BACITRACIN ZINC- bacitracin zinc ointment
Kroger Company

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 Oint USP Kroger part333B

Drug Facts

Active Ingredient (in each gram)

Bacitracin zinc, USP 500 units

Purpose

First aid antibiotic

Uses

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

Warnings

For external use only

Allergy Alert: •do not use if allergic to any of the ingredients

Do not use •in the eyes •over large areas of the body

Ask a doctor before use if you have •deep or puncture wounds •animal bites •serious burns

When using this product •do not use longer than 1 week unless directed by a doctor

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 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

Other information

•Store at room temperature 59°-77°F (15°-25°C). •Before using any medication, read all label directions. Keep carton, it contains important information.

Inactive ingredient white petrolatum

Questions? 1-800-632-6900

PRINCIPAL DISPLAY PANEL

Kroger®

NDC 30142-501-56
Bacitracin Zinc Ointment, USP
First Aid Antibiotic

Prevents Infection in Minor Cuts, Scrapes, and Burns.

• Prevents Infection in Minor Cuts, Scrapes, and Burns.

Net WT 1 OZ (28g)

BACITRACIN ZINC

bacitracin zinc ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:30142-501
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30142-501-56	1 in 1 CARTON	02/23/2012	
1		28 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	02/23/2012	

Labeler - Kroger Company (006999528)

Registrant - Teva Pharmaceuticals USA, Inc. (001627975)

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