

BACITRACIN ZINC- bacitracin zinc ointment
Chain Drug Marketing Association Inc.

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.

Quality Choice 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
- longer than 1 week unless directed by a doctor

Ask Doctor before Use

In care of deep puncture wounds, animal bites, or serious burns.

Stop Use and Ask Doctor 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 Ingredients

Aloe barbadensis leaf juice, mineral oil, petrolatum

Keep out of reach of children

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

Other Information

Store Between 15° to 25°C (59° to 77°F)

Lot No. and Exp. date: see box and tube crimp

Distributed By:

C.D.M.A, Inc. ©

43157 W 9 Mile Rd

Novi, MI. 48375

www.qualitychoice.com

1-800-935-2362

Product of PRC

Packaging

OUTER BOX



INNER TUBE



BACITRACIN ZINC
bacitracin zinc ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-574
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
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
PETROLATUM (UNII: 4T6H12BN9U)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-574-01	1 in 1 BOX	06/11/2020	
1		28.3 g in 1 TUBE; 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	06/11/2020	

Labeler - Chain Drug Marketing Association Inc. (011920774)**Registrant** - Trifecta Pharmaceuticals USA LLC (079424163)