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 400 units

Neomycin sulfate 3.5 mg

Polymyxin B sulfate 5,000 units

Purpose

First aid antibiotic

Active Ingredient

Pramoxine HCL

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

● 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● 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 **Light** Mineral Oil,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 CDMA., Inc.

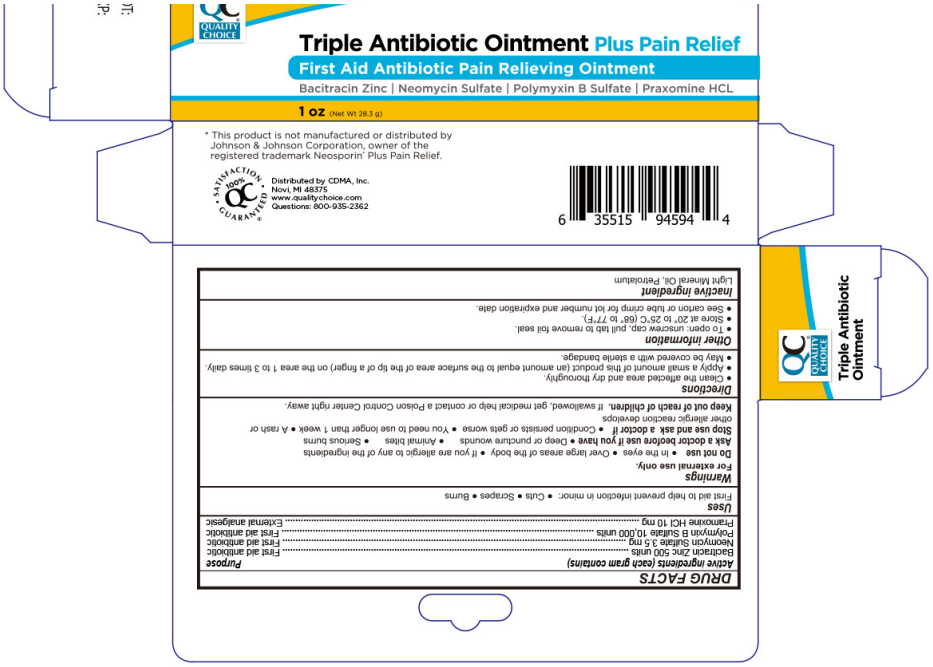
Novi, MI 48375

www.qualitychoice.com

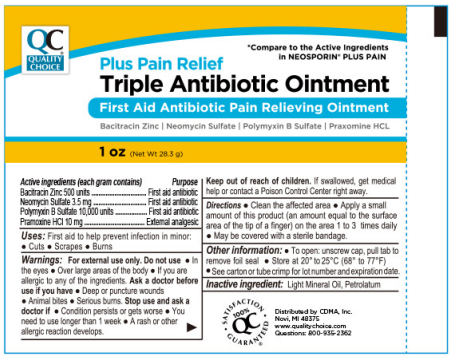
questions: 800-935-2362

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:83324-036
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:J2VZ07J96K)	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
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-036-01	1 in 1 BOX	02/28/2024	
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	02/28/2024	

Labeler - Chain Drug Marketing Association Inc. (011920774)

Registrant - Trifecta Pharmaceuticals USA LLC. (079424163)

Revised: 2/2024 Chain Drug Marketing Association Inc.