TRIPLE ANTIBIOTIC- bacitracin zinc, neomycin sulfate, polymyxin b sulfate 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.

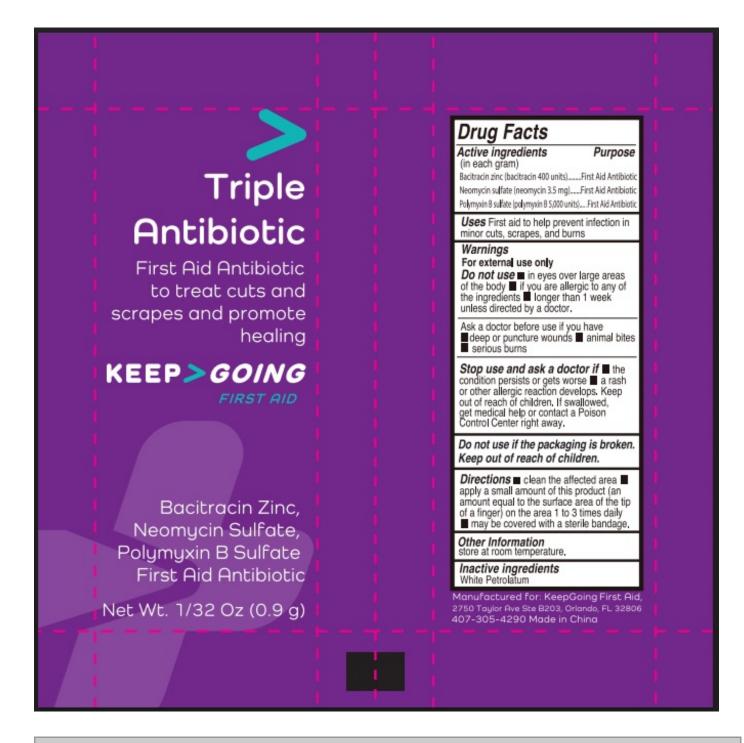
Other Information

store at room temperature

Inactive ingredients

White petrolatum

Product label



TRIPLE ANTIBIOTIC

bacitracin zinc, neomycin sulfate, polymyxin b sulfate cream

BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RW052I)

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82942-1004
Route of Administration	TOPICAL		
Active Ingredient/Active	Moiety		
Ingre	edient Name	Basis (Strength

BACITRACIN

400 [CFU] 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 [CFU] in 1 g

Inactive Ingredients		
Ingredient Name	Strength	
WHITE PETROLATUM (UNII: B6E5W8RQI4)		

ı	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:82942- 1004-1	0.9 g in 1 PACKET; Type 0: Not a Combination Product	07/31/2025		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M004	07/31/2025		

Labeler - J&A Digital Inc. (040268672)

Registrant - J&A Digital Inc. (040268672)

Revised: 8/2025 J&A Digital Inc.