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

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.

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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

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Drug Facts

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Bacitracin 500 units	First aid antibiotic
Neomycin 35 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

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Inactive ingredient white petrolatum

plus pain reliever
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*This product is not manufactured or distributed by Pfizer, Inc. or its affiliates, owner of the registered trademark Neosporin Plus.

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Empowering Healthcare

Another Quality Product Distributed by McKesson
One Post Street
San Francisco, CA 94104
Money Back Guarantee
Please visit us at
www.sunmarkbrand.com

Made in Canada.

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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:J2VZ07J96K)	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	
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 final	part333B	03/31/2012	

Labeler - Strategic Sourcing Services LLC (116956644)

Registrant - Taro Pharmaceuticals U.S.A., Inc. (145186370)

Establishment

Name	Address	ID/FEI	Business Operations
Taro Pharmaceuticals Inc.		206263295	MANUFACTURE(49348-600)