

## **BACITRACIN - bacitracin ointment**

### **A-S Medication Solutions**

*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.*

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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

#### **HOW SUPPLIED**

Product: 50090-0846

NDC: 50090-0846-0 28.4 g in a TUBE

## INACTIVE INGREDIENT

light mineral oil, white petrolatum

## Bacitracin



## BACITRACIN

bacitracin ointment

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50090-0846(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:50090-0846-0	28.4 g in 1 TUBE; Type 0: Not a Combination Product	11/28/2014	

## Marketing Information

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

## Labeler - A-S Medication Solutions (830016429)

## Establishment

Name	Address	ID/FEI	Business Operations
A-S Medication Solutions		830016429	RELABEL(50090-0846)

Revised: 2/2021

A-S Medication Solutions