

TRIPLE ANTIBIOTIC PLUS PAIN RELIEF- bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment
Chain Drug Marketing Association Inc.

Quality Choice Triple Antibiotic Ointment Plus Pain Relief

Active ingredients (each gram contains)

Bacitracin zinc 500 units

Neomycin sulfate 3.5 mg

Polymyxin B sulfate 10,000 units

Purpose

First aid antibiotic

Active Ingredient

Pramoxine HCL 10mg

Purpose

External Analgesic

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Uses first aid to help prevent infection in minor: ● cuts ● scrapes ● burns

Warnings For external use only.

Do not use ● 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 ● condition persists or gets worse

● you need to use longer than 1 week

● a rash or other allergic reaction develops

Directions

- 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.

Inactive ingredient: Petrolatum

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.

Other Information

Distributed by C.D.M.A., Inc.

43157 W 9 Mile Rd

Novi, MI 48375

www.qualitychoice.com

questions: 800-935-2362

Product of PRC

Packaging

OUTSIDE BOX





INNER TUBE



TRIPLE ANTIBIOTIC PLUS PAIN RELIEF

bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83324-264
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)			BACITRACIN	500 [USP'U] 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: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)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-264-01	1 in 1 BOX	08/28/2025	
1		28.3 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	08/28/2025	

Labeler - Chain Drug Marketing Association Inc. (011920774)

Registrant - Trifecta Pharmaceuticals USA LLC. (079424163)