



Vented Vial Adapter Device

Please refer to the device instructions for use (IFU) for complete set of directions for the device.

Intended Use/Indications for Use:

The Vented Vial Adapter device is intended for transfer and mixing of drugs contained in a vial.

Contraindications:

Currently no specific contraindications are known for the Vented Vial Adapter.

Cautions/Precautions/Warnings:

- In case that diluent is not transferred
 - o Do not remove the vial adapter from the package
 - o Do not use the vial adapter if the package is damaged
 - o Do not use the vial adapter if it comes out of package
- Federal (USA) law restricts this device to sale by or on the order of a physician.
- This device is intended to be used for one patient and one application only.
- Single use only.
- Reuse compromises safety and efficacy - it may cause contamination due to loss of sterility.
- Re-sterilization may damage the device.
- Contents are sterile and non-pyrogenic.
- Do not remove the device from vials after use.
- Dispose of used device in accordance with applicable regulations.
- Do not use if package is damaged.

Registration Information

Information herein is for use only in countries with applicable health authority registrations. The Vented Vial Adapter is 510(k) cleared by the United States Food and Drug Administration and carries the CE mark (0344). Products are shown for INFORMATION purposes only and may not be approved for marketing in specific regions. Please contact your West Pharmaceutical Services, inc. (West) representative for product availability.

West and the diamond logo are registered trademarks of West Pharmaceutical Services, Inc. in the U.S. and other jurisdictions.

Date: 09-December-2021

Number: GN-REG-192

Title: Vented Vial Adapter Device - Indications, Safety and Registration Zones

Revision: 1.0