

OLP TRIPLE ANTIBIOTIC PLUS- bacitracin zinc, neomycin sulfate, polymyxin b sulfate, lidocaine 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 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

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 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

Questions

www.ohiolabpharma.us



Net Weight 0.33 oz (9.4g)

NDC#70648-121-01

OLP TRIPLE ANTIBIOTIC PLUS

bacitracin zinc, neomycin sulfate, polymyxin b sulfate, lidocaine ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70648-121
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

NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [USP'U] in 1 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 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:70648-121-01	1 in 1 CARTON	12/06/2017	
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	12/06/2017	

Labeler - Ohio Lab Pharma (080215854)

Establishment

Name	Address	ID/FEI	Business Operations
Ohio Lab Pharma		080215854	manufacture(70648-121)

Revised: 11/2018

Ohio Lab Pharma