

## **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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### **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**

## Bacitracin



## BACITRACIN

bacitracin ointment

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:50090-0845(NDC:45802-060)
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>BACITRACIN</b> (UNII: 58H6RWO52I) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [USP'U] in 1 g

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>LIGHT MINERAL OIL</b> (UNII: N6K5787QVP)	
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:50090-0845-0	144 in 1 CARTON	11/28/2014	
1		.9 g in 1 PACKET; Type 0: Not a Combination Product		

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC monograph final	part333B	06/20/2011	

**Labeler** - A-S Medication Solutions (830016429)**Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
A-S Medication Solutions		830016429	RELABEL(50090-0845)

Revised: 2/2023

A-S Medication Solutions