

**REDICARE TRIPLE ANTIBIOTIC FIRST AID- triple antibiotic first aid ointment ointment**  
**Redicare LLC**

*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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**Triple Antibiotic**

**Active Ingredient**

- Bacitracin Zinc 400 units
- Neomycin sulfate (3.5 mg Neomycin)
- Polymyxin B Sulfate 5000 units

**Purpose**

First Aid Antibiotics

**Uses**

first aid to help prevent infection in minor cuts, scrapes, and burns

**Warnings**

**For external use only**

**Do not use**

- internally
- in eyes
- over large areas of the body or on puncture wounds, animal bites or serious burns
- for more than 1 week unless directed by a doctor
- if you are allergic to any of the ingredients

**Stop use and ask a doctor if**

- a rash or allergic reaction develops
- condition worsens or persists

**Keep out of reach of children**

If ingested, contact a Poison Control Center right away

**Directions**

- clean affected area
- apply a small amount 1 to 3 times daily
- may cover with a sterile bandage

**Inactive Ingredients**

petrolatum



redicare.  
Prepared. The only solution.

**TRIPLE ANTIBIOTIC  
FIRST AID OINTMENT**

NDC 71105-5601-1

**0.5 g**

**Drug Facts**

**Active Ingredient Purpose**

(in each gram)

Bacitracin zinc 400 units  
Neomycin sulfate (3.5mg Neomycin)  
Polymyxin B sulfate First Aid  
5000 units.....Antibiotics

**Uses** first aid to help  
prevent infection in minor  
cuts, scrapes, and burns

**Warnings**

For external use only ▶

**Drug Facts (continued)**

**Warnings (continued)**

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# REDICARE TRIPLE ANTIBIOTIC FIRST AID

triple antibiotic first aid ointment ointment

## Product Information

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:71105-560 |
| <b>Route of Administration</b> | TOPICAL        |                           |               |

## Active Ingredient/Active Moiety

| Ingredient Name  | Basis of Strength | Strength            |
|--|-------------------|---------------------|
| BACITRACIN ZINC (UNII: 89 Y4M234ES) (BACITRACIN - UNII:58 H6RWO52I)    | BACITRACIN        | 400 [USP'U] in 1 g  |
| NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:H16QD7X297)       | NEOMYCIN          | 3.5 mg in 1 g       |
| POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K) | POLYMYXIN B       | 5000 [USP'U] in 1 g |

## Inactive Ingredients

| Ingredient Name               | Strength |
|-------------------------------|----------|
| PETROLATUM (UNII: 4T6H12BN9U) |          |

## Product Characteristics

|              |       |              |  |
|--------------|-------|--------------|--|
| <b>Color</b> | white | <b>Score</b> |  |
| <b>Shape</b> |       | <b>Size</b>  |  |

| <b>Flavor</b>                |  | <b>Imprint Code</b>                                  |                      |                    |
|------------------------------|--|--|----------------------|--------------------|
| <b>Contains</b>              |  |  |                      |                    |
| <b>Packaging</b>             |  |  |                      |                    |
| #                            | Item Code                                | Package Description                                  | Marketing Start Date | Marketing End Date |
| 1                            | NDC:71105-560-12                         | 25 in 1 BOX  | 10/08/2019           |                    |
| 1                            | NDC:71105-560-11                         | 0.5 g in 1 PACKET; Type 0: Not a Combination Product |                      |                    |
| <b>Marketing Information</b> |  |  |                      |                    |
| Marketing Category           | Application Number or Monograph Citation | Marketing Start Date                                 | Marketing End Date   |                    |
| OTC monograph final          | part333B                                 | 10/08/2019   |                      |                    |

**Labeler** - Redicare LLC (800149346)

Revised: 10/2019

Redicare LLC