

SHORT TERM CATHETERS IMPORTANT RISK INFORMATION

SLX®

Indications for Use: The Medcomp® SLX Double Lumen Catheter can be utilized for temporary access for hemodialysis, hemoperfusion, or apheresis therapy. The cannula may be inserted via the Seldinger technique due to the inner Teflon stylet, increasing linear strength. The stylet is removed after insertion, leaving the soft silicone cannula in the body. The flexible silicone make-up conforms well to the vessel anatomy, resulting in higher patient tolerance during extended use.

Contraindications: The Subclavian Approach is NOT recommended for use with the Medcomp® SLX Double Lumen Subclavian-Femoral Catheter in Hemodialysis or Hemoperfusion Procedures used for the management of acute poisoning or other situations in which a ventilator might be used due to risk of traumatic pneumothorax posing a dangerous complication for the patient.

DUO-SPLIT®

Indications for Use: The Medcomp® Duo-Split® Double Lumen Catheter is designed for acute hemodialysis and apheresis. It may be inserted percutaneously and is ideally placed in the internal jugular vein. Although this catheter may be inserted into the subclavian vein or femoral vein, the internal jugular vein is the preferred site.

Contraindications: This catheter is not intended for any use other than that which is indicated. Do not insert catheter in thrombosed vessels.

HEMO-CATH® ST

Indications for Use: The Medcomp® Hemo-Cath® ST Silicone Double Lumen Catheter can be utilized for temporary access for hemodialysis, hemoperfusion, or apheresis therapy. The cannula may be inserted via the Seldinger technique due to the inner Teflon stylet, increasing linear strength. The stylet is removed after insertion, leaving the soft silicone cannula in the body. The flexible silicone make-up conforms well to the vessel anatomy, resulting in higher patient tolerance during extended use.

Contraindications: The Subclavian Approach is NOT recommended for use with the Medcomp® Hemo-Cath® ST Double Lumen Subclavian-Femoral Catheter in Hemodialysis or Hemoperfusion Procedures used for the management of acute poisoning or other situations in which a ventilator might be used due to risk of traumatic pneumothorax posing a dangerous complication for the patient.

DUO-FLOW® 400XL

Indications for Use: The Medcomp® Duo-Flow and Duo-Flow 400XL Double Lumen Catheters are designed for acute hemodialysis and apheresis. They may be inserted percutaneously and are ideally placed in the internal jugular vein. Although these catheters may be inserted into the subclavian or femoral vein, the internal jugular is the preferred site.

Contraindications: This catheter is intended for Short-Term vascular access only and should not be used for any purpose other than indicated in these instructions.

DUO-FLOW®

Indications for Use: The Medcomp® Duo-Flow® Catheter is indicated for use in attaining Short-Term vascular access for Hemodialysis and Apheresis. It may be inserted percutaneously and is primarily placed in the internal jugular vein of an adult patient. Alternate insertion sites include subclavian vein or femoral vein as required. The curved Duo-Flow® Catheter is intended for internal jugular vein insertion. This catheter is indicated for a duration less than (30) days. For femoral placement, monitor catheter condition closely.

Contraindications: This catheter is intended for Short-Term vascular access only and should not be used for any purpose other than indicated in these instructions.

DUO-FLOW® SIDE X SIDE

Indications for Use: The Duo-Flow® Side X Side double lumen catheter is intended for short-term central venous access for hemodialysis, apheresis and infusion.

Contraindications: The catheter is not intended for any purpose other than indicated in these instructions. The Duo-Flow® Side X Side double lumen catheter is intended for short-term (less than 30 days) use. Do not use this catheter in thrombosed vessels or for subclavian puncture when a ventilator is in use. Do not use this catheter when: The patient's body size is insufficient to accommodate the implanted device. The superficial or deep tissue will not permit adequate device stabilization and/or access. There are known physiological limitations that will not allow placement of the device. The patient has known or suspected allergies to any of the materials in the device. The patient has received significant radiation at the intended exit site or tunnel. The patient has severe chronic obstructive lung disease.

