

SUNMARK TRIPLE ANTIBIOTIC PLUS PAIN RELIEF- polymyxin b sulfate, bacitracin zinc, neomycin sulfate, and pramoxine hydrochloride ointment
A-S Medication Solutions

SUNMARK TRIPLE ANTIBIOTIC PLUS PAIN RELIEF

Drug Facts

<i>Active ingredients (in each gram)</i>	<i>Purposes</i>
Bacitracin 500 units	First aid antibiotic
Neomycin 3.5 mg	First aid antibiotic
Polymyxin B 10,000 units	First aid antibiotic
Pramoxine HCl 10 mg	Topical pain reliever

Uses

first aid to help prevent infection and for temporary relief of pain or discomfort in minor:

- cuts
- scrapes
- burns

Warnings

For external use only

Do not use if you are allergic to any of the ingredients

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

When using this product

- do not use in the eyes
- do not apply over large areas of the body

Stop use and ask a doctor if

- you need to use for more than 1 week
- condition persists or gets worse
- symptoms persist for more than 1 week or clear up and occur again within a few days
- 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

- unscrew cap and pull tab to remove foil seal
- adults and children 2 years of age and older:
 - 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
- children under 2 years of age: ask a doctor

Other information

- store at controlled room temperature 15°-30°C (59°-86°F)
- see carton or tube crimp for lot number and expiration date

Inactive ingredient

white petrolatum

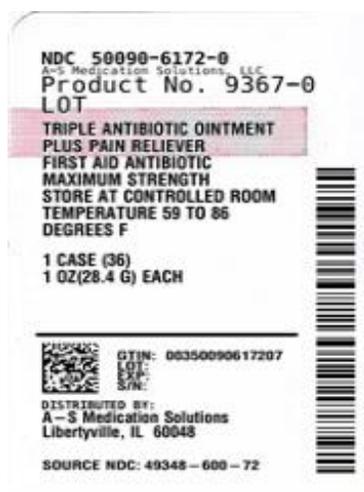
Distributed by McKesson
One Post Street
San Francisco, CA 94104

HOW SUPPLIED

Product: 50090-6172

NDC: 50090-6172-0 28.4 g in a TUBE / 36 in a CASE

polymyxin B sulfate, bacitracin zinc, neomycin sulfate, and pramoxine hydrochloride



SUNMARK TRIPLE ANTIBIOTIC PLUS PAIN RELIEF

polymyxin b sulfate, bacitracin zinc, neomycin sulfate, and pramoxine hydrochloride ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50090-6172(NDC:49348-600)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [iU] 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:J2VZ.07J96K)	POLYMYXIN B	10000 [iU] in 1 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50090-6172-0	36 in 1 CASE	10/14/2022	08/31/2027
1		28.4 g in 1 TUBE; 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	03/31/2012	08/31/2027

Labeler - A-S Medication Solutions (830016429)

Establishment

Name	Address	ID/FEI	Business Operations
A-S Medication Solutions		830016429	RELABEL(50090-6172)