

DRS. ANTIBIOTIC BACITRACIN- bacitracin ointment
OL PHARMA TECH, LLC

Active ingredient

Bacitracin zinc 500 units in one gram

Purpose

First aid antibiotic

Uses

First aid to help prevent infection in :

- minor cuts
- burns
- scrapes

Warnings

For external use only.

Do not use

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

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on deep or puncture wounds, animal bites, or serious burns

Stop use

Stop use and ask a doctor if

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- condition persists or gets worse
- a rash or other allergic reaction develops

Directions

Directions

- clean the affected area and dry thoroughly
- 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

Storage and Handling

- store at room temperature
- see carton or tube crimp for lot number and expiration date

Inactive ingredient

Petrolatum

keep out of reach of children

If swallowed get medical help or contact a poison control center right away

Questions

www.drspharmacyusa.com

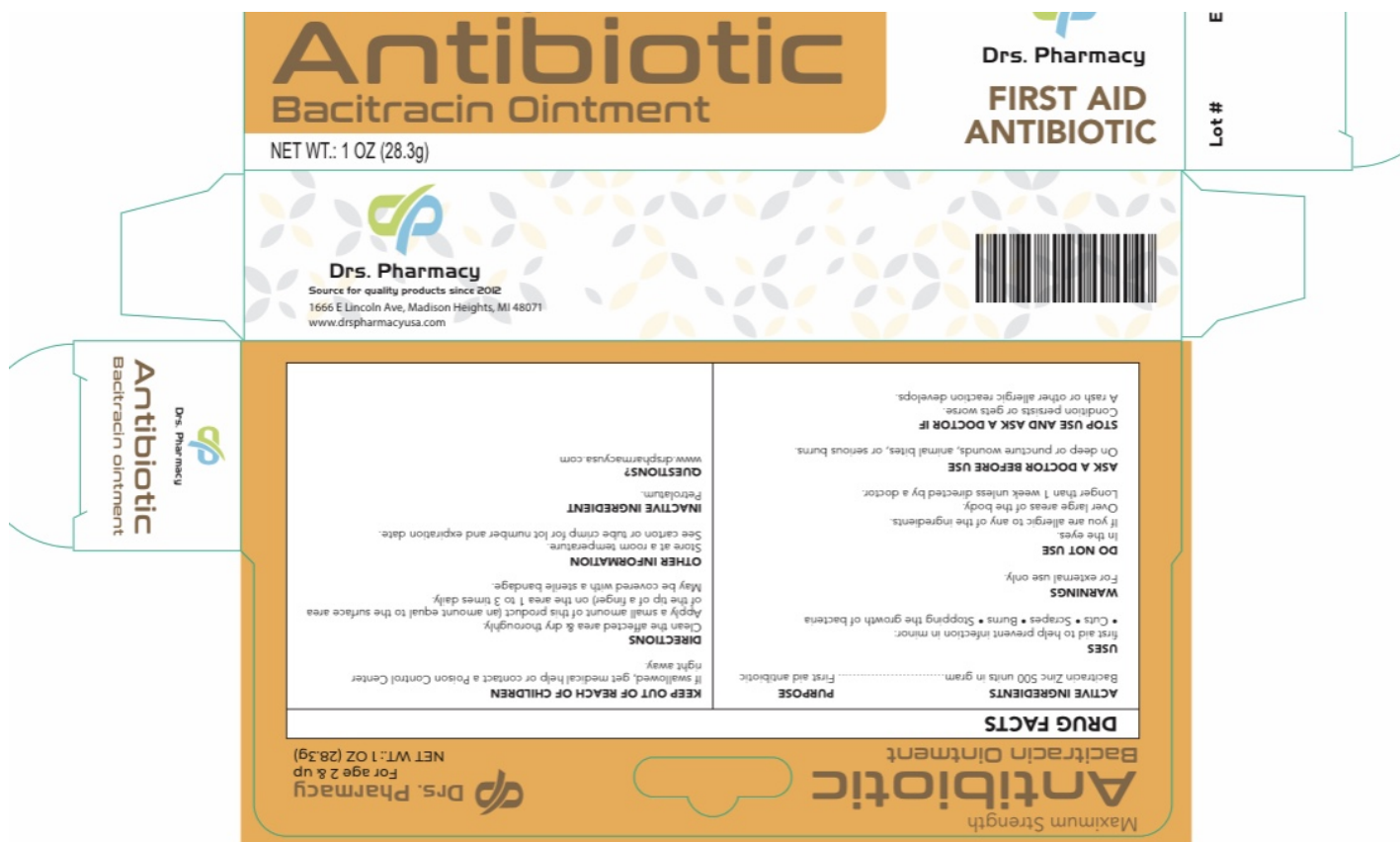


SHREE PACK
CONTAINERS PVT. LTD.

22.03.2021

Tube Ø22 x 134 length
143x32x26mm





Drs. Bacitracin Antibiotic ointment

DRS. ANTIBIOTIC BACITRACIN

bacitracin ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80489-006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [USP'U] in 1 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Product Characteristics

Color	white	Score	
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Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80489-006-01	1 in 1 CARTON	10/01/2021	
1		14 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:80489-006-02	1 in 1 CARTON	10/01/2021	
2		28.3 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	10/01/2021	

Labeler - OL PHARMA TECH, LLC (021170377)

Registrant - OL PHARMA TECH, LLC Drs PHARMACY (021170377)

Establishment

Name	Address	ID/FEI	Business Operations
OL PHARMA TECH, LLC Drs PHARMACY		021170377	manufacture(80489-006)

Revised: 1/2025

OL PHARMA TECH, LLC