward²



Multiple Choices

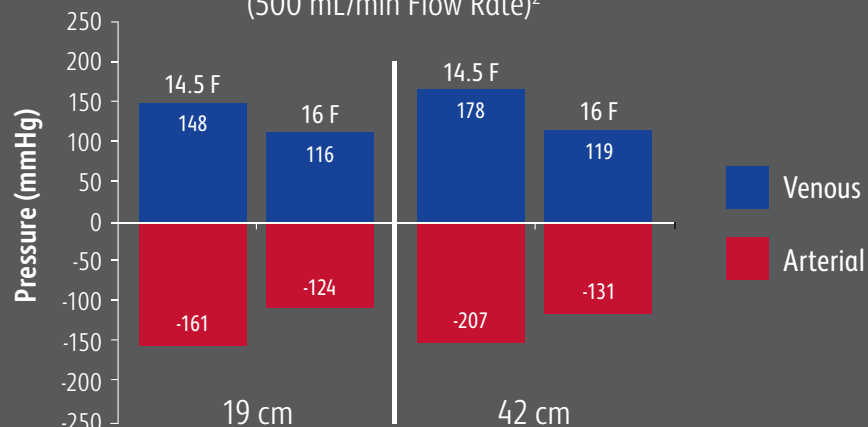
- Wide range of lengths
- Straight or Alphacurve™ Pre-Curved Configurations



Ease of Use

- Preloaded stylet facilitates easier over-the-wire insertion³
- Kits include the AirGuard™ Valved Introducer designed to help reduce the risk of air embolism and blood loss

14.5 F vs. 16 F Catheter Performance (500 mL/min Flow Rate)²



Bench data on file. May not necessarily correlate to clinical performance. Based on simulated testing. Different tests methods may yield different results.

¹ NKF K-DOQI Guideline 12 (<https://www.kidney.org/professionals/guidelines>).

² Tested using 19 cm and 42 cm Equistream™ and Equistream™ XK tip to cuff straight catheters (n=30)

³ Straight codes only.

Large luer connectors designed for durability during long-term use

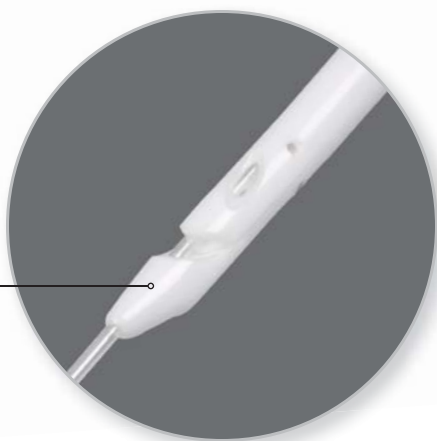
Fixed suture wings to promote and ensure stability

SureCuff™ tissue in-growth cuff

360° multiple side holes provide alternate flow paths

The AlphaCurve™ Pre-curved catheter has 16 mm more arc length on average than the 180° U-curve HemoSpit™ Hemodialysis Catheter, which can provide more tip adjustability*

Nested split tip design enables right atrium placement of both tips



* Data on File. Becton, Dickinson and Company, Tempe, AZ

Equistream™

Long-Term Hemodialysis Catheter

Insertion Length	Catheter Length	Product Code
14.5F, Straight, Polyurethane Catheter, Standard Kit		
15 cm	20 cm	<input type="text"/> 5903150
19 cm	24 cm	<input type="text"/> 5903190
23 cm	28 cm	<input type="text"/> 5903230
27 cm	32 cm	<input type="text"/> 5903270
31 cm	36 cm	<input type="text"/> 5903310
35 cm	40 cm	<input type="text"/> 5903350
42 cm	47 cm	<input type="text"/> 5903420
14.5F, AlphaCurve™, Polyurethane Catheter, Standard Kit		
19 cm	25 cm	<input type="text"/> 5905190
24 cm	29 cm	<input type="text"/> 5905240
28 cm	33 cm	<input type="text"/> 5905280
31 cm	36 cm	<input type="text"/> 5905310
14.5, Straight, Polyurethane Catheter, Microintroducer Kit		
19 cm	24 cm	<input type="text"/> 5904190
23 cm	28 cm	<input type="text"/> 5904230
27 cm	32 cm	<input type="text"/> 5904270

