



PICC Kit with Placement Wire and Integral Needle Protection

REF **PK-01451-TWSP**

4 Fr. | **55** cm catheter length

Contents:

- 1: Indwelling Catheter, Trimmable: 4 Fr. x 55 cm Radiopaque Polyurethane with T-Port Connector, Integral Extension Line, Extension Line Clamp, Integral Suture Wing, Contamination Guard and Placement Wire
- 1: Peel-Away Catheter: 15 Ga. x 1-1/4" (3.18 cm) Radiopaque over 17 Ga. TW Introducer Needle with Integral Needle Protection
- 1: Injection Needle: SafetyGlide™¹ 21 Ga. x 1-1/2" (3.81 cm) TW
- 1: Injection Needle: SafetyGlide™¹ 25 Ga. x 5/8" (1.60 cm) and 3 mL Luer-Lock Syringe
- 1: Syringe: 10 mL Luer-Lock
- 1: Syringe: 10 mL Luer-Slip
- 1: Skin Protectant Prep Pad (non-sterile solution)
- 1: 3 mL Applicator 2% CHG and 70% IPA ChloroPrep®² One-Step Solution with Hi-Lite Orange™ Tint (non-sterile solution)
- 1: Dust Cap: Non-Vented
- 1: SharpsAway II™ Locking Disposal Cup
- 1: SharpsAway® Disposal Cup
- 1: Catheter Trimmer
- 1: Towel
- 1: Towel: 24" x 36" (61 cm x 91 cm)

- 1: Drape: 24" x 36" (61 cm x 91 cm) with 4" (10 cm) fenestration
- 1: Forceps
- 1: Patient ID Card
- 1: Chart Sticker
- 1: Patient Information Booklet
- 1: Dressing: OpSite®³ 14 cm x 10 cm
- 2: Paper Tape Measure
- 1: Tourniquet
- 5: Gauze Pad: 2" x 2" (5 cm x 5 cm)
- 6: Gauze Pad: 4" x 4" (10 cm x 10 cm)
- 1: Dressing: STATLOCK®⁴ Catheter Stabilization Device
- 2: Surgical Apparel: Mask
- 1: Tape: Steri-Strip®⁵

¹A trademark of Becton, Dickinson and Company.
²A registered trademark of CareFusion or one of its subsidiaries.
³A registered trademark of T.J. Smith & Nephew, Limited.
⁴A registered trademark of C. R. Bard, Inc.
⁵A registered trademark of 3M Company.

DuPont™ Tyvek®
packaging

Rx only. **Warning: Read all package insert warnings, cautions and instructions prior to use. Failure to do so may result in severe patient injury or death.**

Sterilized using ethylene oxide unless otherwise indicated in the contents list.

California Prop. 65

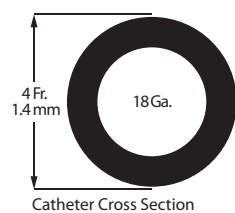
Warning: Cancer and Reproductive Harm www.P65Warnings.ca.gov.

Fluid path components are non-pyrogenic.

Contraindications: This device is contraindicated wherever there is presence of device related infections, previous or current thrombosis. Clinical assessment of patient must be completed to ensure no contraindications exists.

Not made with natural rubber latex

Store between 15 - 30°C (59 - 86°F).
Avoid freezing and excessive heat above 40°C (104°F)



Lumen	Priming Volume* (mL)	Flow Rate† (mL/hr)	Pump Flow Rate‡ (mL/hr)
Distal (18 Ga.)	0.52	850	3700

* Priming volumes are approximate and are done without accessories.
 † Flow rates are done with normal saline, room temperature, 100 cm head height and represent approximate flow capabilities.
 ‡ Pump flow rates are determined at maximum pump pressure of 10 psig.



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Manufacturer:
 Arrow International, Inc.
 Subsidiary of Teleflex Incorporated
 2400 Bernville Road
 Reading, PA 19605 USA

Teleflex®
www.teleflex.com

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