



ARROW

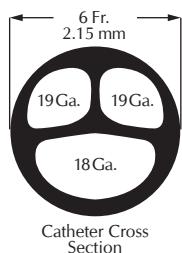
REF PR-45563-BAS

ARROW® PICC with Chlorag⁺ard® Technology

- nl** ARROW PICC met Chlorag⁺ard-technologie
Bewaren beneden 25°C. Vermijd hitte hoger dan 30°C.
- fr** PICC ARROW avec technologie Chlorag⁺ard
Conservez le produit à une température inférieure à 25°C. Veillez à ne pas exposer le produit à une chaleur excessive supérieure à 30°C.
- de** ARROW PICC mit Chlorag⁺ard-Technologie
Unter 25°C lagern. Zu hohe Temperaturen über 30°C vermeiden.
- el** Καθέτηρας ARROW PICC με Τεχνολογία Chlorag⁺ard
Φυλάσσεται κάτω των 25°C. Να αποφεύγεται η υπερβολική θερμότητα άνω των 30°C.
- it** ARROW PICC con tecnologia Chlorag⁺ard
Conservare ad una temperatura inferiore a 25°C. Evitare il calore eccessivo superiore a 30°C.
- pl** Cewnik ARROW PICC z technologią Chlorag⁺ard
Przechowywać w temperaturze poniżej 25°C. Chronić przed temperaturą powyżej 30°C.
- pt** PICC ARROW com Tecnologia Chlorag⁺ard
Conservar a temperaturas inferiores a 25°C. Evitar temperaturas superiores a 30°C.
- ru** ARROW PICC с технологией Chlorag⁺ard
Хранить при температуре ниже 25°C. Избегать нагрева свыше 30°C.
- es** PICC de ARROW con tecnología Chlorag⁺ard
Almacenar a temperaturas inferiores a 25°C. Evite temperaturas excesivas por encima de 30°C.
- sv** ARROW PICC med Chlorag⁺ard-teknologi
Förvaras vid temperatur under 25°C. Undvik höga temperatur över 30°C.
- tr** Chlorag⁺ard Teknolojili ARROW PICC
25°C'in altında saklayın. 30°C'in üzerindeki aşırı ısıdan sakının.

Contents:

- 1: Multi-Lumen Indwelling TaperFree™ CG+™ Antimicrobial/Antithrombogenic Catheter¹: 6 Fr. x 55 cm Pressure Injectable Radiopaque Polyurethane with Blue FlexTip®, T-Port Connector, Extension Line Clamps, Contamination Guard and Placement Wire
 - 1: GlideThru™ Peel-Away Sheath: 6 Fr. x 2-3/4" (7 cm) Radiopaque over 6 Fr. Dilator
 - 1: Guidewire, Nitinol, Marked: .018" (0.46 mm) dia. x 17-3/4" (45 cm) (Straight Soft Tip on One End - Straight Stiff Tip on Other)
 - 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
 - 3: Dust Cap: Non-Vented
 - 1: SecondSite™ Adjustable Hub: Fastener
 - 1: SecondSite™ Adjustable Hub: Catheter Clamp
 - 1: SharpsAway II™ Locking Disposal Cup
 - 1: Catheter Trimmer
 - 1: Safety Scalpel: #11
 - 1: Patient Information Booklet
 - 2: Paper Tape Measure
 - 1: Tourniquet
 - 1: Dressing: STATLOCK^{®2} Catheter Stabilization Device
- ¹ Licensed under US Patent Nos. 6,706,024; 6,872,195 and 7,329,412.
² A registered trademark of C. R. Bard, Inc.
- All components are CE 0086 unless otherwise noted.*



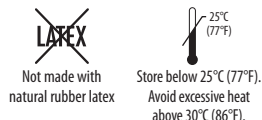
Lumen	Priming Volume* (mL)	Flow Rate† (mL/hr)	Pump Flow Rate‡ (mL/hr)	MAX Pressure Injection Flow Rate** (mL/sec)
Distal (18 Ga.)	0.61	880	5360	6
Medial (19 Ga.)	0.45	244	2090	not rated
Proximal (19 Ga.)	0.47	257	2160	not rated

* Priming volumes are approximate and are done without injection caps.
† Flow rate values are approximate and are determined using deionized water at 100 cm head height.
‡ 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.

Rx only. Warning: Read all package insert warnings, precautions and instructions prior to use. Failure to do so may result in severe patient injury or death. Warning: This product contains a chemical known to the State of California to cause cancer, birth defects or other reproductive harm.

Contraindications: Clinical assessment of the patient must be completed to ensure no contraindications exist. The Arrow® PICC with Chlorag⁺ard® Technology is contraindicated in the following areas: Patients with known hypersensitivity to chlorhexidine; In presence of device related infections; In presence of previous or current thrombosis in the intended vessel or along the catheterized vessel pathway. 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. See additional labeling for product specific contraindications.

Sterile. Single use. Fluid path components are non-pyrogenic.



Manufacturer:
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Subsidiary of Teleflex Incorporated
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