

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

Target Corporation

Target Up &Up Triple Antibiotic Ointment

Drug Facts

Active Ingredients

Bacitracin 400

Neomycin 3.5 mg

Polymyxin B 5,000 units

Purpose

First aid antibiotic

First aid antibiotic

First aid antibiotic

Uses

First aid to help prevent infection in minor:

- cuts
- scrapes
- burns

Warnings

For external use only

Do not use if

- in or near the eyes
- on 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
- a rash or other allergic reaction develops

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

Directions

- clean the affected area
- apply a small amount (equal to surface area of tip of finger) on the area 1 to 3 times daily.
- may be covered with a sterile bandage

Other information

Store at a controlled room temperature 20°-25°C (68°-77°F)

Inactive ingredients

Petrolatum

Questions and Comments ? 1-800-910-6874

Principal Display Panel

Target Up&Up NDC 11673-892-56

Tripel Antibiotic Ointment

Polymyxin B Sulfate

Bacitracin Zinc

Neomycin Sulfate

First aid Antibiotic Ointment

NET WT 1oz(28g)

Total NET WT 2OZ (56g)



Original Strength Antibiotic Ointment

Bacitracin Zinc/Neomycin Sulfate/
Polymyxin B Sulfate

First aid antibiotic ointment

NET WT 1 OZ (28 g)

NDC 11673-892-56

Active ingredients (in each gram)

	<i>Purpose</i>
Bacitracin 400 units.....	First aid antibiotic
Neomycin 3.5 mg	First aid antibiotic
Polymyxin B 5,000 units	First aid antibiotic

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 or near 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 **Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center immediately. **Directions** • clean affected area • apply a small amount (equal to the surface area of tip of finger) on area 1 to 3 times daily • may be covered with a sterile bandage **Other information** • store at controlled room temperature 20°-25°C (68°-77°F).

Questions or comments? 1-800-910-6874

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Inactive ingredient
 White Petrolatum

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Compare to active ingredients in Neosporin®*

Original Strength Antibiotic Ointment

Bacitracin Zinc/
 Neomycin Sulfate/
 Polymyxin B Sulfate
 First aid antibiotic ointment

Original Strength Antibiotic Ointment

Distributed by Target Corporation
 Minneapolis, MN 55403
 Made in U.S.A. of U.S. and imported ingredients and components
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 Johnson & Johnson Consumer Products Company
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 owner of the registered trademark Neosporin®.

NDC 11673-892-56
 245 07 0076 R00 C-002262-01-035-0000

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TWO 1 OZ (28 g) TUBES,
 TOTAL NET WT 2 OZ (56 g)

LOT
 EXP

TRIPLE ANTIBIOTIC			
bacitracin zinc, neomycin sulfate, and polymyxin b sulfate ointment			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-892
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
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ.07J96K)	POLYMYXIN B	5000 [USP'U] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-892-56	2 in 1 CARTON	01/31/2024	
1		28 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	01/31/2024	

Labeler - Target Corporation (006961700)

Revised: 1/2024

Target Corporation