

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

Acme United Corporation

First Aid Only Triple Antibiotic Ointment

• ACTIVE INGREDIENT

Bacitracin zinc 400 units

Neomycin sulfate 5 mg (equivalent to 3.5 mg. of neomycin base)

Polymyxin - B sulfate 5000 units

Active ingredient (in each gram)	Purpose
Bacitracin Zinc 400 Units	First Aid Antibiotic
Neomycin Sulfate 5mg (Equivalent to 3.5 mg of Neomycin base)	First Aid Antibiotic
Polymyxin B Sulfate 5000 Units	First Aid Antibiotic

• USES

First aid to help prevent infection in minor:

- cuts
- scrapes
- burns

WARNINGS

For external use only

DO NOT USE

Do not use • in the eyes • over large areas of the body • if you are allergic to any of the ingredients

• longer than 1 week unless directed by a doctor • on deep lacerations or puncture wounds, animal bites, or serious burns

STOP USE AND ASK A DOCTOR IF

- condition persists or gets worse
- a rash or allergic reaction occurs
- condition persists for more than 7 days

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

DIRECTIONS

Adults and Children Clean the affected area. Apply a small amount of product (equal to the surface area of a fingertip) 1 to 3 times daily. May be covered with a sterile bandage.

OTHER INFORMATION

Store at 15° - 30°C (59° - 86°F)

INACTIVE INGREDIENT

White Petrolatum

Questions 1.800.835.2263

Carton Image

Drug Facts	
Active Ingredients	Purpose
Bacitracin zinc 400 units.....	First Aid antibiotic
Neomycin sulfate 5 mg (equivalent to 3.5 mg of neomycin).....	First Aid antibiotic
Polymyxin - B sulfate 5000 units.....	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 ■ longer than 1 week unless directed by a doctor ■ on deep lacerations or puncture wounds, animal bites, or serious burns	
Stop use and ask a doctor if ■ condition persists or gets worse ■ a rash or allergic reaction occurs ■ condition persists for more than 7 days	
Keep out the reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.	
Directions Adults & Children Clean the affected area. Apply a small amount of this product (equal to the surface area of a fingertip) 1 to 3 times daily. May be covered with sterile bandage.	
Other information Store at 15° - 30°C (59° - 86°F)	
Inactive ingredient white petrolatum	
Questions 1.800.835.2263	

12-001
ANTISEPTICS

Triple Antibiotic Ointment



FIRST AID ONLY.

Triple Antibiotic Ointment
12 Packets, 0.5g each

12-001
ANTISEPTICS

12-001
ANTISEPTICS

Triple Antibiotic Ointment

Meets ANSI/ISEA Z308.1-2015 Standard

Manufactured for:
Acme United Corporation
1 Waterview Dr., Shelton, CT 06484
www.FirstAidOnly.com
Made in India
© 2021 Acme United Corporation.
GUJ/DRUGS/G/28/1551



807709-revA

TRIPLE ANTIBIOTIC

bacitracin zinc, neomycin sulfate, and polymyxin b sulfate ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0924-5610(NDC:50382-023)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN SULFATE	5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ.07J96K)	POLYMYXIN B	5000 [iU] in 1 g

Inactive Ingredients

Ingredient Name	Strength

PETROLATUM (UNII: 4T6H12BN9U)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-5610-02	25 in 1 CARTON	08/05/2022	
1		0.5 g in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:0924-5610-00	0.5 g in 1 PACKET; Type 0: Not a Combination Product	08/05/2022	
3	NDC:0924-5610-01	12 in 1 CARTON	08/05/2022	
3		0.5 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	M004	08/04/2022	

Labeler - Acme United Corporation (001180207)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	repack(0924-5610) , relabel(0924-5610)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		080119599	repack(0924-5610) , relabel(0924-5610)

Revised: 11/2024

Acme United Corporation