WARRANTY: B. BRAUN INTERVENTIONAL SYSTEMS INC. WARRANTS THAT THIS PRODUCT WAS MANUFACTURED ACCORDING TO APPLICABLE STANDARDS AND SPECIFICATIONS. PATIENT CONDITION, CLINICAL TREATMENT, AND PRODUCT MAINTENANCE MAY AFFECT THE PERFORMANCE OF THIS PRODUCT. USE OF THIS PRODUCT SHOULD BE IN ACCORDANCE WITH THE INSTRUCTIONS PROVIDED AND AS DIRECTED BY THE PRESCRIBING PHYSICIAN.

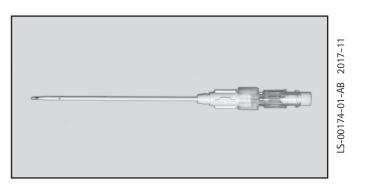
Interventional Systems B BRAUN



Manufacturer: pfm medical, inc. 1916 Palomar Oaks Way, Suite 150 Carlsbad, CA 92008 Distributed by: **B. Braun Interventional Systems Inc.**824 Twelfth Avenue
Bethlehem, PA 18018
www.bisusa.org

ACCEL® Centesis Catheter Percutaneous Fluid Drainage Catheters

INSTRUCTIONS FOR USE



Rx only

Do not reuse

Do not use

if package is

damaged

Store at room temperature

Do not

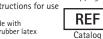
resterilize







Consult instructions for use



Not made with natural rubber latex



Keep away





number

STERILE EO Sterilized using ethylene oxide

DEHIP

Not made

with DEHP

Interventional Systems

B BRAUN

Distributed by:

B. Braun Interventional Systems Inc. 824 Twelfth Avenue Bethlehem, PA 18018

www.bisusa.org

Customer Service, ordering TEL: (877) 836-2228 FAX: (610) 849-1334

Technical Support TEL: (800) 443-VENA (8362)

Made in U.S.A.

INDICATIONS FOR USE:

The ACCEL® Centesis Catheter is for short term percutaneous fluid drainage.

CONTRAINDICATIONS:

This product should only be used by qualified personnel who are familiar with the technique. During insertion avoid contact with bone, cartilage, and scar tissue which can damage the catheter tip.

CAUTION:

- For Single Patient Use Only. Do not attempt to clean or resterilize this product.
- After use, this product may be a potential biohazard.
- Dispose in accordance with applicable laws and regulations.
- Handle in a manner that will prevent contamination.
- Do not use a device that has been damaged.
- Do not use any device if the package is opened or damaged.

WARNING:

- This product contains chemicals known to the state of California to cause cancer and birth defects or other reproductive harm.
- Federal Law (USA) restricts this device to sale by or on the order of a physician.

NOTE:

These instructions for the ACCEL Centesis Catheter are not meant to define or suggest any medical or surgical technique. The individual physician is responsible for the proper procedure and techniques to be used with this product.

| French Size | Needle O.D. | Sheath Length | Catheter O.D. | Catheter I.D. |
|----------------|----------------|------------------|------------------|------------------|
| 4 | 20g (.09 mm) | 7cm | 1.4mm | 1.0mm |
| 4 | 20g (.09 mm) | 10cm | 1.4mm | 1.0mm |
| 5 | 19g (1.1 mm) | 7cm | 1.7mm | 1.2mm |
| 5 | 19g (1.1 mm) | 10cm | 1.7mm | 1.2mm |
| 5 | 19g (1.1 mm) | 15cm | 1.7mm | 1.2mm |

PROCEDURE

(Single step catheterization through a skin incision)

- 1. Perform a skin incision under local anesthesia.
- 2. Introduce the ACCEL Centesis Catheter into the cavity using ultrasound guidance, CT or fluoroscopy.
- 3. Unlock the needle locking collar and remove the needle from the catheter.
- 4. Connect a syringe or suction tube to the catheter hub.

ORDERING INFORMATION

| Reference Number | Description | Quantity |
|---------------------|--|------------|
| 613210 | ACCEL Centesis Catheter 4Fr, 7cm Length | 5 per pack |
| 613211 | ACCEL Centesis Catheter 4Fr, 10cm Length | 5 per pack |
| 613212 | ACCEL Centesis Catheter 5Fr, 7cm Length | 5 per pack |
| 613213 | ACCEL Centesis Catheter 5Fr, 10cm Length | 5 per pack |
| 613214 | ACCEL Centesis Catheter 5Fr, 15cm Length | 5 per pack |