

**BACITRACIN- bacitracin ointment**  
**NuCare Pharmaceuticals, Inc.**

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**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**

**NuCare Pharmaceuticals, Inc.**

NDC: 68071-5253-3  
**Bacitracin**  
**1oz Ointment**  
 Active ingredient (in each gram)  
 Bacitracin 500 units  
 See manufacturer's label  
 for full list of ingredients.

Product #: R0236030

GTIN 00368071525335  
 Serial# 0000000002  
 Exp. Date 00-00  
 LOT#: 00000

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

WARNING: KEEP OUT OF REACH OF CHILDREN STORE AT CONTROLLED TEMPERATURE 68-77°F.

<b>BACITRACIN</b>				
bacitracin ointment				
<b>Product Information</b>				
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:68071-5253(NDC:0713-0280)	
<b>Route of Administration</b>	TOPICAL			
<b>Active Ingredient/Active Moiety</b>				
	Ingredient Name	Basis of Strength	Strength	
	BACITRACIN (UNII: 58H6RWO52I) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [USP'U] in 1 g	
<b>Inactive Ingredients</b>				
	Ingredient Name	Strength		
	LIGHT MINERAL OIL (UNII: N6K5787QVP)			
	PETROLATUM (UNII: 4T6H12BN9U)			
<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date

<b>1</b>	NDC:68071-5253-3	28.4 g in 1 BOX; Type 0: Not a Combination Product	05/08/2020	
<b>Marketing Information</b>				
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>		<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M014		01/10/1995	

**Labeler - NuCare Pharmaceuticals, Inc. (010632300)**

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

Revised: 7/2024

NuCare Pharmaceuticals, Inc.