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

Active ingredients

Bacitracin zinc 500 units
Neomycin 3.5 mg
Polymyxin B sulfate 10,000 units

Pramoxine hydrochloride 10 mg

Uses

first aid to help prevent infection in minor
• cuts
• scrapes
• burns

Purpose

- first aid antibiotic
- External Analgesic

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

For external use only

Ask a doctor before use if you have

• deep or puncture wounds
• animal bites
• 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

Petrolatum

www.drspharmacyusa.com

DRS. PHARMACY TRIPLE ANTIBIOTIC AND BURN RELIEF

bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80489-919
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 1000 mg
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1000 mg

POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	10000 [USP'U] in 1000 mg
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1000 mg

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:80489-919-01	1 in 1 CARTON	09/05/2021	
1		14000 mg in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:80489-919-02	1 in 1 CARTON	09/05/2021	
2		28300 mg 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	09/05/2021	

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

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

Establishment

Name	Address	ID/FEI	Business Operations
OL PHARMA TECH, LLC Drs PHARMACY		021170377	manufacture(80489-919)