CURETECH BACITRACIN FIRST AID- bacitracin zinc ointment
Curetech Skincare

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.

Curetech Bacitracin Zinc

Active Ingredient (in each gram)

Bacitracin Zinc (equal to 500 bacitracin 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

■ Not for opthalmic use **Do not use** ■ in the eyes ■ if you are allergic to any of the ingredients ■ over large areas of the body longer than 1 week unless directed by a doctor

Ask a doctor before use in case of

■ deep or puncture wounds

■ animal bites ■ serious burns

Stop use and ask a doctor if

■ the condition persists or gets worse

■ a rash or other allergic reaction develops

Keep out of reach of children

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

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

Other Information

■ store at room temperature 15-30C (59-86F)

■ avoid excessive heat and humidity

Inactive Ingredients

hard paraffin, liquid paraffin, white soft paraffin, lanolin

Package Label

CURETECH BACITRACIN FIRST AID

bacitracin zinc ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73622-3035
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**
PARAFFIN (UNII: I9O0E3H2ZE)	
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	
LANOLIN (UNII: 7EV65EAW6H)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73622-3035-2	0.9 g in 1 PACKET; Type 0: Not a Combination Product	05/30/2013	
2	NDC:73622-3035-3	14 g in 1 TUBE; Type 0: Not a Combination Product	05/30/2013	
3	NDC:73622-3035-4	28 g in 1 TUBE; Type 0: Not a Combination Product	05/30/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	05/30/2013	

Labeler - Curetech Skincare (677682180)

Establishment

Name	Address	ID/FEI	Business Operations
curetech skincare		677682180	manufacture(73622-3035)

Revised: 7/2020 Curetech Skincare