

**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.*

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**Active Ingredients (in each gram)**

**Purpose**

|                                       |                      |
|---------------------------------------|----------------------|
| Bacitracin zinc 500 units.....        | First aid antibiotic |
| Neomycin 3.5 mg.....                  | First aid antibiotic |
| 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 Poison 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, propylparaben


DISTRIBUTED BY:

C.D.M.A., INC.


43157 W. NINE MILE

NOVI, MI 48376-0995 USA

Made in Korea



NDC 63868-910-01



**Maximum Strength**  
**Triple Antibiotic Ointment**  
 Bacitracin Zinc | Neomycin Sulfate | Polymyxin B Sulfate | Pramoxine Hydrochloride

**First Aid Antibiotic with Pain Relief**

To Help Prevent Infection in Minor Cuts, Scrapes and Burns


**Helps Prevent Infection in  
Minor Cuts, Scrapes and Burns**

**Maximum Strength**  
**Triple Antibiotic Ointment**  
**First Aid Antibiotic with Pain Relief**


Maximum Strength  
**Triple Antibiotic Ointment**  
 First Aid Antibiotic with  
 Pain Relief

**1** oz. (28 g) Net Weight

V-894



6 35515 94594 4

100%  
  
 SATISFACTION  
 GUARANTEED

Distributed by C.D.M.A., Inc. ©  
 43157 W. Nine Mile  
 Novi, MI 48376-0995  
 www.qualitychoice.com  
 Questions: 248-449-9300

**TAMPER-EVIDENT:** Do not use if "FOIL SEAL" is broken or missing.  
 \*This product is not manufactured or distributed by Johnson & Johnson Consumer Companies, Inc., owner of the registered trademark Neosporin® Plus.

MADE IN KOREA

LOT & EXP.

**Drug Facts**  
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 Neomycin Sulfate 3.5 mg.....First aid antibiotic  
 Polymyxin B Sulfate 10,000 units.....First aid antibiotic  
 Pramoxine Hydrochloride.....First aid analgesic

**Uses**  
 Helps prevent infection and temporarily relieves pain due to minor cuts, scrapes, and burns.

**Warnings**  
 For external use only.  
 Allergy alert: do not use if allergic to any of the ingredients.  
 Do not use if 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.

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# QUALITY CHOICE MAXIMUM STRENGTH TRIPLE ANTIBIOTIC

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

## Product Information

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:63868-910 |
| <b>Route of Administration</b> | TOPICAL        |                           |               |

## Active Ingredient/Active Moiety

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

## Inactive Ingredients

| Ingredient Name                                | Strength |
|--|----------|
| <b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)        |          |
| <b>PARAFFIN</b> (UNII: I9O0E3H2ZE)             |          |
| <b>PETROLATUM</b> (UNII: 4T6H12BN9U)           |          |
| <b>POLYOXYL 40 STEARATE</b> (UNII: 13A4J4NH9I) |          |
| <b>PROPYLPARABEN</b> (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)