QUALITY CHOICE MAXIMUM STRENGTH TRIPLE ANTIBIOTIC- bacitracin zinc, neomycin sulfate, polymyxin b sulfate, and 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.

QC Max Strength Triple Antibiotic Ointment 1 oz. 94594 CMI 2019

Active ingredient (in each gram) Purposes

Bacitracin zinc 500 units.....First aid antibiotic

Neomycin sulfate 3.5 mg.....First aid antibiotic

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

Pramoxine HCI 10 mg.....External analgesic

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 or nose
- 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
- rash or other allergic reaction develops
- symptoms persist for more than 1 week, or clear up and occur again within a few days
- condition persists or gets worse
- redness, irritation, swelling, or pain persists or increases

If pregnant or breast-feeding, ask a health professional before use

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 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

- children under 2 years of age: ask a doctor

Other information

- store at room temperature 20-25°C (68-77°F)

Inactive ingredients

cetyl alcohol, light mineral oil, polyoxyl 40 stearate, stearic acid, white petrolatum

Distributed by:

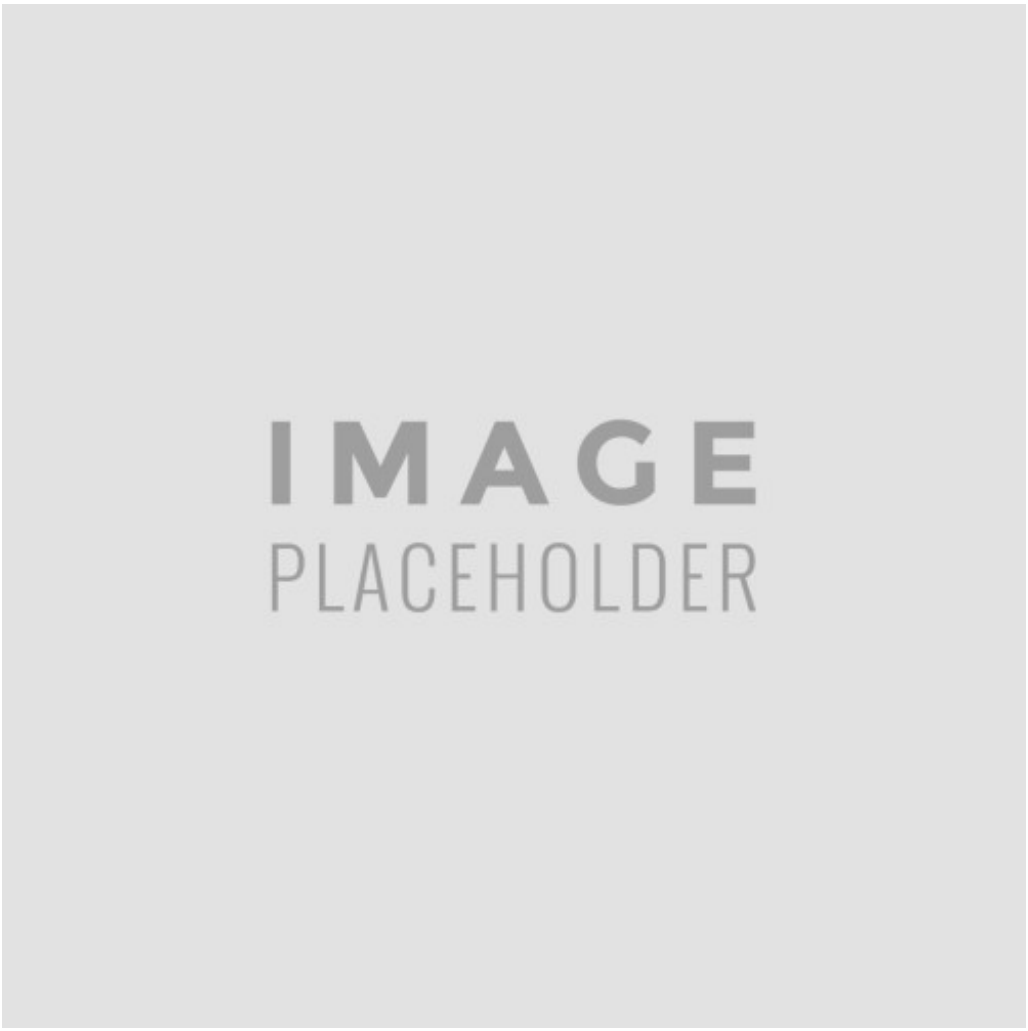
C.D.M.A., Inc.

43157 W 9 Mile Rd

Novi, MI 48375

www.qualitychoice.com

Made in South Korea



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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-009
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN SULFATE	3.5 mg in 1 g
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [USP'U] in 1 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	10000 [USP'U] in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
POLYOXYL 40 STEARATE (UNII: 13A4J4NH9I)	
PETROLATUM (UNII: 4T6H12BN9U)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
CETYL ALCOHOL (UNII: 936JST6JCN)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63868-009-28	1 in 1 CARTON	12/19/2019	
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	12/19/2019	

Labeler - Chain Drug Marketing Association Inc (011920774)

Revised: 12/2019 Chain Drug Marketing Association Inc