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

First Aid Only Triple Antibiotic Ointment

Drug Facts

Active Ingredients (in each gram)

Bacitracin zinc 400 units

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

Polymyxin B sulfate 5000 units

Purpose

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
- the condition persists for more than 7 days

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center immediately.

Directions

- clean the affected area
- apply small amount 1 to 3 times daily

- may be covered with a sterile bandage

Other Information

Store at room temperature

Inactive Ingredient

white petrolatum

Questions

1.800.835.2263

Box Label

FIRST AID ONLY TRIPLE ANTIBIOTIC

bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0924-5618(NDC:61040-226)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 U in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 U in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
WHITE PETROLATUM (UNII: B6E5W8RQJ4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-5618-04	60 in 1 BOX	02/24/2023	
1		0.5 g in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:0924-5618-00	0.5 g in 1 POUCH; Type 0: Not a Combination Product	02/24/2023	
3	NDC:0924-5618-02	12 in 1 BOX	06/12/2023	
3		0.5 g in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:0924-5618-03	25 in 1 BOX	06/12/2023	
4		0.5 g in 1 POUCH; 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	02/24/2023	

Labeler - Acme United Corporation (001180207)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		080119599	relabel(0924-5618) , repack(0924-5618)

Revised: 11/2024 Acme United Corporation