TRIPLE ANTIBIOTIC- polymyxin b sulfate, bacitracin zinc and neomycin sulfate ointment
A-S Medication Solutions

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

ACTIVE INGREDIENT (in each gram)

Bacitracin zinc 400 units
Neomycin 3.5 mg
Polymyxin B sulfate 5,000 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
• if you are allergic to any of the ingredients
• in the eyes
• over large areas of the body
• longer than 1 week unless directed by a doctor

Ask a doctor before use if you have
• deep or puncture 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 INGREDIENT

light mineral oil, white petrolatum

HOW SUPPLIED

Product: 50090-0234

NDC: 50090-0234-0 28.4 g in a TUBE

TRIPLE ANTIBIOTIC OINTMENT

TRIPLE ANTIBIOTIC

polymyxin b sulfate, bacitracin zinc and neomycin sulfate ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50090-0234(NDC:0713-0268)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [USP'U] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [USP'U] in 1 g

Inactive Ingredients

Ingredient Name	Strength
Light Mineral Oil (UNII: N6K5787QVP)	
Petrolatum (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50090-0234-0	28.4 g in 1 TUBE; Type 0: Not a Combination Product	11/28/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	01/11/1995	

Labeler - A-S Medication Solutions (830016429)

Establishment

Name	Address	ID/FEI	Business Operations
A-S Medication Solutions		830016429	RELABEL(50090-0234)

Revised: 3/2023 A-S Medication Solutions