QUALITY CHOICE MAXIMUM STRENGTH TRIPLE ANTIBIOTIC- bacitracin zinc, neomycin, polymyxin b sulfate, pramoxine hydrochloride ointment Chain Drug Marketing Association Inc

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.

Active Ingredients (in each gram) Purpose

Polymyxin B sulfate 10,000 units...... First aid antibiotic

Pramoxine hydrochloride 10 mg...... First aid analgesic

Uses

Helps prevent infection and temporarily relieves pains due to:

- minor cuts
- scrapes
- burns

Warnings

For external use only

Allergy alert: do not use if allergic to any of the ingredients

Do not use

- in or near the eyes
- over large areas of the body
- longer than 1 week

Ask a doctor before use in case of deep or puncture wounds, animal bites, or serious burns

Stop use and ask a doctor if:

- condition persists or gets worse
- symptoms last for more than 7 days or clear up and comes back within a few days
- a rash or other allergic reaction occurs

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

Directions

Adults and children 2 years of age and older:

- clean the affected area
- apply a small amount equal to surface area of a finger tip, on 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 20° to 25°C (68° to 77°F)
- Lot No. & Exp. Date: see box or crimp or tube

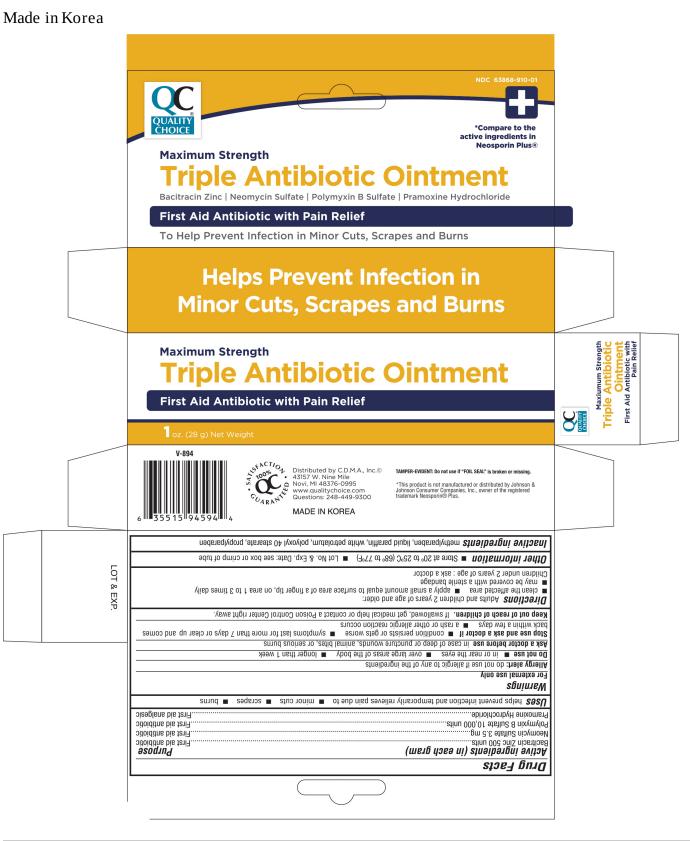
Inactive ingredients

methylparaben, liquid paraffin, white petrolatum, polyoxyl 40 stearate, proplyparaben **DISTRIBUTED BY:**

C.D.M.A., INC.

43157 W. NINE MILE

NOVI, MI 48376-0995 USA



QUALITY CHOICE MAXIMUM STRENGTH TRIPLE ANTIBIOTIC

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

Product Information	oduct Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-910			
Route of Administration	TOPICAL					

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
BACITRACIN ZINC (UNII: 89 Y4M234ES) (BACITRACIN - UNII:58 H6 RWO 52I)	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:J2VZ07J96K)	POLYMYXIN B	10000 [USP'U] in 1 g		
PRAMO XINE HYDRO CHLO RIDE (UNII: 88 AYB867L5) (PRAMO XINE - UNII: 068 X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 g		

Inactive Ingredients			
Ingredient Name	Strength		
METHYLPARABEN (UNII: A218 C7H19 T)			
PARAFFIN (UNII: 19 O 0 E 3 H 2 Z E)			
PETROLATUM (UNII: 4T6 H12BN9 U)			
POLYOXYL 40 STEARATE (UNII: 13A4J4NH9I)			
PROPYLPARABEN (UNII: Z8IX2SC1OH)			

]	Packaging				
#	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:63868-910-01	1 in 1 CARTON	10/12/2016		
1		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 final	part333B	03/25/2014		

Labeler - Chain Drug Marketing Association Inc (011920774)

Revised: 10/2016 Chain Drug Marketing Association Inc