TRIPLE ANTIBIOTIC PLUS PAIN RELIEF MAXIMUM STRENGTH- bacitracin zinc, neomycin sulfate, polymyxin b sulfate and pramoxine hydrochloride ointment Chain Drug Consortium, LLC

Triple Antibiotic Plus Pain Relief Maximum Strength Ointment - Premier Value

Drug Facts

Active ingredients (in each gram)

Bacitracin zinc, USP 500 units Neomycin 3.5 mg Polymyxin B sulfate, USP 10,000 units Pramoxine hydrochloride, USP 10 mg

Purposes

First aid antibiotic

Pain reliever

Uses

First aid to help prevent infection and for the temporary relief of pain 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 one week
- condition persists or gets worse
- rash or other allergic reaction develops

 symptoms persist for more than one week, or clear up and occur again within a few days

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

- Adults and children 2 years of age and older
 - clean 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 15°-30°C (59°-86°F)
- Before using any medication, read all label directions. Keep carton, it contains important information.

Inactive ingredient

white petrolatum

Questions or comments?

Call 1-877-753-3935 Monday-Friday 9AM-5PM EST

Principal display panel

COMPARE TO THE ACTIVE INGREDIENTS IN NEOSPORIN® + PAIN RELIEF OINTMENT* maximum strength

Triple Antibiotic

+ PAIN RELIEF

Bacitracin Zinc • Neomycin Sulfate • Polymyxin B Sulfate • Pramoxine HCl

Soothe Painful Cuts, Burns and Scrapes

24 Hour Infection Protection

Dual Action

*This product is not manufactured or distributed by Johnson & Johnson Consumer, Inc., Distributor of Neosporin® + Pain Relief Ointment.

Distributed by: Pharmacy Value Alliance, LLC 407 East Lancaster Avenue, Wayne, PA 19087

Package label



maximum strength

Triple Antibiotic

+ PAIN RELIEF

Bacitracin Zinc • Neomycin Sulfate • Polymyxin B Sulfate • Pramoxine HCI



Soothes Painful Cuts, Burns and Scrapes 24 Hour Infection Protection Dual Action



maximum strength

Triple Antibiotic

+ PAIN RELIEF

Bacitracin Zinc • Neomycin Sulfate • Polymyxin B Sulfate • Pramoxine HCI

NET WT. 1 OZ (28 g)



Distributed by: Pharmacy Value Alliance, LLC 407 East Lancaster Avenue, Wayne, PA 19087

Product of India



If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.

Code: GO/DRUGS/362 ADP3A/02 PLD-B692B FC007304



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SprinsW For external use only.	Directions ■ souts and children 2 years of age and older ■ shows a small amount of this product (an amount equal to the surface area 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
Uses Rist aid to help prevent infection and for the temporary relief of pain in minor m cuts m scrapes m burns	
Meomycin 3.5 mg 2.6 mg last aud antibiodic Polymycin S sulfate, USP 10,000 units Pramoxine hydrochloride, USP 20 mg	
Active ingredients (in each gram) Bactive ingredients (in each gram) Bactive ingredients (in each gram)	Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.
Drug Facts	Drug Facts (continued)

PREMIER VALUE Maximum Strength Triple Antibiotic + Pain Relief

TRIPLE ANTIBIOTIC PLUS PAIN RELIEF MAXIMUM STRENGTH

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

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-715
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RW052I)	BACITRACIN	500 [USP'U] in 1 g	
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:116QD7X297)	NEOMYCIN	3.5 mg in 1 g	
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII: J2VZ 07J96K)	POLYMYXIN B	10000 [USP'U] 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)	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-715- 01	1 in 1 CARTON	07/10/2015	
,		28 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	07/10/2015	

Labeler - Chain Drug Consortium, LLC (101668460)

Revised: 3/2024 Chain Drug Consortium, LLC