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Peripherally Inserted Central Catheter Kit with Blue FlexTip® Catheter and Placement Wire

Towel

Forceps

Patient ID Card

Chart Sticker

Tourniquet

1:

1:

1:

1:

2:

1:

Towel: 24" x 36" (61 cm x 91 cm)

Patient Information Booklet

Paper Tape Measure

Surgical Apparel: Mask

Dressing: OpSite®2 14 cm x 10 cm

Gauze Pad: 2" x 2" (5 cm x 5 cm)

Gauze Pad: 4" x 4" (10 cm x 10 cm)

Surgical Tape: Transpore™ 1" x 54" Roll

completed to ensure no contraindications exists.

¹A registered trademark of CareFusion or one of its subsidiaries.

Dressing: STATLOCK®3 Catheter Stabilization Device

Drape: 24" x 36" (61 cm x 91 cm) with 4" (10 cm) fenestration

REF PK-01451-WS

4 Fr. 55 cm catheter length

Contents:

- Indwelling Catheter: 4 Fr. x 55 cm Radiopaque Polyurethane with Blue FlexTip®, T-Port Connector, Integral Extension Line, Extension Line Clamp, Integral Suture Wing, Contamination Guard and Placement Wire
- GlideThru™ Peel-Away Sheath: 4 Fr. x 2-3/4" (7 cm) Radiopaque over 4 Fr. Dilator
- Spring-Wire Guide: .018" (0.46 mm) dia. x 13-1/8" (33 cm) (Straight Soft Tip on
- 1: Catheter: 22 Ga. x 1-3/4" (4.45 cm) Radiopaque over 25 Ga. RW Introducer Needle
- Injection Needle: 18 Ga. x 1-1/2" (3.81 cm) 1:
- Introducer Needle: 21 Ga. x 1-1/2" (3.81 cm) TW 1:
- Introducer Needle: Echogenic 21 Ga. x 2-3/4" (7 cm) TW 1:
- 1: Syringe: 10 mL Luer-Slip
- Skin Protectant Prep Pad (non-sterile solution) 1:
- 3 mL Applicator 2% CHG and 70% IPA ChloraPrep®1 One-Step Solution with 1: Hi-Lite Orange™ Tint (non-sterile solution)
- Dust Cap: Non-Vented
- SecondSite™ Adjustable Hub: Catheter Clamp 1:
- SecondSite™ Adjustable Hub: Fastener 1:
- SharpsAway® Disposal Cup

²A registered trademark of T.J. Smith & Nephew, Limited. ³A registered trademark of C. R. Bard, Inc.

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.















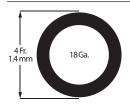


Contraindications: This device is contraindicated wherever there is presence of device related infections, previous or current thrombosis. Clinical assessment of patient must be

Not made with natural rubber latex



Store between 20 - 25°C (68 - 77°F) Avoid excessive heat above 40°C (104°F).



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

[‡] Pump flow rates are determined at maximum pump pressure of 10 psig.

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



IK-01451-107K (9/17)

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^{*} 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.