TRIPLE ANTIBIOTIC- triple antibiotic ointment
Front Line Safety

FL-2009

Active Ingredient (in each gram)

Bacitracin Zinc 400 units

Purpose

First Aid Antibiotic

Active Ingredient (in each gram)

Neomycin Sulfate 5 mg (equivalent to 3.5 mg Neomycin)

Purpose

First Aid Antibiotic

Active Ingredient (in each gram)

Polymyxin B Sulfate 5000 units

Purpose

First Aid Antibiotic

Use(s)

First aid to help prevent infection in minor cuts, scrapes, and burns

Warnings

For External Use Only

Do not use

- In the eyes or apply 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 puncture wounds, animal bites, or serious burns

Stop use and ask a doctor if

• The condition perists 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 (1-800-222-1222) 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 between 15º-30ºC (59º-86ºF)

• Tamper evident. Do not use if packet is torn, cut, or opened.

Inactive Ingredient

White Petrolatum

Questions?

1-888-900-2920 Monday - Friday 8AM-4PM PST.

Label

FL-2009

TRIPLE ANTIBIOTIC

triple antibiotic ointment

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:58228-6209
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	5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [iU] in 1 g

Inactive Ingredients

Ingredient Name	Strength
WHITE PETROLATUM (UNII: B6E5W8RQJ4)	

Product Characteristics

Color		**Score**	
Shape	FREEFORM	**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58228-6209-1	1800 in 1 CASE	01/26/2024	
1		25 in 1 BOX		
1		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	01/26/2024	

Labeler - Front Line Safety (061263699)

Revised: 12/2025 Front Line Safety