TRIPLE ANTIBIOTIC- bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment

Trifecta Pharmaceuticals Usa Llc

Triple Antibiotic Ointment

DRUG FACTS

Active ingredient (each gram contains)

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

Purpose

First aid antibiotic

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control

Center right away.

Uses first aid to help prevent infection in minor: ● cuts ● scrapes ● burns

Warnings For external use only.

Do not use ● in the eyes ● over large areas of the body

if you are allergic to any of the ingredients.

Ask a doctor before use if you have

◆ deep or puncture wounds◆ animal bites◆ serious burns.

Stop use and ask a doctor if ● condition persists or gets worse

- you need to use longer than 1 week
- a rash or other allergic reaction develops

Directions

- clean the affected area and dry thoroughly.
- 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

Inactive ingredient Light Mineral Oil, Petrolatum

Other information

- To open: unscrew cap, pull tab to remove foil seal
- store at 20° to 25°C (68° to 77°F)

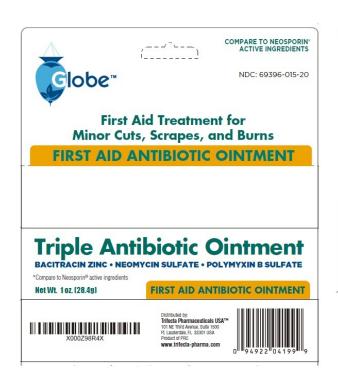
see carton or tube crimp for lot number and expiration date

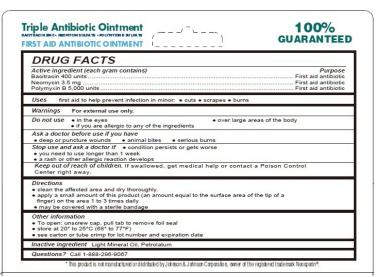
Questions? Call 1-888-296-9067

Distributed by:

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Packaging





TRIPLE ANTIBIOTIC

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Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69396-015
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RW052I)	BACITRACIN	400 [USP'U] in 1 g		
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:116QD7X297)	NEOMYCIN	3.5 mg in 1 g		
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII: J2VZ 07J96K)	POLYMYXIN B	5000 [USP'U] in 1 g		

Inactive Ingredients			
Ingredient Name	Strength		
LIGHT MINERAL OIL (UNII: N6K5787QVP)			
PETROLATUM (UNII: 4T6H12BN9U)			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:69396-015- 05	1 in 1 BOX	03/18/2016			
1		15 g in 1 TUBE; Type 0: Not a Combination Product				
2	NDC:69396-015- 20	1 in 1 BOX	03/18/2016			
2		28.4 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 Drug	M004	03/18/2016		

Labeler - Trifecta Pharmaceuticals Usa Llc (079424163)

Revised: 12/2024 Trifecta Pharmaceuticals Usa Llc