Dimensions:
Width: 8-1/2”
Length: 11

PB-40002-100B

ARROW

REF **PR-35052-HPHNM** (IPNXXXXXX)

5 Fr. **2** Lumen **50** cm catheter length **.018** inch dia. spring-wire guide

Pressure Injectable Two-Lumen PICC with 80 cm Wire

Contents:

1: Two Lumen TaperFree® Catheter: 5 Fr. (1.81 mm OD) x 50 cm, Pressure Injectable, Blue FlexTip® and Contamination Guard origin:xxxxxxxxxxxxxxxxxxxxxx
1: GlideThru™ Peel-Away Sheath: 5 Fr. x 4" (10 cm) Radiopaque over 5 Fr. Dilator origin:xxxxxxxxxxxxxxxxxxxxx
1: 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) origin:xxxxxxxxxxxxxxxxxxxxx
1: Guidewire, Marked: .018" (0.46 mm) dia. x 80 cm Hydrophilic-Coated Nitinol with Soft Tip Tungsten Coil (CE 0120) origin:xxxxxxxxxxxxxxxxxxxxx
1: Introducer Needle: Echogenic 21 Ga. x 2-3/4" (7 cm) TW origin:xxxxxxxxxxxxxxxxxxxxx
1: Syringe: 10 mL Luer-Lock origin:xxxxxxxxxxxxxxxxxxxxx
2: Dust Cap: Non-Vented origin:xxxxxxxxxxxxxxxxxxxxx
1: SecondSite™ Adjustable Hub: Fastener origin:xxxxxxxxxxxxxxxxxxxxx
1: SecondSite™ Adjustable Hub: Catheter Clamp origin:xxxxxxxxxxxxxxxxxxxxx
1: Catheter Trimmer origin:xxxxxxxxxxxxxxxxxxxxx
1: Safety Scalpel: #11 origin:xxxxxxxxxxxxxxxxxxxxx
1: Patient ID Card origin:xxxxxxxxxxxxxxxxxxxxx
1: Chart Sticker origin:xxxxxxxxxxxxxxxxxxxxx
1: Patient Information Booklet origin:xxxxxxxxxxxxxxxxxxxxx
1: Checklist/CLIP Sheet origin:xxxxxxxxxxxxxxxxxxxxx
1: Paper Tape Measure origin:xxxxxxxxxxxxxxxxxxxxx
1: Dressing: STATLOCK®[1] PICC Plus Catheter Stabilization Device origin:xxxxxxxxxxxxxxxxxxxxx

[1]A registered trademark of C. R. Bard, Inc.
All components are CE 2797 unless otherwise noted.

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. www.teleflex.com/IFU
Not made with natural rubber latex.
Fluid path components are non-pyrogenic.
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.

Lumen	Priming Volume* (mL)	Gravity Flow Rate† (mL/hr)	Pump Flow Rate†† (mL/hr)	MAX Pressure Injection Flow Rate** (mL/sec)
Distal (18 Ga.)	0.48	488	3860	5
Proximal (18 Ga.)	0.5	488	3870	5

* Priming volumes are approximate and are done without accessories.
† 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.

EU Authorized Representative and Importer:
EC REP Teleflex Medical
IDA Business and Technology Park
Dublin Road, Athlone
CE 2797 Co. Westmeath, Ireland

All components are CE 2797 unless otherwise noted.

MD

STERILE EO

LATEX

CAS # 7440-48-4

LOT LotNumber

YYYY-MM-DD

YYYY-MM-DD

(01)XXXXXXXXXXXXXX
(17)YYMMDD
(11)YYMMDD
(10)LotNumber

Arrow International LLC
Subsidiary of Teleflex Incorporated
3015 Carrington Mill Blvd.
Morrisville, NC 27560 USA

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