

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.

Globe Bacitracin Zinc Ointment

Active Ingredient

Bacitracin Zinc 500 Units

Purpose

First Aid Antibiotic

Uses

First Aid to help prevent infection in

- minor cuts
- scrapes
- burns

Warnings

For External Use Only.

Do Not Use

- in eyes
- over large areas of the body
- if you are allergic to any of the ingredients

Stop Use and Ask Doctor if

- Before use in case of deep puncture wounds, animal bites or serious burns
- The condition persists or gets worse
- A rash or allergic reaction develops

Directions

- Clean the affected area and dry thoroughly.
- Apply a small amount of the 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

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center Immediately.

Inactive Ingredients

Mineral oil, white petrolatum

Questions

Call 1-888-296-9067

Fax: 1-888-878-3609

Online: www.trifecta-pharma.com

Reorder No. 1013SL

Storage Information

Store at room temperature 15° to 25°C (59° to 77°F).

Other Information

Distributed By:

Trifecta Pharmaceuticals USA®

Ft. Lauderdale, FL. 33301 USA

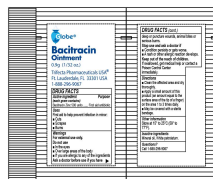
www.trifecta-pharma.com

Packaging

OUTSIDE BOX



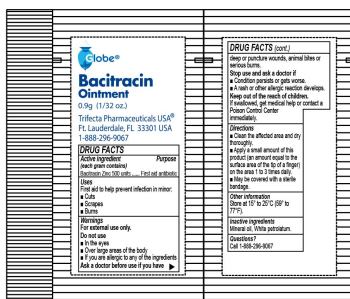
INNER PACKET



OUTSID BOX



INNER PACKET



BACITRACIN ZINC
bacitracin zinc ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69396-104
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [USP'U] in 1 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	
MINERAL OIL (UNII: T5L8T28FGP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69396-104-09	144 in 1 CARTON	04/21/2020	
1		0.9 g in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:69396-104-25	25 in 1 CARTON	03/14/2022	
2		0.9 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	04/21/2020	

Labeler - Trifecta Pharmaceuticals USA LLC (079424163)

Revised: 9/2023

Trifecta Pharmaceuticals USA LLC