

VanishPoint® I.V. Catheter

Automated Retraction Technology

VanishPoint® IV-Katheter

Automatisierte Retraktionstechnologie

Cathéter intraveineux VanishPoint®

Technologie de Rétraction Automatisée

Cateter endovena VanishPoint®

Tecnologia di Retrazione Automatizzata

Catéter I.V. Vanishpoint®

Tecnologia de Retracción Automatizada

VanishPoint® I.V. Katheter

Geautomatiseerde Retractietechnologie

Cateter IV VanishPoint®

Tecnologia de Retração Automática

VanishPoint® I.V. カテーテル

自動引き込み技術

VanishPoint® I.V. Kateter

Otomatik Geri Çekme Teknolojisi

Cewnik dożylny VanishPoint®

Tecnologia Automatycznego Wycofywania

Intravenøst VanishPoint® - kateter

Automatiseret Retraktionsteknologi

VanishPoint® I.V.-Kateter

Teknologi för Automatisk Tillbakadragning av Nål

REAL SAFETY™

Rx Only

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SINGLE USE

STERILE EO
STERILIZED BY
ETHYLENE OXIDE

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PATENTS--
WWW.RETRACTABLE.COM/PATENTS

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VanishPoint® I.V. Catheter

Automated Retraction Technology

Description:

- Contents are sterile, non-toxic, and non-pyrogenic.
- The VanishPoint® I.V. catheter is designed to minimize exposure to the contaminated introducer needle and prevent needlestick injuries. Upon activation, the introducer needle is automatically retracted into the needle housing.
- Color coding for the catheter gauge can be found on the catheter hub and at the end of the needle housing. VanishPoint® I.V. catheters are color coded according to international standards.

GAUGE	14G	16G	18G	20G	22G	24G
COLOR	Orange	Medium Grey	Deep Green	Pink	Deep Blue	Yellow

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).
- Do not use if product or package is damaged.
- Not made with natural rubber latex.
- To prevent exposure to contaminated needle, activate safety mechanism prior to withdrawing needle from catheter.
- Do not use scissors or sharp implements near I.V. catheters.
- Never reinsert the introducer needle into the catheter.
- Clinicians must follow instructions for use and be trained on proper product use.

- During catheter insertion, maintenance, replacement, and removal, follow current “Guidelines for the Prevention of Intravascular Catheter-Related Infections”, Centers for Disease Control and Prevention, Atlanta, GA.
- Notify physician immediately and follow facility protocols in the event patients exhibit symptoms of infusion-related complications.

Indications:

- For access to a peripheral vein or artery.
- May be used for any patient population with consideration given to patient size, infusate, and duration of therapy.
- 18-24 gauge I.V. catheters are suitable for use with power injectors up to 325 psi.

Maximum Flow Rates with High Pressure Injection (up to 325 psi)

	24G	22G	20G	18G
Maximum Flow Rate (mL/sec) with Omnipaque™ 300 at Room Temperature	5.0	6.3	7.1	7.7
Maximum Flow Rate (mL/sec) with Omnipaque™ 350 at Room Temperature	3.4	4.3	5.3	6.3

Instructions for Use:

- Select and prepare site using aseptic technique according to institutional policy. Based on patient allergy information, select an antiseptic solution to disinfect the insertion site. Recommended solutions include: 2% chlorhexidine, 10% povidone-iodine, 70% alcohol.
 - Remove needle cover and inspect catheter unit.
1. Perform venipuncture, observing for blood return. Advance the catheter slightly to assure full catheter entry into the vein.
 2. Holding the flashback chamber stationary, disengage the hub from the introducer needle and advance the catheter into the vein. **DO NOT REINSERT THE NEEDLE INTO THE CATHETER AT ANY TIME.**
 3. **Before withdrawing the introducer needle from the catheter**, depress the color coded end of the needle housing to activate the automated retraction of the introducer needle. To activate the retraction mechanism using one hand, stabilize the color coded end of the needle housing against the proximal portion of the palm, while pulling back the housing using the finger grips.
 4. The introducer needle will retract into the housing. In the event that needle retraction mechanism does not activate, discard catheter introducer in an appropriate sharps container per protocol of institution. Do not recap contaminated needles.

