DRS. PHARMACY TRIPLE ANTIBIOTIC- bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment ointment
OL PHARMA TECH, LLC

Active Ingredients

- Bacitracin Zinc 400 units
- Neomycin Sulfate 3.5mg
- Polymyxin B Sulfate 5,000 units

Purpose

- First aid antibiotic

Uses

First aid to help prevent infection in minor:

- cuts
- scrapes
- burns

Warnings

For external use only

Ask a doctor

Ask a doctor before use if you have

- serious burns
- deep or puncture wounds
- animal bites

Stop use and ask a doctor

Stop use and ask a doctor if

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

Do not use

- do not use in the eyes
- do not apply over large areas of the body
- if you are allergic to any of the ingredients
- longer than one week unless directed by your doctor

keep out of the 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 tot the surface area of the tip of a finer) on the area 1 to 3 times daily
- may be covered with a sterile bandage
- store at room temperature

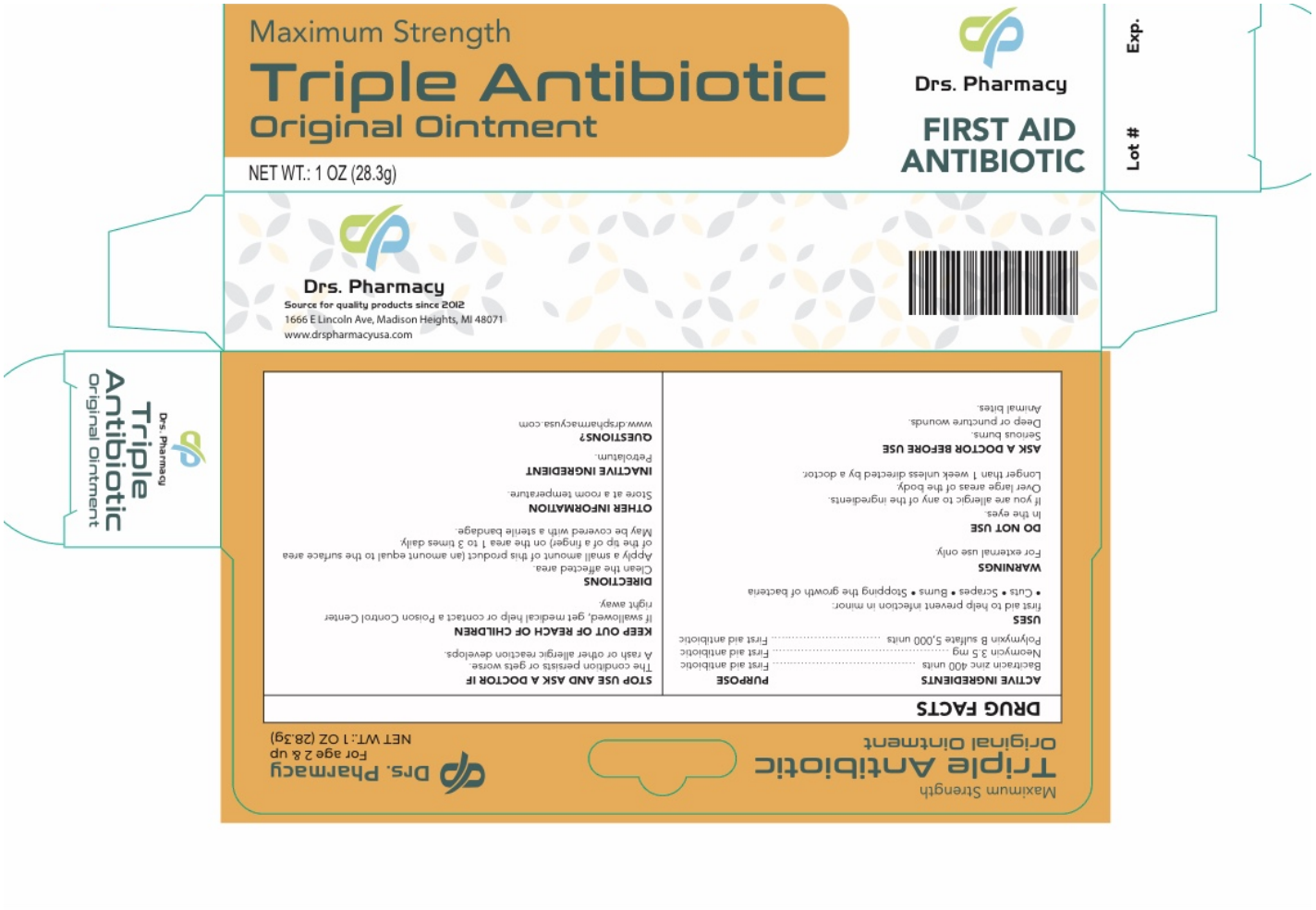
Inactive ingredient

Petrolatum

Questions

www.drspharmacyusa.com

Principal display panel



Maximum Strength

DRS. PHARMACY TRIPLE ANTIBIOTIC

bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment ointment

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80489-012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN ZINC	400 [USP'U] in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [USP'U] in 1 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Product Characteristics			
Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:80489-012-01	1 in 1 CARTON	02/16/2021	
1		14 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:80489-012-02	1 in 1 CARTON	02/16/2021	
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	02/16/2021	

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(80489-012)

Revised: 1/2025 OL PHARMA TECH, LLC