

BACITRACIN ZINC- bacitracin zinc ointment

A-S Medication Solutions

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

Drug Facts

Active ingredient (each gram contains)

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 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 doctor

Ask a doctor before use

- on deep or puncture wounds, animal bites, or serious burns

Stop use and ask a doctor if

- condition persists or gets worse
- a rash or other allergic reaction develops

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

Directions

- clean the affected area and dry thoroughly
- 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

Other information

- To open: unscrew cap, pull tab to remove foil seal
- Store at 20° - 25°C (68° - 77°F)
- see carton or tube crimp for lot number and expiration date

Inactive ingredients

mineral oil, white petrolatum

Questions?

Call **1-866-923-4914**

Distributed by:

**Taro Pharmaceuticals
U.S.A., Inc.**

Hawthorne, NY 10532

HOW SUPPLIED

Product: 50090-4706

NDC: 50090-4706-0 15 g in a TUBE / 1 in a CARTON

Bacitracin Zinc



BACITRACIN ZINC

bacitracin zinc ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50090-4706(NDC:51672-2075)
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
mineral oil (UNII: T5L8T28FGP)	
petrolatum (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50090-4706-0	1 in 1 CARTON	11/11/2019	
1		15 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	05/11/2006	

Labeler - A-S Medication Solutions (830016429)

Establishment

Name	Address	ID/FEI	Business Operations
A-S Medication Solutions		830016429	RELABEL(50090-4706)

Revised: 2/2023

A-S Medication Solutions