en English — Non-Coring Needle

This product is indicated for the administration into or withdrawal of fluids from implanted ports. It is recommended that this product be changed in accordance with U.S. Centers for Disease Control (CDC) guidelines for administration sets, local or country specific guidelines, professional standards of practice, and/or according to your institution's policy. Do not use this product if it appears damaged or if the package has been previously opened or damaged. This product contains a nonvented cap.

Do Not Reuse: Medical devices require specific material characteristics to perform as intended. These characteristics have been verified for single use only. Any attempt to reprocess the device for subsequent re-use may adversely affect the integrity of the device or lead to deterioration in performance.

WARNINGS:

- Verify needle length is correct for portal/patient; if too long, needle and/or portal may be damaged at insertion; if too short, needle may not completely pierce portal septum, and medication may be delivered into surrounding tissue and/or needle may be blocked.
- Fully tighten Y-site injection cap before each use.
- Y-site septum: Failure to attach an appropriate needleless device after removing the injection cap can result in air embolism.

CAUTIONS:

- Do not use with high pressure or with power injectors. Pressure above 45 psi may cause damage and leakage.
- **Follow standard infection control precautions as specified by the CDC (USA) or local equivalent.**

Use aseptic technique. Refer to the illustrations when preparing and using this device.

1. Prepare the portal site for sterile needle insertion ❶.
2. Remove the needle guard and flush the set ❷. If appropriate, remove injection cap from luer lock Y-site and attach appropriate needleless device.

   WARNING: Failure to attach an appropriate needleless device after removing the injection cap can result in air embolism.
3. Grasp the GRIPPER® tab and insert the needle through the skin and portal septum at a 90° angle to the septum. **Remove the GRIPPER® tab ❸.**
4. Apply a semi-permeable dressing over the GRIPPER® base, ensuring that a minimum 4 cm area surrounding the base is covered ❹.

Dispose of used needles in a sharps container, in accordance with CDC guidelines (USA) and/or your institution's bloodborne pathogens program.

Additional Instructions for Use can be printed at www.smiths-medical.com

fr Français — Aiguille Atraumatique

Ce produit est indiqué pour l'administration ou le prélèvement de fluides à partir de chambres implantables. Il est recommandé de remplacer ce produit conformément aux recommandations du Centre de lutte contre les maladies (CDC - USA) ou à la politique en vigueur dans votre établissement. Ne pas utiliser le produit s'il apparaît endommagé ou si l'emballage a été initialement ouvert ou endommagé. Ce produit est doté d'un bouchon non ventilé.

Ne pas réutiliser: Les dispositifs médicaux requièrent des caractéristiques matérielles spécifiques pour être utilisés comme prévu. Ces caractéristiques ont été avérées uniquement en cas d'usage unique. Toute tentative de retraitement du dispositif en vue d'une réutilisation peut gravement compromettre