

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

**Sun Pharmaceutical Industries, Inc.**

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**Triple Antibiotic**

**Drug Facts**

<b>Active ingredients (in each gram)</b>	<b>Purpose</b>
Bacitracin zinc USP 400 units	First aid antibiotic
Neomycin sulfate USP 3.5 mg	First aid antibiotic
Polymyxin B sulfate USP 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
- in the eyes
- over large areas of the body

**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
- 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**

- 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

**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 ingredients**

cocoa butter, cottonseed oil, olive oil, sodium pyruvate, vitamin E, white petrolatum

**Questions?**

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

Distributed by:

**Taro Pharmaceuticals U.S.A., Inc.**

Hawthorne, NY 10532

**PRINCIPAL DISPLAY PANEL - 28.4 g Tube Carton****Original Strength****Triple Antibiotic  
Ointment**

*Bacitracin Zinc USP • Neomycin Sulfate USP • Polymyxin B Sulfate USP*

**First Aid Antibiotic Ointment**

**NET WT 1 oz (28.4 g)**

NDC 51672-2120-2

Compare to the active ingredients in Neosporin®\*

# Triple Antibiotic Ointment



Bacitracin Zinc USP • Neomycin Sulfate USP • Polymyxin B Sulfate USP

First Aid Antibiotic Ointment



Helps Prevent Infection in Minor Cuts, Scrapes, and Burns.

LPK-7939-0  
0915-0  
M603

# Triple Antibiotic Ointment

Original Strength

Bacitracin Zinc USP • Neomycin Sulfate USP • Polymyxin B Sulfate USP

First Aid Antibiotic Ointment

NET WT 1 oz (28.4 g)

Triple Antibiotic Ointment  
Bacitracin Zinc USP • Neomycin Sulfate USP  
Polymyxin B Sulfate USP  
First Aid Antibiotic Ointment

T51  
B503  
ENG19.13



Distributed by:  
Taro Pharmaceuticals U.S.A., Inc.  
Hawthorne, NY 10532

TARO is a registered trademark of Taro Pharmaceuticals U.S.A., Inc.  
Made in Canada



NO VARNISH/ NO AQ  
NO COPY/ NO COLOR  
THIS FLAP FOR LOT #  
AND EXP DATE PRINT

\*This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Neosporin®

<p><b>Drug Facts</b> (continued)</p> <p><b>Keep out of reach of children.</b> If swallowed, get medical help or contact a Poison Control Center right away.</p> <p><b>Directions</b></p> <ul style="list-style-type: none"> <li>• apply a small amount of the product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily</li> <li>• may be covered with a sterile bandage</li> </ul> <p><b>Other Information</b></p> <ul style="list-style-type: none"> <li>• To open: unscrew cap, pull tab to remove bil seal</li> <li>• store at 20° to 25° C (68° to 77° F)</li> <li>• see carton or tube crimp for lot number and expiration date</li> </ul> <p><b>Inactive Ingredients</b></p> <p>cocoa butter, colored oil, olive oil, sodium pyruvate, vitamin E, white petrolatum</p> <p><b>Questions?</b> Call 1-866-923-4914</p>	<p><b>Drug Facts</b></p> <p><b>Active ingredients (in each gram) Purpose</b></p> <p>Bacitracin zinc USP 400 units.....First aid antibiotic</p> <p>Neomycin sulfate USP 3.5 mg.....First aid antibiotic</p> <p>Polymyxin B sulfate USP 5,000 units.....First aid antibiotic</p> <p><b>Uses</b> First aid to help prevent infection in minor:</p> <ul style="list-style-type: none"> <li>• cuts • scrapes • burns</li> </ul> <p><b>Warnings</b></p> <p>For external use only.</p> <p><b>Do not use</b> • If you are allergic to any of the ingredients</p> <ul style="list-style-type: none"> <li>• in the eyes</li> <li>• over large areas of the body</li> </ul> <p><b>Ask a doctor before use if you have</b></p> <ul style="list-style-type: none"> <li>• deep or puncture wounds</li> <li>• animal bites</li> <li>• serious burns</li> </ul> <p><b>Stop use and ask a doctor if</b></p> <ul style="list-style-type: none"> <li>• you need to use longer than 1 week</li> <li>• condition persists or gets worse</li> <li>• rash or other allergic reaction develops</li> </ul>
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Triple Antibiotic Ointment  
Bacitracin Zinc USP • Neomycin Sulfate USP  
Polymyxin B Sulfate USP  
Original Strength  
First Aid Antibiotic Ointment

## TRIPLE ANTIBIOTIC

bacitracin zinc, neomycin sulfate, and polymyxin b sulfate ointment

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:51672-2120
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>BACITRACIN ZINC</b> (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [iU] in 1 g
<b>NEOMYCIN SULFATE</b> (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g
<b>POLYMYXIN B SULFATE</b> (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ.07J96K)	POLYMYXIN B	5000 [iU] in 1 g

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>COCOA BUTTER</b> (UNII: 512OYT1CRR)	
<b>COTTONSEED OIL</b> (UNII: H3E878020N)	
<b>OLIVE OIL</b> (UNII: 6UYK2W1W1E)	
<b>SODIUM PYRUVATE</b> (UNII: POD38AIF08)	
<b>.ALPHA.-TOCOPHEROL</b> (UNII: H4N855PNZ1)	
<b>PETROLATUM</b> (UNII: 4T6H12BN9U)	

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:51672-2120-1	1 in 1 CARTON	12/15/2015	
1		14.2 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:51672-2120-2	1 in 1 CARTON	12/15/2015	
2		28.4 g in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:51672-2120-3	1 in 1 CARTON	12/15/2015	
3		60 g in 1 TUBE; Type 0: Not a Combination Product		
4	NDC:51672-2120-6	6 in 1 CARTON	09/20/2021	
4		0.9 g in 1 APPLICATOR; Type 0: Not a Combination Product		
5	NDC:51672-2120-8	15 in 1 CARTON	09/20/2021	
5		0.9 g in 1 APPLICATOR; Type 0: Not a Combination Product		
6	NDC:51672-2120-9	144 in 1 CARTON	09/20/2021	
6		0.9 g in 1 APPLICATOR; 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	12/15/2015	

**Labeler** - Sun Pharmaceutical Industries, Inc. (146974886)

## Establishment

Name	Address	ID/FEI	Business Operations
Sun Pharma Canada Inc.		243339023	manufacture(51672-2120)

Revised: 7/2025

Sun Pharmaceutical Industries, Inc.