Product and Packaging Are Not Made with Natural Rubber Latex

Equistream™ XK

Long-Term Hemodialysis Catheter

Insertion Length	Catheter Length	Product Code
16F, Straight, Polyurethane Catheter, Standard Kit		
19 cm	24 cm	<input type="text"/> 5913190
23 cm	28 cm	<input type="text"/> 5913230
27 cm	32 cm	<input type="text"/> 5913270
31 cm	36 cm	<input type="text"/> 5913310
35 cm	40 cm	<input type="text"/> 5913350
42 cm	47 cm	<input type="text"/> 5913420
16F, AlphaCurve™, Polyurethane Catheter, Standard Kit		
19 cm	25 cm	<input type="text"/> 5915190
24 cm	29 cm	<input type="text"/> 5915240
28 cm	33 cm	<input type="text"/> 5915280
<div> <div>REPRESENTATIVE'S NAME</div> <div>CONTACT PHONE NO.</div> <div>PHYSICIAN'S SIGNATURE</div> </div>		

Standard Kit Contents - 14.5F or 16F (XK)

- 2 Each Adhesive Dressings
- 1 Each AirGuard™ Valved Introducer – 15F (16.5F if XK)
- 1 Each Dilator – 8F
- 1 Each Dual Lumen Catheter – 14.5F (16F if XK)
- 1 Each Dualator™ Vessel Dilator – 10-12F
- 2 Each End Caps
- 1 Each Guidewire – 70 cm x 0.038 in.
- 1 Each Introducer Needle – 18G
- 1 Each Tunneler
- 1 Each Insertion Stylet

Additional XK Standard Kit Contents - 16F (XK)

- 1 Each Dualator™ Vessel Dilator – 14-16F
- 1 Each Dualator™ Vessel Dilator – 15.5-17.5F
- 1 Each Scalpel

Microintroducer Kit Contents - 14.5F Only

- 2 Each Adhesive Dressings
- 1 Each AirGuard™ Valved Introducer – 15F
- 1 Each Dilator – 8F
- 1 Each Dual Lumen Catheter – 14.5F
- 1 Each Dualator™ Vessel Dilator – 10-12F
- 1 Each Dualator™ Vessel Dilator – 14-16F
- 1 Each Dualator™ Vessel Dilator – 15.5-17.5F
- 2 Each End Caps
- 1 Each Guidewire – 45 cm x 0.018 in.
- 1 Each Guidewire – 120 cm x 0.038 in.
- 1 Each Introducer Needle – 21G
- 1 Each MicroIntroducer – 5F
- 1 Each Tunneler
- 1 Each Insertion Stylet

Equistream™ and Equistream™ XK Long-Term Hemodialysis Catheters Indications For Use

The Equistream™ and Equistream™ XK Long-Term Hemodialysis Catheters are indicated for use in attaining short-term or long-term vascular access for hemodialysis, hemoperfusion, or apheresis therapy. Access is attained via the internal jugular vein, external jugular vein, subclavian vein, or femoral vein. Catheters longer than 40 cm are intended for femoral vein insertion.

Contraindication

This device is contraindicated for patients exhibiting severe, uncontrolled thrombocytopenia or coagulopathy.

Warnings

WARNING: Percutaneous insertion of the catheter should be made into the axillary-subclavian vein at the junction of the outer and mid-thirds of the clavicle lateral to the thoracic outlet. The catheter should not be inserted into the subclavian vein medially, because such placement can lead to compression of the catheter between the first rib and clavicle and can lead to damage or fracture and embolization of the catheter.¹ Fluoroscopic or radiographic confirmation of catheter tip placement should be helpful in demonstrating that the catheter is not being pinched by the first rib and clavicle.¹ • Alcohol or alcohol-containing antiseptics (such as chlorhexidine) may be used to clean the catheter/skin site; however, care should be taken to avoid prolonged or excessive contact with the solution(s). Solutions should be allowed to completely dry before applying dressing. • Acetone and Polyethylene Glycol (PEG)-containing ointments can cause failure of this device and should not be used with polyurethane catheters. Chlorhexidine patches or bacitracin zinc ointments (e.g., Polysporin™ ointment) are the preferred alternative. • Follow Universal Precautions when inserting and maintaining this device. • Cardiac arrhythmias may result if the guidewire and/or stylet touches the walls of the right atrium. Use cardiac rhythm monitoring to detect arrhythmias. • Close all clamps used in the center of the extension legs. Extensions may develop cuts or tears if subjected to excessive pulling or contact with rough edges. Repeated clamping near or on the Luer-lock connectors may cause tubing fatigue and possible disconnection. • Catheters should be implanted carefully. • Any sharp or acute angles that could compromise the opening of the catheter lumens need to be avoided. • To prevent air embolism and/or blood loss put patient in Trendelenburg position and always place thumb over the exposed orifice of the sheath introducer. • To avoid damage to vessels and viscous, infusion pressures should not exceed 25 psi (172 kPa). The use of a 10 mL or larger syringe is recommended because smaller syringes generate more pressure than larger syringes. Note: A three pound (13.3 Newton) force on the plunger of a 3 mL syringe generates pressure in excess of 30 psi (206 kPa) whereas the same three pound (13.3 Newton) force on the plunger of a 10

