TRIPLE ANTIBIOTIC PLUS PAIN RELIEF- bacitracin zinc, neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride ointment
Sun Pharmaceutical Industries, Inc.

Triple Antibiotic Plus Pain Relief

Active ingredients (each gram contains)	***Purpose***
Bacitracin zinc 500 units	First aid antibiotic
Neomycin sulfate 3.5 mg	First aid antibiotic
Polymyxin B sulfate 10,000 units	First aid antibiotic
Pramoxine HCl 10 mg	External analgesic

Uses

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

- cuts
- scrapes
- burns

Warnings

For external use only.

Do not use

- in the eyes
- over large areas of the body
- 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.

Stop use and ask a doctor if

- you need to use longer 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
- 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

Adults and children 2 years of age and older:

- clean the affected area and dry thoroughly
- 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

- To open: unscrew cap, pull tab to remove foil seal
- store at 20° to 25°C (68° to 77°F)
- see carton or tube crimp for lot number and expiration date

Inactive ingredient

white petrolatum

Questions?

Call **1-866-923-4914**

PRINCIPAL DISPLAY PANEL - 28.4 g Tube Label

First Aid Antibiotic
Pain Relieving Ointment

Maximum Strength
Triple Antibiotic
Ointment + Pain
Relief

Bacitracin Zinc • Neomycin Sulfate
Polymyxin B Sulfate • Pramoxine Hydrochloride

NET WT 1 oz (28.4 g)

TRIPLE ANTIBIOTIC PLUS PAIN RELIEF

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

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51672-2027
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN ZINC	500 [USP'U] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN SULFATE	3.5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	10000 [USP'U] in 1 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE -	PRAMOXINE	10 mg in 1 g

UNII:068X84E056)	HYDROCHLORIDE	10 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WHITE PETROLATUM (UNII: B6E5W8RQJ4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51672-2027-1	1 in 1 CARTON	03/31/2012	
1		14.2 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:51672-2027-2	1 in 1 CARTON	03/31/2012	
2		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	

Labeler - Sun Pharmaceutical Industries, Inc. (146974886)

Establishment

Name	Address	ID/FEI	Business Operations
Sun Pharma Canada Inc.		243339023	manufacture(51672-2027)

Revised: 7/2025 Sun Pharmaceutical Industries, Inc.