

**ANTIBIOTIC APPLICATION- bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment**  
**CMC Group, Inc.**

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**Antibiotic Application**

**Drug Facts**

**Active ingredients (in each gram)**

Bacitracin zinc (bacitracin 400 units)

Neomycin sulfate (neomycin 3.5mg)

Polymyxin B sulfate (polymyxin B 5,000 units)

**Purpose**

First aid antibiotic

**Use**

- First aid to help prevent infection in minor cuts, scrapes, and 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 ingredient • 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 ingredients

Mineral oil, petrolatum, purified water

## Package Labeling:

**REFILL INSTRUCTIONS**

1. Remove old box from tabbed grid by slightly squeezing the front of box and pull straight out.
2. Insert the new product box into the grid with the removable panel facing the front, pushing it past the tabs, locking it into place.
3. Remove the upper half of the box front panel by hooking your finger into the hole, and pulling back. This will tear along the perforated edge and give access to the first aid product.

**Drug Facts**

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Bacitracin zinc (bacitracin 400 units)	First aid antibiotic
Neomycin sulfate (neomycin 3.5 mg)	
Polymyxin B sulfate (polymyxin B 5,000 units)	

**Use**  
 First aid to help prevent infection in minor cuts, scrapes, and burns.

**Warnings**  
 For external use only.  
 Do not use in the eyes or over large areas of the body.  
 If you are allergic to any of the ingredients or longer than 1 week unless directed by a doctor.  
 Ask a doctor before use if you have deep or puncture wounds, animal bites, or serious burns.  
 Stop use and ask a doctor if the condition persists or gets worse, a rash or other allergic reaction develops, or if you are pregnant, planning to get pregnant, or breastfeeding. 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.  
 Store at room temperature.

**Other information**  
 Inactive ingredients: Mineral oil, petrolatum, purified water.

ANSI-HSEA Z308.1-2015  
**DayMark**  
 Safety Systems  
 Antibiotic Application  
 Locion Antibiotica  
 10 packs IT113401

083491314405

## ANTIBIOTIC APPLICATION

bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:49687-0013
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BACITRACIN ZINC</b> (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
<b>NEOMYCIN SULFATE</b> (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g
<b>POLYMYXIN B SULFATE</b> (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ.07J96K)	POLYMYXIN B	5000 [iU] in 1 g

### Inactive Ingredients

Ingredient Name	Strength
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	
<b>WATER</b> (UNII: 059QF0KO0R)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49687-0013-0	10 in 1 KIT	08/06/2016	
1		0.9 g in 1 PACKAGE; Type 0: Not a Combination Product		

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	08/06/2016	

**Labeler** - CMC Group, Inc. (117201448)