

SUNMARK TRIPLE ANTIBIOTIC PLUS PAIN RELIEF- polymyxin b sulfate, bacitracin zinc, neomycin sulfate, and pramoxine hydrochloride ointment
Strategic Sourcing Services LLC

sunmark™
triple antibiotic plus pain relief

Drug Facts

Active ingredients (in each gram)	Purposes
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

PRINCIPAL DISPLAY PANEL - 28.4 g Tube Carton

sunmark™

triple antibiotic ointment

plus pain reliever

**polymyxin B sulfate • bacitracin zinc
neomycin sulfate • pramoxine hydrochloride**

First Aid Antibiotic

MAXIMUM STRENGTH

NET WT 1 OZ (28.4 g)

sunmark™

COMPARE TO NEOSPORIN® PLUS ACTIVE INGREDIENTS*

NDC 49348-600-72

Helps prevent infection in minor cuts, scrapes & burns
Maximum strength pain relief

MAXIMUM STRENGTH

LPK-4655-0
1003-0
M105

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First Aid Antibiotic



This product is not manufactured or distributed by Pfizer, Inc. or its affiliates, owner of the registered trademark. M5KES5SON Impression Number
Another Quality Product Distributed by McKesson One Post Street San Francisco, CA 94104
Please visit us at www.sunmark.com
Made in Canada

Drug Facts

Active ingredients (in each gram)

Bacitracin 500 units
Neomycin 35 mg
Polymyxin B 10,000 units
Pramoxine HCl 10 mg

First aid antibiotic
First aid antibiotic
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

Directions

• unseal 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

plus pain reliever
polymyxin B sulfate • bacitracin zinc
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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	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:49348-600-72	1 in 1 CARTON	02/13/2013	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 - Strategic Sourcing Services LLC (116956644)**Establishment**

Name	Address	ID/FEI	Business Operations
Sun Pharma Canada Inc.		243339023	manufacture(49348-600)

