OLP TRIPLE ANTIBIOTIC- bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment
Ohio Lab Pharma

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 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 before use if you have

- serious burns
- deep or puncture wounds
- animal bites

Stop use and ask a doctor if

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 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

OTHER INFORMATION

- store at room temperature

Inactive Ingredient

Petrolatum

QUESTIONS

www.ohiolabpharma.us

ORIGINAL OINTMENT

NET WEIGHT: 0.33OZ (9.4g)

NDC#70648-555-01

OLP TRIPLE ANTIBIOTIC

bacitracin zinc, polymyxin b sulfate, neomycin sulfate ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70648-555
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 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [USP'U] in 1 g

NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg 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:70648-555-01	1 in 1 CARTON	10/18/2017	
1		9.4 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 final	part333B	10/18/2017	

Labeler - Ohio Lab Pharma (080215854)

Establishment

Name	Address	ID/FEI	Business Operations
Ohio Lab Pharma		080215854	manufacture(70648-555)

Revised: 11/2018 Ohio Lab Pharma