DUO-FLOW® SOFT-LINE®

Indications for Use: The Medcomp® Duo-Flow® Soft-Line® and Raulerson/Duo-Flow® Double Lumen Internal Jugular Catheters are designed for acute hemodialysis and apheresis. They may be inserted percutaneously and are ideally placed in the jugular vein.

Contraindications: This catheter is not intended for any use other than that which is indicated. Do not implant catheter in thrombosed vessels.

T-3® CT

Indications for Use: The Medcomp® T-3® CT Catheter is a triple lumen catheter indicated for use in attaining short-term vascular access for hemodialysis, apheresis. The third internal lumen is intended for infusion, power injection of contrast media and central venous pressure monitoring. The catheter is intended to be inserted in the jugular, femoral or subclavian vein as required. The maximum recommended infusion rate is 5ml/sec for power injection of contrast media.

Contraindications: This catheter is intended for short-term (less than 30 days) vascular access only and should not be used for any purpose other than indicated in these instructions. This device is also contraindicated: When the presence of device related infection, bacteremia, or septicemia is known or suspected. When the patient's body size is insufficient to accommodate the size of the implanted device. When the patient is known or is suspected to be allergic to materials contained in the device. If the prospective insertion site has been previously irradiated. If the prospective placement site has previously suffered episodes of venous thrombosis or vascular surgical procedures. If local tissue factors may prevent proper device stabilization and/or access.

TRI-FLOW

Indications for Use: The 12F Tri-Flow Triple Lumen Catheter is indicated for use in attaining Short-Term vascular access for Hemodialysis and Apheresis. It may be inserted percutaneously and is primarily placed in the internal jugular vein of an adult patient. Alternate insertion sites include subclavian vein or femoral vein as required. The 12F Tri-Flow Triple Lumen Catheter is intended to be used less than (30) days.

Contraindications: This catheter is intended for Short-Term vascular access only and should not be used for any purpose other than indicated in these instructions. Do not insert catheter in thrombosed vessels.

TRIO-CT®

Indications for Use: The Trio-CT® Triple Lumen Catheter is indicated for use in attaining short-term (less than 30 days) vascular access for hemodialysis and apheresis. The third internal lumen is intended for infusion, power injection of contrast media and central venous pressure monitoring. The catheter is intended to be inserted in the jugular, femoral or subclavian vein as required. The maximum recommended infusion rate is 5ml/sec for power injection of contrast media.

Contraindications: This catheter is intended for short-term (less than 30 days) vascular access only and should not be used for any purpose other than indicated in these instructions. This device is also contraindicated: When the presence of device related infection, bacteremia, or septicemia is known or suspected. When the patient's body size is insufficient to accommodate the size of the implanted device. When the patient is known or is suspected to be allergic to materials contained in the device. If the prospective insertion site has been previously irradiated. If the prospective placement site has previously suffered episodes of venous thrombosis or vascular surgical procedures. If local tissue factors may prevent proper device stabilization and/or access.

Refer to Instructions for Use provided with the product for complete instructions, warnings, precautions, and contraindications. Observe all instructions for use prior to using products. Failure to do so may result in patient complications.



DOUBLE "D" INTERNAL LUMEN DESIGN

Polyurethane
Material

Split Tip
Tip Design

13F
French Size

Straight,
Pre-Curved,
Raulerson IJ
Configurations



SPLIT TIP DESIGN



ARTERIOVENOUS WEAVE WITH DUAL STYLET



RAULERSON IJ CONFIGURATION



CATHETER ONLY		
DSP134C	13F X 12CM DUO-SPLIT®, STRAIGHT	10/BOX
DSP136C	13F X 15CM DUO-SPLIT®, STRAIGHT	10/BOX
DSP138C	13F X 20CM DUO-SPLIT®, STRAIGHT	10/BOX
DSP139C	13F X 24CM DUO-SPLIT®, STRAIGHT	10/BOX
DSP130C	13F X 30CM DUO-SPLIT®, STRAIGHT	10/BOX
DSP134PC	13F X 12CM DUO-SPLIT®, PRE-CURVED	10/BOX
DSP136PC	13F X 15CM DUO-SPLIT®, PRE-CURVED	10/BOX
DSP138PC	13F X 20CM DUO-SPLIT®, PRE-CURVED	10/BOX
DSP139PC	13F X 24CM DUO-SPLIT®, PRE-CURVED	10/BOX
DDSP136IJC	13F X 15CM DUO-SPLIT®, RAULERSON IJ	10/BOX
DDSP138IJC	13F X 20CM DUO-SPLIT®, RAULERSON IJ	10/BOX
DDSP139IJC	13F X 24CM DUO-SPLIT®, RAULERSON IJ	10/BOX