mL syringe generates less than 15 psi (103 kPa) of pressure. • Accessories and components used in conjunction with this catheter should incorporate Luer-lock adapters. • The heparin solution must be aspirated out of both lumens immediately prior to using the catheter to prevent systemic heparinization of the patient. • Failure to clamp extensions when not in use may lead to air embolism. • In the rare event of a leak, the catheter should be clamped immediately. Necessary remedial action must be taken prior to resuming dialysis or infusion procedure. • The risk of infection is increased with femoral vein insertion. • Do not resterilize the catheter or components by any method. The manufacturer will not be liable for any damages caused by reuse of the catheter or accessories. • Cannulation of the left internal jugular vein was reportedly associated with a higher incidence of complications compared to catheter placement in the right internal jugular vein.² • Alcohol should not be used to lock, soak or decontaminate polyurethane catheters over time with repeated and prolonged exposure. • Intended for Single Use. DO NOT REUSE. Reuse and/or repackaging may create a risk of patient or user infection, compromise the structural integrity and/or essential material and design characteristics of the device, which may lead to device failure, and/or lead to injury, illness or death of the patient.

Cautions

• Repeated over tightening of blood lines, syringes and caps will reduce connector life and could lead to potential connector failure. In case of damage, clamp the catheter between the patient and the damaged area with a smooth-edged, atraumatic clamp. • Sterile and non-pyrogenic only if packaging is not opened, damaged or broken. • Read the instructions for use carefully before using this device. • CAUTION: Federal (USA) law restricts this device to sale by or on the order of a physician. • Left sided placement in particular, may provide unique challenges due to the right angles formed by the innominate vein and at the left brachiocephalic junction with the SVC.^{3,4} • Care should be taken NOT to force the dilator sheath introducer assembly into the vessel during insertion as vessel damage including perforation could result. • Stylet is intended for use over a guidewire to aid in placement. Inserting the stylet into the venotomy without tracking over a guidewire could result in vessel damage including perforation. • Failure to retract the stylet when inserting the tunneler into the catheter tip can result in damage to the stylet. • Ensure that the catheter does not move out of the vein while removing the insertion stylet. • Care should be taken not to advance the split sheath too far into vessel as a potential kink would create an impasse to the catheter. • Ensure that the introducer sheath is only torn externally. Catheter may need to be further pushed into the vessel as sheath is torn. • For optimal product performance, do not insert any portion of the cuff into the vein. • If the microintroducer

guidewire must be withdrawn while the needle is inserted, remove both the needle and wire as a unit to prevent the needle from damaging or shearing the guidewire. • Before attempting the insertion of Equistream™ Catheters, ensure that you are familiar with the complications listed below and their emergency treatment should any of them occur. • The complications listed below as well as other complications are well documented in medical literature and should be carefully considered before placing the catheter. Placement and care of Equistream™ Catheters should be performed by persons knowledgeable of the risks involved and qualified in the procedures.

Possible Complications

The use of an indwelling central venous catheter provides an important means of venous access for critically ill patients; however, the potential exists for serious complications including the following: • Air Embolism • Arterial Puncture • Bleeding • Brachial Plexus Injury • Cardiac Arrhythmia • Cardiac Tamponade • Catheter or Cuff Erosion Through the Skin • Catheter Embolism • Catheter Occlusion • Catheter Occlusion, Damage or Breakage due to Compression Between the Clavicle and First Rib¹ • Catheter-related Sepsis • Endocarditis • Exit Site Infection • Exit Site Necrosis • Extravasation • Fibrin Sheath Formation • Hematoma • Hemomediastinum • Hemothorax • Hydrothorax • Inflammation, Necrosis or scarring of skin over implant area • Intolerance Reaction to Implanted Device • Laceration of Vessels or Viscus • Perforation of Vessels or Viscus • Pneumothorax • Thoracic Duct Injury • Thromboembolism • Venous Stenosis • Venous Thrombosis • Ventricular Thrombosis • Vessel Erosion • Risks Normally Associated with Local and General Anesthesia, Surgery, and Post-Operative Recovery.

References

- Aitken, D.R. and Minton, J.P. "The Pinch-Off Sign: A Warning of Impending Problems with Permanent Subclavian Catheters", American Journal of Surgery, Vol. 148, Nov. 1984, pp.633-638.
- Mickley, V., "Central venous catheters: many questions: few answers", Nephrol Dial Transplant, (2002) 17:1368-1373.
- Sulek, CA., Blas, ML., Lobato, EB., "A randomized study of left versus right internal jugular vein cannulation in adults." J Clin Anesth. 2000 Mar;12(2):142-5
- Tan, P.L., Gibson, M., "Central Venous Catheters: the role of radiology", Clin Rad. 2006, 61:13-22

Please consult package inserts for more detailed safety information and instructions for use.