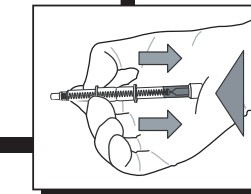
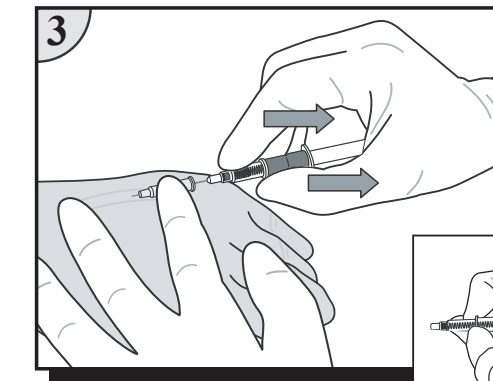
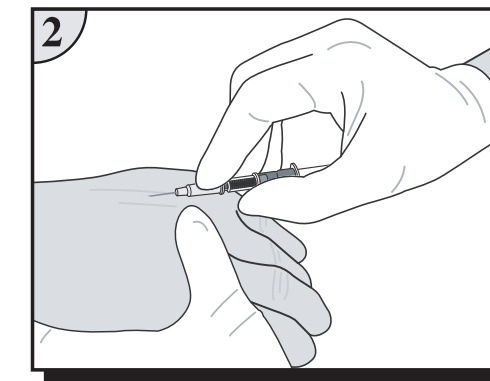
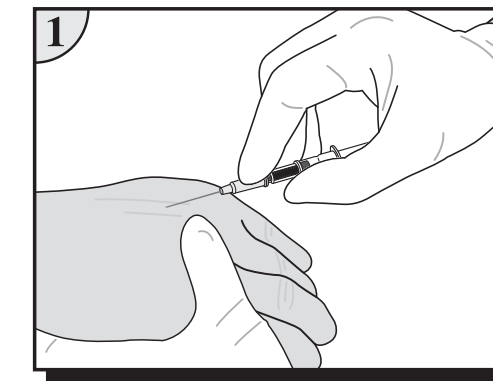
VanishPoint® IV-Katheter

Cathéter intraveineux VanishPoint®

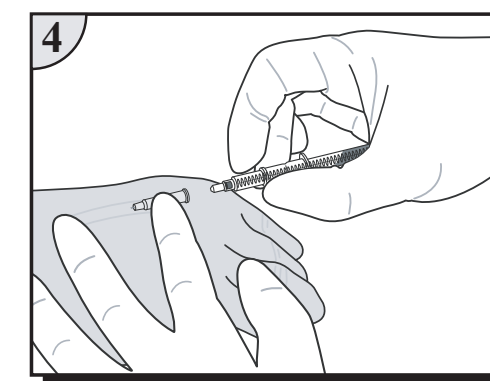
Cateter endovena VanishPoint®

Catéter I.V. VanishPoint®

- Dispose of the introducer needle in an appropriate sharps container per protocol of institution.
- Securely connect tubing or adapter to the catheter hub, following the manufacturer’s instruction for the device.
- Secure catheter and dress site per protocol of institution.



View from underside of palm



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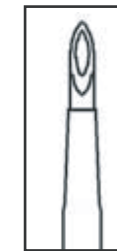
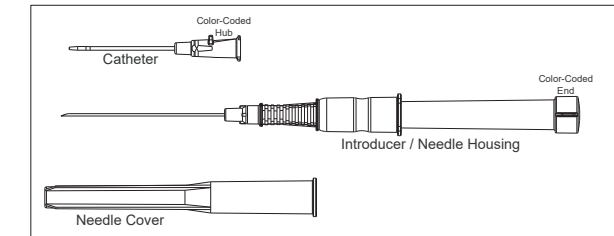
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Distal End Configuration

SYMBOL KEY

	EN - Do not use if package is damaged DE - Nicht verwenden, wenn die Verpackung beschädigt ist FR - Ne pas utiliser si l'emballage est endommagé IT - Non utilizzare se la confezione è danneggiata ES - No utilizar si el producto está dañado NL - Niet gebruiken als de verpakking is beschadigd PT - Não utilize se a embalagem estiver danificada JP - 包装が破損している場合は使用しないでください TR - Ambalajı zarar görmüşse kullanmayınız PL - Nie używać, jeżeli opakowanie jest uszkodzone DK - Må ikke anvendes, hvis emballagen er beskadiget SE - Får ej användas om förpackningen är skadad		EN - Batch Code TR - Parti Kodu
	EN - Sterilized by Ethylene Oxide TR - Etilen oksit ile sterilize edilmiştir		EN - Use by Date TR - Son Kullanma Tarihi
	EN - EU Authorized Representative TR - AB Yetkili Temsilcisi		EN - Catalogue Number TR - Katalog Numarası
	EN - Single Use TR - Tek Kullanımlıktır		EN - Manufacturer TR - Üretici
	EN - See Instructions for Use TR - Kullanım talimatlarına bakınız		EN - Date of Manufacture TR - Üretim Tarihi
	EN - Rx Only TR - Sadece Reçeteye Satılır		

