TRIPLE ANTIBIOTIC- bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment Acme United Corporation

First Aid Only Triple Antibiotic Ointment Drug Facts

Active Ingredient (in each gram)

Bacitracin Zinc (400 units Bacitracin)

Neomycin Sulfate (3.5 mg Neomycin)

Polymyxin B Sulfate (Polymyxin B 5000 units)

Purpose

First Aid Antibiotic

First Aid Antibiotic

First Aid Antibiotic

Uses

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
- on puncture wounds, animal bites, or serious burns
- longer than 1 week unless directed by a doctor
- if you are allergic to any of the ingredients

Stop use and ask a doctor if:

- a rash or allergic reaction develops
- condition worsens or persists

Keep out the reach of children.

Directions

- Clean the affected area.
- Apply a small amount of product 1 to 3 times daily

• may be covered with a sterile bandage

Inactive Ingredient

White Petrolatum

Questions

1-800-835-2263

Box Label





Manufactured for:
Acme United Corporation
1 Waterview Dr, Shelton, CT 06484
www.FirstAidOnly.com
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213460-001-revA

TRIPLE ANTIBIOTIC

bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0924-5604(NDC:61010-5603)	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RW052I)	BACITRACIN	400 [iU] in 1 g	
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:116QD7X297)	NEOMYCIN	3.5 mg in 1 g	
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII: J2VZ 07J96K)	POLYMYXIN B	5000 [iU] in 1 g	

Inactive Ingredients			
Strength			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-5604- 00	0.5 g in 1 POUCH; Type 0: Not a Combination Product	04/24/2023	
2	NDC:0924-5604- 01	6 in 1 BAG	04/24/2023	
2		0.5 g in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:0924-5604- 03	12 in 1 BOX	04/24/2023	
3		0.5 g in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:0924-5604- 02	10 in 1 BOX	04/24/2023	
4		0.5 g in 1 PACKET; Type 0: Not a Combination Product		
5	NDC:0924-5604- 04	20 in 1 BOX	04/24/2023	
5		0.5 g in 1 PACKET; Type 0: Not a Combination Product		
6	NDC:0924-5604- 05	25 in 1 BOX	04/24/2023	
6		0.5 g in 1 PACKET; Type 0: Not a Combination Product		
7	NDC:0924-5604- 06	60 in 1 BOX	04/24/2023	
7		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	04/24/2023	

Labeler - Acme United Corporation (001180207)

Establishment			
Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	relabel(0924-5604), repack(0924-5604)

Establishment			
Name	Address	ID/FEI	Business Operations
Acme United Corporation		080119599	relabel(0924-5604), repack(0924-5604)

Revised: 11/2024 Acme United Corporation