BACITRACIN ZINC- bacitracin zinc ointment Trifecta Pharmaceuticals Usa Llc

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Bacitracin ZINC OINTMENT USP

Drug Facts

Active ingredient Bacitracin Zinc 500 units

Purpose

First Aid Antibiotic

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

Uses

• first aid to help prevent infection in • minor cuts • scrapes • burns

Warnings

For external use only.

Do not use • in the eyes • over large areas of the body • if you are allergic to any of the ingredients • longer than 1 week unless directed by a healthcare professional

Ask a healthcare professional before use in care of deep or puncture wounds, animal bites, or serious burns

Stop use and ask a healthcare professional if • the condition persists or gets worse • a rash or allergic reaction develops

Directions

• clean the affected area

• apply a small amount of this product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily

• may be covered with a sterile bandage

Inactive ingredient:

aloe vera gel, light mineral oil, white petrolatum

Other information

• store between15° to 25°C (59° to 77°F) • Lot No & Expiration Date: See box and tube crimp.

100% GUARANTEED

Distributed by:

Trifecta Pharmaceuticals USA™

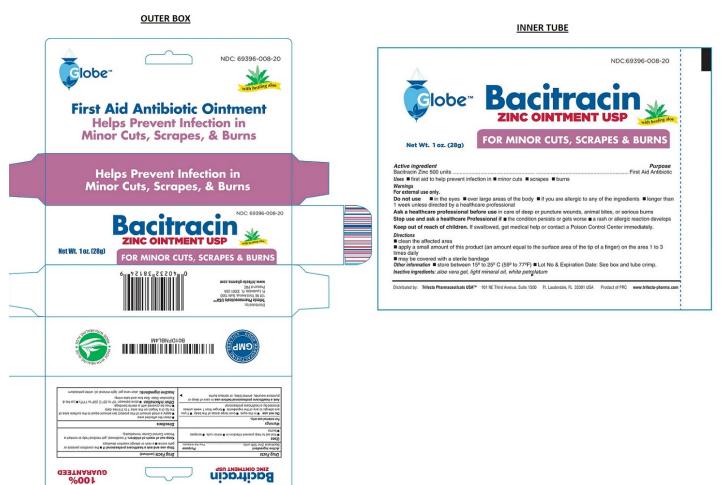
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Product of PRC

www.trifecta-pharma.com

Packaging



BACITRACIN ZINC bacitracin zinc ointment Product Information Product Type HUMAN OTC DRUG Route of Administration TOPICAL Active Ingredient/Active Moiety

	Ingredient Name	Basis of Streng	gth Strength
BACITRACIN ZINC (UNII: 89 Y4M234ES) (BACITRACIN - UNII:58 H6 RWO52I) BACITRACIN		500 [USP'U] in 1 g	
Inactive Ingredier	its		
Ingredient Name			Strength
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)			
LIGHT MINERAL OIL (UNII: N6K5787QVP)			
PETROLATUM (UNII: 4T6H12BN9U)			
Packaging # Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:69396-008-20	1 in 1 BOX	04/07/2016	
1	28 g in 1 TUBE; Type 0: Not a Combination Product		
Marketing Info	rmation		
Marketing Info		Marketing Start Date	Marketing End Date
Marketing Info Marketing Category OTC monograph final	rmation Application Number or Monograph Citation part333B	Marketing Start Date 04/07/2016	Marketing End Date

Labeler - Trifecta Pharmaceuticals Usa Llc (079424163)

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