CVS TRIPLE ANTIBIOTIC- bacitracin, neomycin, polymyxin b ointment CVS 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.

Package Labeling

Drug Facts

Active Ingredients (in each gram)

Bacitracin 400 units

Neomycin 3.5 mg

Polymyxin B 5,000 units

Uses

first aid to help prevent infection in minor:

• cuts • scrapes • burns

Warnings

For external use only.

Do not use

- if you are allergic to any of the ingredients
- in the eyes
- over large areas of the body

Ask a doctor before use if you have

• deep or puncture wounds • animal bites • serious burns

Stop use and ask a doctor if

- you need to use longer than 1 week
- condition persists or gets worse
- rash or other allergic reaction develops side effects occur. You may report side effects to FDA at 1-800-FDA-1088.

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 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 Ingredients

white petrolatum

Purpose

First Aid Antibiotic

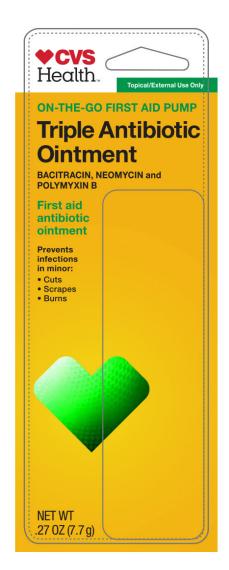
Other information

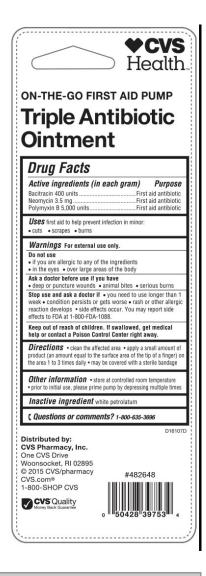
- •store at controlled room temperature
- prior to initial use, please prime pump by depressing multiple times

Questions or Comments?

1-800-635-3696







CVS TRIPLE ANTIBIOTIC

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Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69842-610

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BACITRACIN ZINC (UNII: 89 Y4M234ES) (BACITRACIN - UNII:58 H6 RWO 52I)	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:J2VZ07J96K)	POLYMYXIN B	5000 [USP'U] in 1 g	

Inactive Ingredients				
Ingredient Name	Strength			
PETROLATUM (UNII: 4T6H12BN9U)				

Packaging					
# Item Code	Package Description	Marketing Start Date	Marketing End Date		
1 NDC:69842-610-71	1 in 1 PACKAGE	11/0 1/20 0 8			
1	7.7 g in 1 BOTTLE; 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	11/01/2008		

Labeler - CVS Pharmacy (062312574)

Revised: 1/2017 CVS Pharmacy