Clik-FIX® Peripheral Catheter Securement

REF PCSK-1011B

DESCRIPTION - The B. Braun Medical Clik-FIX Peripheral Catheter Securement Device is an integrated securement device (ISD) designed to protect the insertion site, cushion under the hub, and secure the



catheter. The Clik-FIX Peripheral device is compatible with most catheters with a Luer lock connection.

INDICATIONS - The Clik-FIX Peripheral Catheter Securement Device is indicated for use with most intravascular catheters with a Luer lock connection.

CONTRAINDICATIONS - Do not use the device on compromised skin (i.e., open wounds, burns, lesions, rash, or other irritation and/or patients with known tape or adhesive allergies or hypersensitive skin).

ADVERSE REACTIONS - Skin irritation.

- WARNING AND PRECAUTIONS
- Follow facility protocol when placing and removing the device.
 Observe universal precautions.
- Do not use a Clik-FIX device on patients who are allergic to adhesives and on non-adherent skin.
- Do not apply alcohol, acetone, or other liquid agent directly on the Clik-FIX device when in use, as it can weaken the device.
- During application and removal of the Clik-FIX device, take care to minimize manipulation and movement of the catheter.
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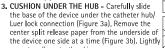
 Prepare the skin surface according to facility protocol. Remove oil and moisture from application area and allow the area to
- completely dry.

 Routinely inspect the Clik-FIX device, insertion site, and catheter position according to facility protocol and best practice quidelines.
 - Replace the Clik-FIX® device at least every 7 days or per facility protocol.

B. Braun Medical Inc. Bethlehem, PA 18018-3524 USA **B** BRAUN

INSTRUCTIONS FOR USE:

- 1. PREP Prepare and cleanse insertion site and surrounding securement area according to facility protocol. Allow the area to completely dry. Establish vascular access according to facility protocol or remove previous dressing from catheter.
- 2. PROTECT THE INSERTION SITE Carefully remove the release paper from the horseshoe window portion of the device (Figure 2a). While holding the base of the device, ensure exposed adhesive is facing the patient's skin. Align the transparent dressing so the insertion site is covered and the Luer lock connection is centered under the foam base pad (Figure 2b). The bordered bio-occlusive dressing should completely cover the entry site and surrounding





the Luer lock connection (Figure 4). Lightly press to adhere. Carefully remove the other half of the release papers from the securement strap and lightly press to adhere. The extra foam tape strip can be used to secure the extension tubing and/or provide additional securement. Date and time per facility protocol.

REMOVAL: Follow your facility's protocol.

Sterile when packaged in a sealed sterile pouch or when included as a component in a sterile kit.

Not made with Consult Instructions for Use natural rubber www.bbraunusa.com Pat.: www.bbraunusa.com/patents

Single.

latex or phthalates, A4809167-2

Do not use if

package is

Rx only

damaged

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