TRIPLE ANTIBIOTIC- bacitracin zinc, neomycin sulfate, polymyxin b sulfate, polymyxin b cream
J&A Digital Inc.

Triple Antibiotic

Drug Facts

***Active ingredients*(in each gram)**

Bacitracin zinc (bacitracin 400 units)

Neomycin sulfate (neomycin 3.5 mg)

Polymyxin B sulfate

(polymyxin B 5,000 units)

Purpose

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 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

Ask a doctor

before use if you have

- deep or puntucre wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- the condition persists or gets worse
- 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

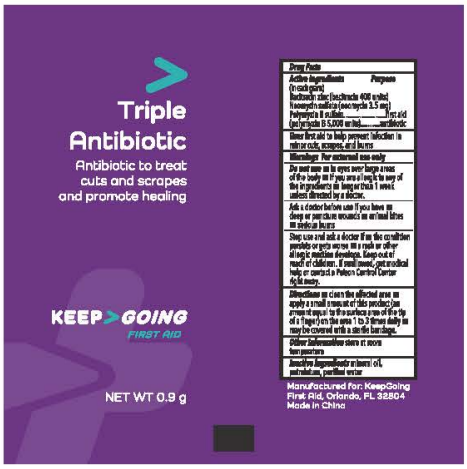
- clean the affected area
- apply a small amount of this product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily
- may be covered with a sterile bandage.

Other Information

store at room temperature

Inactive ingredients

Mineral oil, petrolatum, purified water



TRIPLE ANTIBIOTIC

bacitracin zinc, neomycin sulfate, polymyxin b sulfate, polymyxin b cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82942-1002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [CFU] 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 [CFU] in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:82942-1002-1	0.9 g in 1 PACKET; Type 0: Not a Combination Product	10/01/2022	

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
OTC Monograph Drug	M004	10/01/2022	

Labeler - J&A Digital Inc. (040268672)

Registrant - J&A Digital Inc. (040268672)

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