

**TRIBIOTIC- bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment**  
**HART Health**

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**TRIBIOTIC**

Active Ingredients (in each gram)

Bacitracin Zinc 400 units

Neomycin Sulfate 5 mg (equivalent to 3.5mg neomycin base)

Polymyxin B Sulfate 5,000 units

Purpose:

Topical Antibiotic

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Uses: First aid to help prevent infection and temporarily relieve pain in

- minor cuts
- scrapes
- burns
- abrasions

Warnings: For external use only

Serious burns require immediate medical attention

Allergy alert: Do not use if you are allergic to any of the ingredients

do not use

- in or near the eyes
- over large portions of the body
- longer than 7 days unless directed by a doctor
- more than 1 to 3 times daily

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- condition worsens or lasts for more than 7 days
- a rash or other allergic reaction occurs

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

Directions:

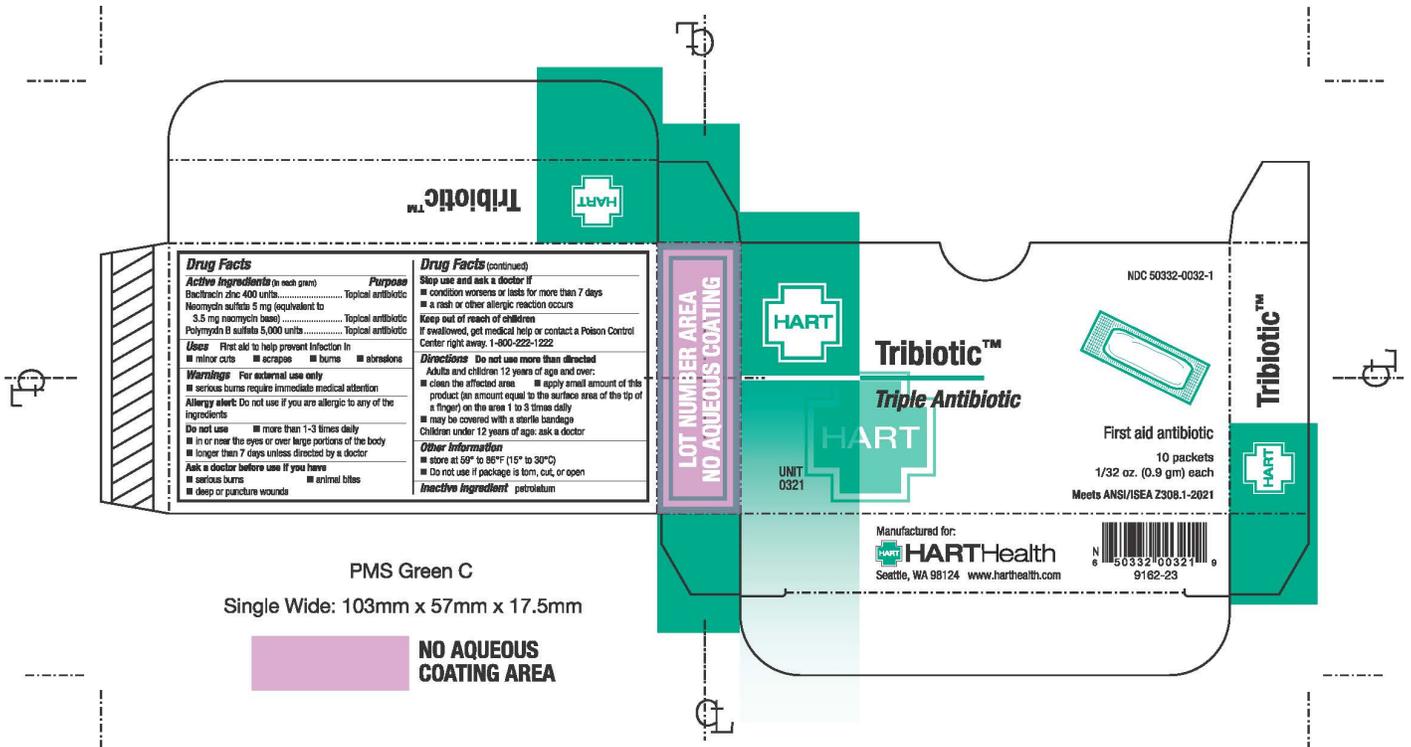
Do not use more than directed.

Adults and children 12 years of age and over:

- clean the affected area
- apply 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

Children under 12 years of age: ask a doctor

Petrolatum



## TRIBIOTIC

bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:50332-0032
<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:J2VZ07J96K)	POLYMYXIN B	5000 [iU] in 1 g

### Inactive Ingredients

Ingredient Name	Strength
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50332-0032-6	10 in 1 BOX, UNIT-DOSE	10/03/2005	
1		.5 g in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:50332-0032-3	25 in 1 BOX, UNIT-DOSE	10/03/2005	
2		.5 g in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:50332-0032-5	144 in 1 BOX, UNIT-DOSE	10/03/2005	
3		.5 g in 1 PACKET; Type 0: Not a Combination Product		
4	NDC:50332-0032-1	10 in 1 BOX, UNIT-DOSE	10/03/2005	
4		.9 g in 1 PACKET; Type 0: Not a Combination Product		
5	NDC:50332-0032-2	25 in 1 BOX, UNIT-DOSE	10/03/2005	
5		.9 g in 1 PACKET; Type 0: Not a Combination Product		
6	NDC:50332-0032-4	144 in 1 BOX, UNIT-DOSE	10/03/2005	
6		.9 g in 1 PACKET; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M	10/03/2005	

**Labeler** - HART Health (069560969)