



Medical Devices, Inc.
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HalochestSeals.Com



Medical Device



Do Not Reuse

-34°C
-30°F



Designated Temperature

60°C
140°F



Not Sterilized



Do Not Use if Package is Damaged or Open



Not Manufactured with Natural Rubber Latex



Instructions Available



Translation



[Http://WWW.HALOCHESTSEALS.com/IFU](http://WWW.HALOCHESTSEALS.com/IFU)

Product Description:

Halo Seal and Halo XL consists of an adhesive hydrogel layer and an adhesive, non-latex, hypoallergenic, backing layer. It is impermeable to liquids and seals the wound until the patient can be transported to a Health Care Facility.

Indications for Usage:

Halo Products are designed to aid treatment of thoracic wounds by using standard dressing application techniques that aid Civilian or Emergency Responders. Designed to cover and protect wounds. Do not use the dressing as a replacement for sutures and other primary wound closure methods. Halo Products are a single use product. Attempts to reuse a Halo Products may introduce contamination and infection to a secondary patient. No indications of usage, safety, risk, or efficacy are established for post surgical applications.

Warnings

Ensure the Halo completely seals the perimeter of the wound to prevent air, liquids, and debris entering. Persons who have thoracic injuries should be immediately transported to a healthcare or trauma facility for evaluation by a Clinician for post care and treatment.

Application

- Clean and dry wound.
- Remove from release liner. Do not place on secondary surface before application to prevent contamination.
- With Gel side down, place over center of wound.
- Use light pressure - Do not stretch during application.
- Use additional Halo Seals, if other wounds are present.



Application Video

Transport Patient to Emergency Care Facility for Post Treatment and Evaluation

Dispose after use, per local medical device waste guidelines.

Contact Information



Manufacturer : Medical Devices , Inc.



Please contact or report any serious incidents regarding this product to the manufacturer and regulatory authorities.

NB#
2862

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CLEAN AND DRY WOUND



REMOVE PRODUCT FROM PACKAGING. REMOVE RELEASE LINER



APPLY PRODUCT WITH LIGHT PRESSURE



Transport Patient to Emergency Care Facility for Evaluation and Post Treatment