FULL TRAY		
DSP136T	13F X 15CM DUO-SPLIT®, STRAIGHT	5/BOX
DSP138T	13F X 20CM DUO-SPLIT®, STRAIGHT	5/BOX
DSP139T	13F X 24CM DUO-SPLIT®, STRAIGHT	5/BOX
DSP130T	13F X 30CM DUO-SPLIT®, STRAIGHT	5/BOX
DSP134PT	13F X 12CM DUO-SPLIT®, PRE-CURVED	5/BOX
DSP136PT	13F X 15CM DUO-SPLIT®, PRE-CURVED	5/BOX
DSP138PT	13F X 20CM DUO-SPLIT®, PRE-CURVED	5/BOX
DSP139PT	13F X 24CM DUO-SPLIT®, PRE-CURVED	5/BOX
DSP134IJT	13F X 12CM DUO-SPLIT®, RAULERSON IJ	5/BOX
DSP136IJT	13F X 15CM DUO-SPLIT®, RAULERSON IJ	5/BOX
DSP138IJT	13F X 20CM DUO-SPLIT®, RAULERSON IJ	5/BOX
DSP139IJT	13F X 24CM DUO-SPLIT®, RAULERSON IJ	5/BOX

BASIC SET		
DDSP134S	13F X 12CM DUO-SPLIT®, STRAIGHT	5/BOX
DDSP136S	13F X 15CM DUO-SPLIT®, STRAIGHT	5/BOX
DDSP138S	13F X 20CM DUO-SPLIT®, STRAIGHT	5/BOX
DDSP139S	13F X 24CM DUO-SPLIT®, STRAIGHT	5/BOX
DDSP130S	13F X 30CM DUO-SPLIT®, STRAIGHT	5/BOX
DDSP134PS	13F X 12CM DUO-SPLIT®, PRE-CURVED	5/BOX
DDSP136PS	13F X 15CM DUO-SPLIT®, PRE-CURVED	5/BOX
DDSP138PS	13F X 20CM DUO-SPLIT®, PRE-CURVED	5/BOX
DDSP139PS	13F X 24CM DUO-SPLIT®, PRE-CURVED	5/BOX
DDSP134IJS	13F X 12CM DUO-SPLIT®, RAULERSON IJ	5/BOX
DDSP136IJS	13F X 15CM DUO-SPLIT®, RAULERSON IJ	5/BOX
DDSP138IJS	13F X 20CM DUO-SPLIT®, RAULERSON IJ	5/BOX
DDSP139IJS	13F X 24CM DUO-SPLIT®, RAULERSON IJ	5/BOX

CATHETER ONLY CONTENTS:

- (1) Catheter with Dual Stylets
- (2) Dilators
- (2) End Caps

BASIC SET CONTENTS:

- (1) Catheter with Dual Stylets
- (1) Introducer Needle
- (1) Guidewire
- (2) Dilators
- (1) Adhesive Wound Dressing
- (2) End Caps

FULL TRAY CONTENTS:

- (1) Catheter with Dual Stylets
- (1) Introducer Needle
- (1) Guidewire
- (2) Dilators
- (1) Scalpel
- (1) Peelable Introducer
- (1) Adhesive Wound Dressing
- (2) End Caps
- (3) 4" x 4" Gauze
- (1) Chloraprep
- (1) 5cc Syringe
- (2) Pre-Filled Saline Syringes
- (1) 2-0 Silk Suture
- (3) Safety Hypodermic Needles
- (1) Hemostat
- (1) Fenestrated Drape
- (1) 5cc Lidocaine
- (1) Lidocaine/Saline Stickers
- (1) Filter Straw
- (1) Surgical Gloves
- (1) Needle Stick Pad

Refer to the Table of Contents for Important Risk Information regarding this device.