**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:66267-988(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:66267-988-30 | 30 g in 1 TUBE; Type 0: Not a Combination Product | 07/18/2017 | |

| 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(66267-988) |

Revised: 1/2021 NuCare Pharmaceuticals,Inc.