

Antimicrobial Biofilm Wound Gel

Powered by XBio™ Technology

Description

BLASTX Antimicrobial Biofilm Wound Gel has antimicrobial properties that inhibit the growth of microorganisms. BlastX disrupts biofilm structures while maintaining a moist wound environment that is conducive to wound healing. BLASTX is a white and virtually odorless product.

BLASTX contains benzalkonium chloride 0.13%, macrogol 400, macrogol 3350, sodium citrate, citric acid, water.

Indications for use

BLASTX is intended for the management of wounds such as stage I-IV pressure ulcers, partial and full thickness wounds, diabetic foot and leg ulcers, post-surgical wounds, first and second degree burns, grafted and donor sites.

Contraindications

BLASTX should not be used if there is hypersensitivity to any of the ingredients.

Warnings and precautions

For external topical use only. Do not ingest. Do not cover with alginate dressings. Potential burning or discomfort may occur. Benzalkonium chloride may irritate the skin. Must not be used in or near the eyes. BLASTX can be used during pregnancy and in the lactation period under medical supervision. Do not apply BLASTX to skin of the breast during lactation, because the baby may ingest the product. BLASTX should not be used if there is a history of allergy to any of the ingredients. No specific drug interaction studies have been undertaken; therefore use with other topical medications is not recommended.

Instructions for use

BLASTX should be applied directly to the wound and then covered with an appropriate dressing. Cover entire wound surface with gel at a minimum thickness of 3 mm. After opening, BlastX may be used until the labelled expiration date.

Storage conditions

BLASTX should be stored between 15°C and 25°C.

Manufacturer

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Distributed by

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