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

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.

Active ingredients

Bacitracin zinc 400 units
Neomycin 3.5 mg
Polymyxin B sulfate 5,000 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

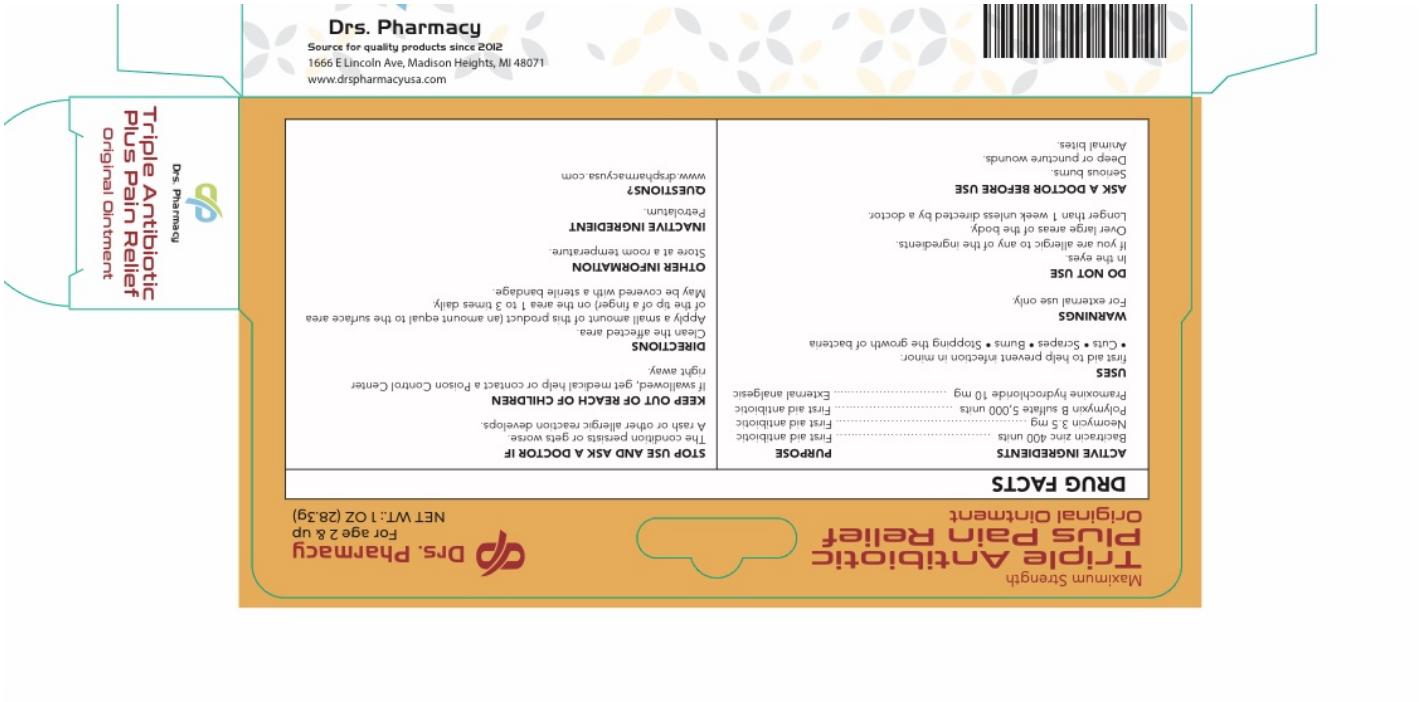
Inactive Ingredient

Petrolatum

www.drspharmacyusa.com

Tube Ø22 x 134 length

143x32x26mm



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Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80489-757
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [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	5000 [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:80489-757-01	1 in 1 CARTON	09/05/2021	
1		14000 mg in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:80489-757-02	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 final	part333B	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(80489-757)