

(IPNXXXXXX)

55 cm catheter length

Pressure Injectable Two-Lumen PICC



Contents:

- Two Lumen TaperFree® Catheter‡‡‡: 5 Fr. (1.81 mm OD) x 55 cm, Pressure Injectable, Stiffening Stylet (0.36 mm dia. x 75 cm with straight distal tip) and T-
- 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
- Introducer Needle: 21 Ga. x 1-1/2" (3.81 cm) TW
- Injection Needle: Eclipse™1 25 Ga. x 1" (2.54 cm)
- Safety Introducer Needle: Echogenic 21 Ga. x 2-3/4" (7 cm) TW
- Introducer Needle: Echogenic 21 Ga. x 2-3/4" (7 cm) TW
- Filter Needle: 18 Ga. TW x 1-1/2" (3.81 cm) with Blunt Fill Tip
- Svringe: 10 mL Luer-Lock
- Syringe: 3 mL Luer-Lock
- Dust Cap: Non-Vented
- SecondSite™ Adjustable Hub: Fastener
- SecondSite™ Adjustable Hub: Catheter Clamp
- SharpsAway® Disposal Cup
- SharpsAway® II Locking Disposal Cup
- Catheter Trimmer
- Towel: 24" x 36" (61 cm x 91 cm)
- Maximal Barrier Drape™ with two 13 cm (5") x 15 cm (6") fenestrations
- Safety Scalpel: #11
- Patient ID Card
- Sterile Procedure Sign
- 1: Patient Information Booklet

- 2: Paper Tape Measure
- 1: Tourniquet
- Gauze Pad: 2" x 2" (5 cm x 5 cm)
- 10: Gauze Pad: 4" x 4" (10 cm x 10 cm)
- Surgical Apparel: Impervious Gown
- Dressing: GRIP-LOK®2 Catheter Fixation Device
- Dressing: I.V. Advanced 8.5 cm x 11.5 cm Transducer Cover with Gel and Rubber Bands
- Surgical Apparel: Mask with Eye Shield
- Surgical Apparel: Bouffant Cap Surgical Apparel: Mask
- Tape: Steri-Strip®3
- 2: Valve: MicroCLAVE™4 Neutral Displacement Connector

¹A trademark of Becton, Dickinson and Company

- ²A registered trademark of TIDI.
- ³A registered trademark of 3M Company
- ⁴A trademark of ICU Medical, Inc.

All components are CE 2797 unless otherwise noted.











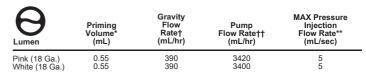




###MR Safe

Not made with natural rubber latex.

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 the patient must be completed to ensure no contraindications exist.



^{*} Priming volumes are approximate and are done without accessories.





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[†] Flow rate values are approximate and are determined using deionized water at 100 cm head

^{††} Pump flow rates are determined at maximum pump pressure of 10 psig and represent approximate flow capabilities.

^{**} Pressure injection flow rates are determined at the injector pressure setting of 300 psi maximum using media of 11.8 centipoise viscosity, with 152 cm pressure tubing.