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.

Bacitracin Ointment

ACTIVE INGREDIENT

Bacitracin 500 units

PURPOSE

First aid antibiotic

USES

first aid to help prevent infection in minor cuts, scrapes and burns

WARNINGS

For external use only
Do not use
• if you are allergic to any of the ingredients
• in the eyes
• 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, or 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

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

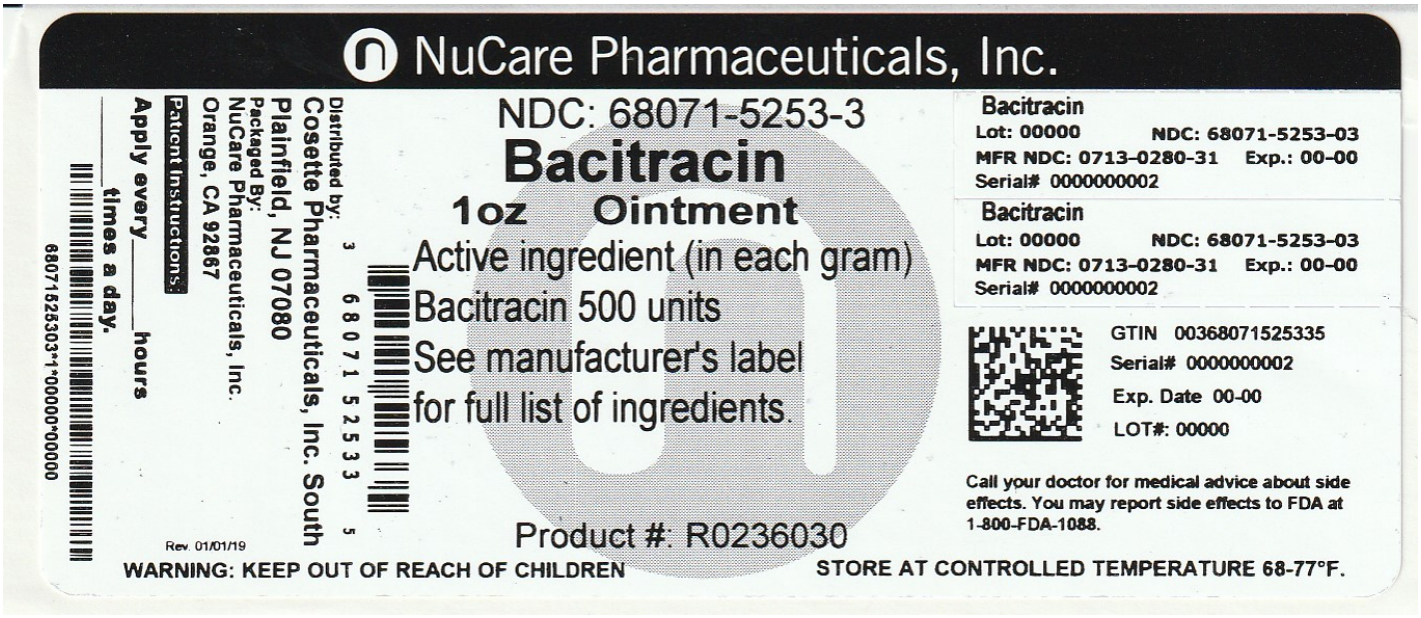
OTHER INFORMATION

store at room temperature

INACTIVE INGREDIENT

light mineral oil, white petrolatum

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



BACITRACIN

bacitracin ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68071-5253(NDC:0713-0280)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BACITRACIN (UNII: 58H6RWO52I) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [USP'U] in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-5253-3	28.4 g in 1 BOX; Type 0: Not a Combination Product	05/08/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	01/10/1995	

Labeler - NuCare Pharmaceuticals,Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals,Inc.		010632300	relabel(68071-5253)

Revised: 8/2023 NuCare Pharmaceuticals,Inc.