SUNMARK TRIPLE ANTIBIOTIC- polymyxin b sulfate, bacitracin zinc, and neomycin sulfate ointment
McKesson

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

Drug Facts

Active ingredients (in each gram)	*Purposes*
Bacitracin 400 units	First aid antibiotic
Neomycin 3.5 mg	First aid antibiotic
Polymyxin B 5,000 units	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

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™

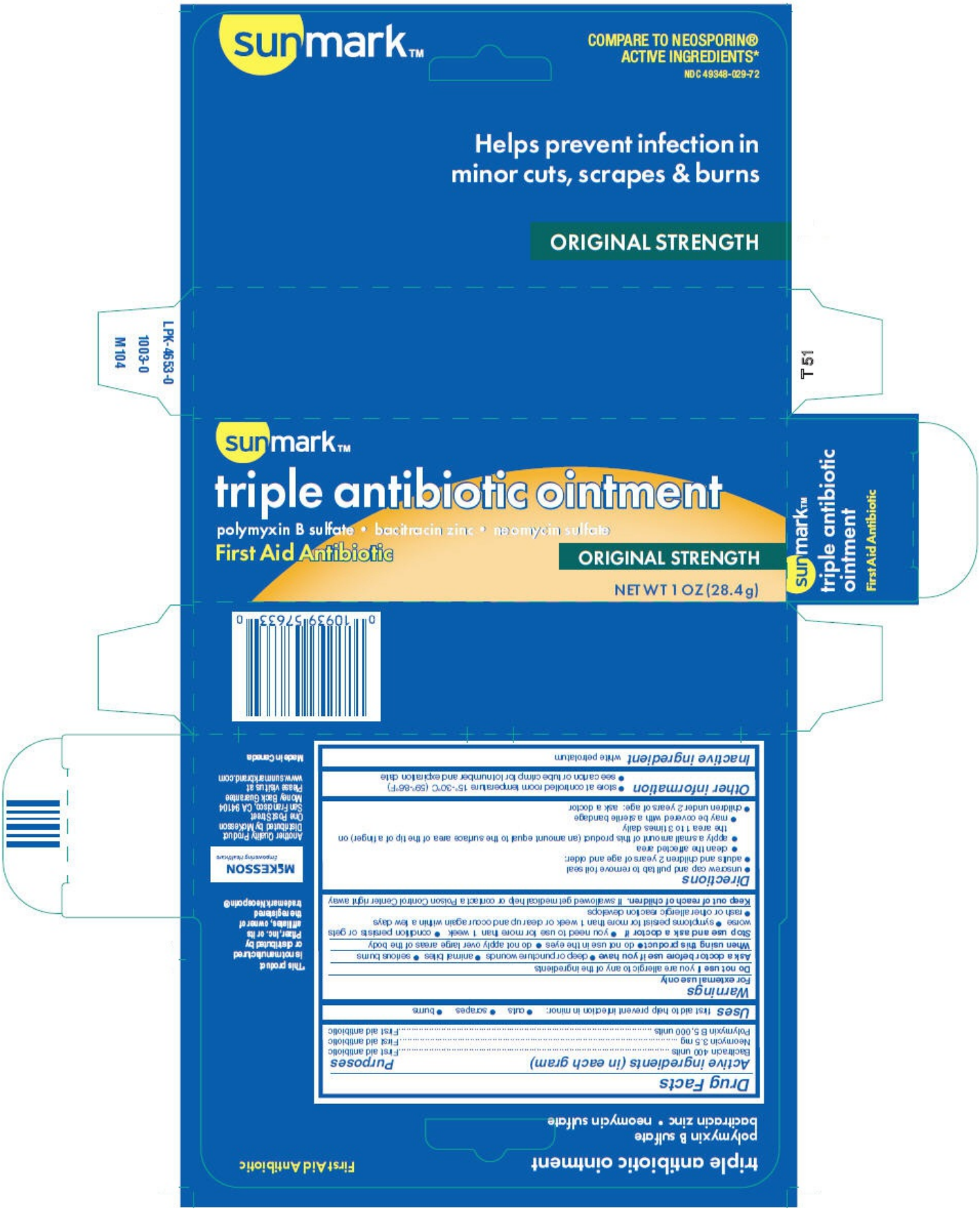
triple antibiotic ointment

polymyxin B sulfate • bacitracin zinc • neomycin sulfate

First Aid Antibiotic

ORIGINAL STRENGTH

NET WT 1 OZ (28.4 g)



SUNMARK TRIPLE ANTIBIOTIC

polymyxin b sulfate, bacitracin zinc, and neomycin sulfate ointment

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49348-029
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	3.5 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
petrolatum (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49348-029-72	1 in 1 CARTON		
1		28.4 g in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	12/14/1995	

Labeler - McKesson (177667227)

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

Establishment

Name	Address	ID/FEI	Business Operations
Taro Pharmaceuticals Inc.		206263295	MANUFACTURE(49348-029)

Revised: 4/2013 McKesson