

REF CDA-34552-HPK1A

Contents: _

- Two-Lumen Indwelling TaperFree[™] Catheter: 5 Fr. x 45 cm Pressure Injectable Radiopaque Polyurethane with Blue FlexTip[®], T-Port Connector, Extension Line Clamps, Contamination Guard and Placement Wire
- GlideThru[™] Peel-Away Sheath: 5 Fr. x 2-3/4"
 (7 cm) Radiopaque over 5 Fr. Dilator
- Spring-Wire Guide, Nitinol, Marked: .018" (0.46 mm) dia. x 17-3/4" (45 cm) (Straight Soft Tip on One End – Straight Stiff Tip on Other) with Arrow Advancer[™]
- 1: Injection Needle: SafetyGlide^{™1} 25 Ga. x 5/8" (1.60 cm) RW
- 1: Safety Introducer Needle: Echogenic 21 Ga. x 2-3/4" (7 cm) TW
- 1: Introducer Needle: Echogenic 21 Ga. x 2-3/4" (7 cm) TW

- 1: Syringe: 10 mL Luer-Lock
- 1: Syringe: 3 mL Luer-Lock
- 2: Dust Caps: Non-Vented
- 1: SecondSite[™] Adjustable Hub: Fastener
- 1: SecondSite[™] Adjustable Hub: Catheter Clamp
- 1: SharpsAway II[™] Locking Disposal Cup
- 1: Catheter Trimmer
- 1: Drape: 24" x 36"
- 1: Drape: 60" x 76"
- 1: Drape: 36" x 41" with 3" x 5" fenestration
- 2: CSR Wraps
- 1: Towel
- 1: Filter: 5 Micron Straw
- 1: Safety Scalpel: #11
- 1: Flow Rate Information Card with Injection Log
- 1: Sterile Procedure Sign
- 1: Patient ID Card

- 1: Chart Sticker
- 1: Patient Information Booklet
- 1: Checklist/CLIP Sheet
- 2: Paper Tape Measures
- 1: Tourniquet
- 5: Gauze Pads: 2" x 2"
- 5: Gauze Pads: 4" x 4"
- 1: Surgical Apparel: Mask
- 1: Dressing: Tegaderm^{®2} 15.5 cm x 10 cm
- 1: Surgical Apparel: Impervious Gown
- 1: Surgical Apparel: Mask with Eye Shield
- 1: Surgical Apparel: Bouffant Cap
- 1: Tape: Steri-Strip®2
- 1: HemoHopper[®] Fluid Receptacle
- ¹ A trademark of Becton, Dickinson and Company.
- ² A registered trademark of 3M Company.

All components are CE 0086 unless otherwise noted.

Warning: This product contains a chemical known to the State of California to cause cancer, birth defects or other reproductive harm.

Contraindications: The Pressure Injectable PICC is contraindicated wherever there is presence of device related infections or presence of thrombosis in the intended insertion vessel or catheter pathway. Clinical assessment of patient must be completed to ensure no contraindications exist. See additional labeling for product specific contraindications.





Erg[®]Pack