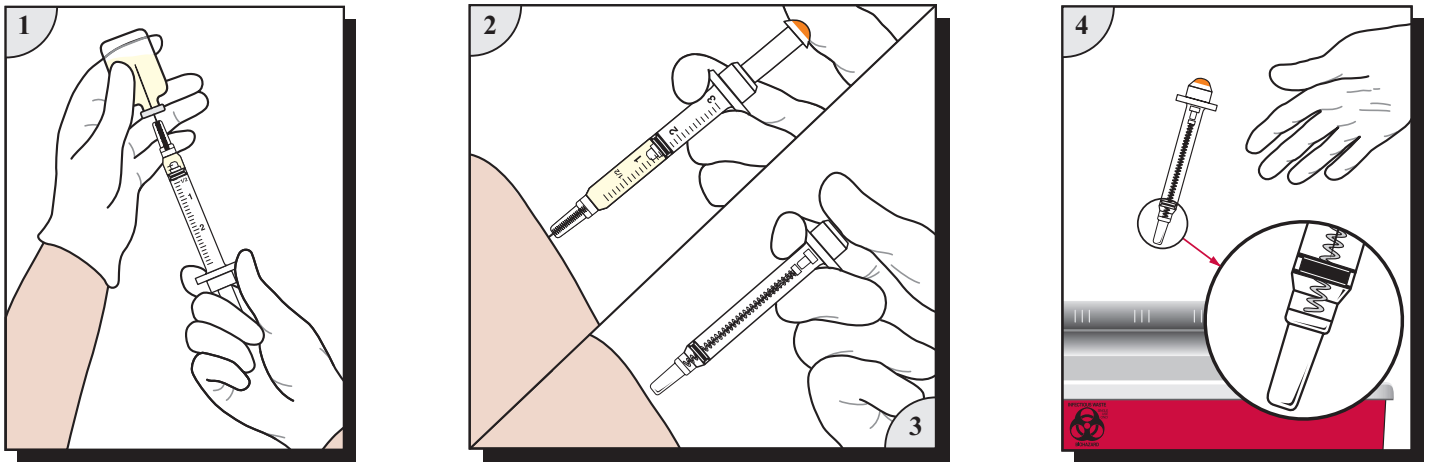
CE 0086

VANISHpoint®

Syringe

Standard Draw Procedure | In-Patient, Automated Retraction | Non-Reusable



Product Usage Information:

1. Prepare and give injection using aseptic technique according to institutional policy.
2. For injection into patients, continue depressing plunger to activate automatic needle retraction ***while needle is still in patient.*** For injection into IV ports, continue depressing plunger to activate automatic needle retraction and ***immediately remove needle from port. Full dose is administered only when needle retraction is activated.***
3. Needle will automatically retract into syringe, preventing exposure to contaminated needle and rendering syringe non-reusable. In the event that needle retraction mechanism does not activate, discard syringe in an appropriate sharps container per protocol of institution. Do not recap contaminated needles.
4. Dispose of VanishPoint® syringe in an appropriate sharps container per protocol of institution.

Precautions:

- Single use only. Reuse of this device may result in exposure to bloodborne pathogens, including Hepatitis B virus (HBV), Hepatitis C virus (HCV), and human immunodeficiency virus (HIV).
- Contents are sterile, non-toxic, and non-pyrogenic. Do not use if product or package is damaged.
- Not made with natural rubber latex.
- Use only with attached needle. Needle cannot be changed.
- Automated needle retraction occurs only when barrel is emptied and plunger is fully depressed.
- For applications where full dose is not administered, expel remaining contents according to institutional policy and activate needle retraction.
- U.S. Federal Law restricts this device to sale by or on the order of a physician.

RETRACTABLE TECHNOLOGIES, INC.®
www.vanishpoint.com

511 Lobo Lane
Little Elm, Texas 75068-0009
Tel: (972) 294-1770 Fax: (972) 294-4400
Toll Free 1-888-703-1010

M001 REV. 9 EN, JUL 2013