

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 Vial Adapter device is intended for the transfer and mixing of drugs contained in vials.

Contraindications:

Currently no specific contraindications are known for the Vial Adapter.

Cautions/Precautions/Warnings:

- Don't use product if fell 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.
- Contents are sterile and non-pyrogenic.
- Reuse compromises safety and efficacy it may cause contamination due to loss of sterility.
- Re-sterilization may damage the device
- Do not remove the device from vials after use.
- Do not use if package is damaged / Do not use if not intact.
- Dispose of used device in accordance with applicable regulations.

Registration Information

Information herein is for use only in countries with applicable health authority registrations. The 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-191

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

Revision: 1.0