BACITRACIN- bacitracin ointment NuCare Pharmaceuticals,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.

Perrigo Bacitracin Drug Facts

Active ingredient (each gram contains)

Bacitracin 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

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- you need to use longer than 1 week
- condition persists or gets worse
- 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
- 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 20°-25°C (68°-77°F)

Inactive ingredients

light mineral oil, white petrolatum

Questions or comments?

1-800-719-9260

Principal Display Panel



BACITRACIN bacitracin ointment							
Product Information							
Product Type Route of Administration	HUMAN OTC DRUG	ltem Code (Source)	NDC:68071-2395(NDC:45802-060)				
Active Ingredient/Active Moiety							

	Ingredient Name	Basis of Strength	Strength			
ACITRACIN (UNII:	58H6RWO52I) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [USP'U] in 1 g			
nactive Ingre	dients					
		Strength				
IGHT MINERAL O						
ETROLATUM (UN						
Packaging						
# Item Code	Package Description	Marketing Start	Marketing End Date			
		Date	Date			
NDC:68071- 2395-5	1 in 1 CARTON	04/30/2021	Date			
NDC:68071- 2395-5			Date			
NDC:68071-	1 in 1 CARTON 14 g in 1 TUBE; Type 0: Not a Combination		Date			
1 NDC:68071- 2395-5	1 in 1 CARTON 14 g in 1 TUBE; Type 0: Not a Combination Product		Date			
1 NDC:68071- 2395-5	1 in 1 CARTON 14 g in 1 TUBE; Type 0: Not a Combination		Date			
NDC:68071- 2395-5	1 in 1 CARTON 14 g in 1 TUBE; Type 0: Not a Combination Product	04/30/2021	Date Marketing End Date			

Labeler - NuCare Pharmaceuticals,Inc. (010632300)

Establishment						
Name	Address	ID/FEI	Business Operations			
NuCare Pharmaceuticals, Inc.		010632300	relabel(68071-2395)			

Revised: 4/2021

NuCare Pharmaceuticals, Inc.