

AKK()V Peripherally Inserted Central Catheter Set with Trimmable Catheter Seldinger Technique with 130 cm Marked Wire

REF PR-07051-TLW

Contents:

- 1: Indwelling Catheter, Trimmable: 16 Ga. x 70 cm Radiopaque Polyurethane with Integral Extension Line, Extension Line Clamp, Integral Suture Wing
- GlideThru[™] Peel-Away Sheath: 5 Fr. x 4" (10 cm) Radiopaque over 1. 5 Fr. Dilator
- Spring-Wire Guide, Marked: .018" (0.46 mm) dia. x 51-3/16" 1: (130 cm) (Straight Soft Tip on One End - Straight Stiff Tip on Other)
- Introducer Needle: Echogenic 21 Ga. x 2-3/4" (7 cm) TW 1:
- 1: Syringe: 5 mL Luer-Slip
- 1: Skin Protectant Prep Pad (non-sterile solution)

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.















Not made with natural rubber latex

DuPont" Tyvek[®]



Catheter Cross Section

Lumen	Priming	Flow	Pump
	Volume*	Rate [†]	Flow Rate [‡]
	(mL)	(mL/hr)	(mL/hr)
Distal (16 Ga.)	0.76	1450	6050

Flow rates are done with normal saline, room temperature, 100 cm head height and represent approximate flow capabilities.

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

Above values are for untrimmed catheter. Trimming catheter will alter priming volumes and flow rates.

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



IR-07051-101J (7/17)



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ARROW 70^{cm}_{catheter} 0 inch dia. spring-wire

5 Fr. 70 cm cm catheter length catheter

- Dust Cap: Non-Vented Catheter Trimmer Patient ID Card
- 1: 1: Chart Sticker

1:

1:

- Patient Information Booklet 1:
- Paper Tape Measure 1:
- Dressing: STATLOCK®1 Catheter Stabilization Device 1:

¹A registered trademark of C. R. Bard, Inc.

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.