

TRIPLE ANTIBIOTIC- bacitracin zinc, neomycin sulfate , polymyxin b sulfate. ointment
Chain Drug Marketing Association

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.

Triple Antibiotic Ointment
First Aid Antibiotic

Drug Facts

Active ingredients (each gram contains)

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

Purpose

First Aid Antibiotic Ointment

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

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

Ask Doctor before use if you have

- Deep or puncture wounds
- Animal bites
- Serious burns

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

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.

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.

Inactive ingredient Mineral Oil, Petrolatum

Distributed by C.D.M.A., Inc. ©

43157 W 9 Mile Rd.

Novi, MI. 48375

www.qualitychoice.com

Questions: 800-935-2362

Other Information:

This product is not manufactured or distributed by

Johnson & Johnson Corporation, owner of the registered trademark Neosporin®

Packaging

OUTER BOX

NDC 63868-459-10



*Compare to the Active Ingredients in NEOSPORIN®

Triple Antibiotic Ointment

First Aid Antibiotic

Bacitracin Zinc | Neomycin Sulfate | Polymyxin B Sulfate

First Aid to Help Prevent Infection in:
Minor Cuts | Scrapes | Burns



Triple Antibiotic Ointment

First Aid Antibiotic

Bacitracin Zinc | Neomycin Sulfate | Polymyxin B Sulfate

1 oz (Net Wt 28.3 g)



Triple Antibiotic Ointment

* This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Neosporin®.



Distributed by C.D.M.A., Inc.®
43157 W 9 Mile Rd
Novi, MI 48375
www.qualitychoice.com
Questions: 800-935-2362



6 35515 99258 0

LOT:
EXP:

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 First aid antibiotic
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 Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.
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.
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.
Inactive ingredient Light Mineral Oil, Petrolatum

OUTER BOX

NDC 63868-459-05



Compare to the Active Ingredients in NEOSPORIN

Triple Antibiotic Ointment

First Aid Antibiotic

Bacitracin Zinc | Neomycin Sulfate | Polymyxin B Sulfate

First Aid to Help Prevent Infection in:
Minor Cuts | Scrapes | Burns



Triple Antibiotic Ointment

First Aid Antibiotic

Bacitracin Zinc | Neomycin Sulfate | Polymyxin B Sulfate

.5 oz (Net Wt. 14.2 g)



Triple Antibiotic Ointment

* This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Neosporin*.



Distributed by C.D.M.A., Inc.©
43157 W 9 Mile Rd
Novi, MI 48375
www.qualitychoice.com
Questions: 800-935-2362



LOT:
EXP:

DRUG FACTS (continued)	
Purpose	Active ingredient (each gram contains) Bacitracin Zinc (400 units)..... First aid antibiotic Neomycin Sulfate (3.5 mg)..... First aid antibiotic Polymyxin B Sulfate (5,000 units)..... First aid antibiotic
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.
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 or 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
Uses	First aid to help prevent infection in minor: • Cuts • Scrapes • Burns
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 cimp for lot number and expiration date.
Inactive Ingredients	Mineral oil, Petrolatum

TRIPLE ANTIBIOTIC

bacitracin zinc, neomycin sulfate , polymyxin b sulfate. ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-459
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
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	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
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-459-10	1 in 1 BOX	05/24/2019	
1		28.3 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:63868-459-05	1 in 1 BOX	03/26/2019	
2		14.2 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 final	part333B	03/26/2019	

Labeler - Chain Drug Marketing Association (011920774)**Registrant** - Trifecta Pharmaceuticals USA (079424163)