**DRS. PHARMACY TRIPLE ANTIBIOTIC PLUS- bacitracin zinc, neomycin sulfate, polymyxin b sulfate, lidocaine ointment**
**OL PHARMA TECH, LLC Drs PHARMACY**

*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

Bacitracin zinc 400 units
Neomycin 3.5 mg
Polymyxin B sulfate 5,000 units

Pramoxine hydrochloride 10 mg

## Uses

first aid to help prevent infection in minor
• cuts
• scrapes
• burns

## Purpose

- first aid antibiotic
- External Analgesic

## Do Not Use

Do not use
• if you are allergic to any of the ingredients
• in the eyes
• over large areas of the body
• longer than 1 week unless directed by a doctor

For external use only

## Ask a doctor before use if you have

• deep or puncture wounds
• animal bites
• serious burns

Stop use and ask a doctor if
• the condition persists or gets worse
• a rash or other allergic reaction develops

KEEP OUT OF REACH OF CHILDREN

If swallowed, get medical help or contact a Poison Control Center right away.

DIRECTIONS

• 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

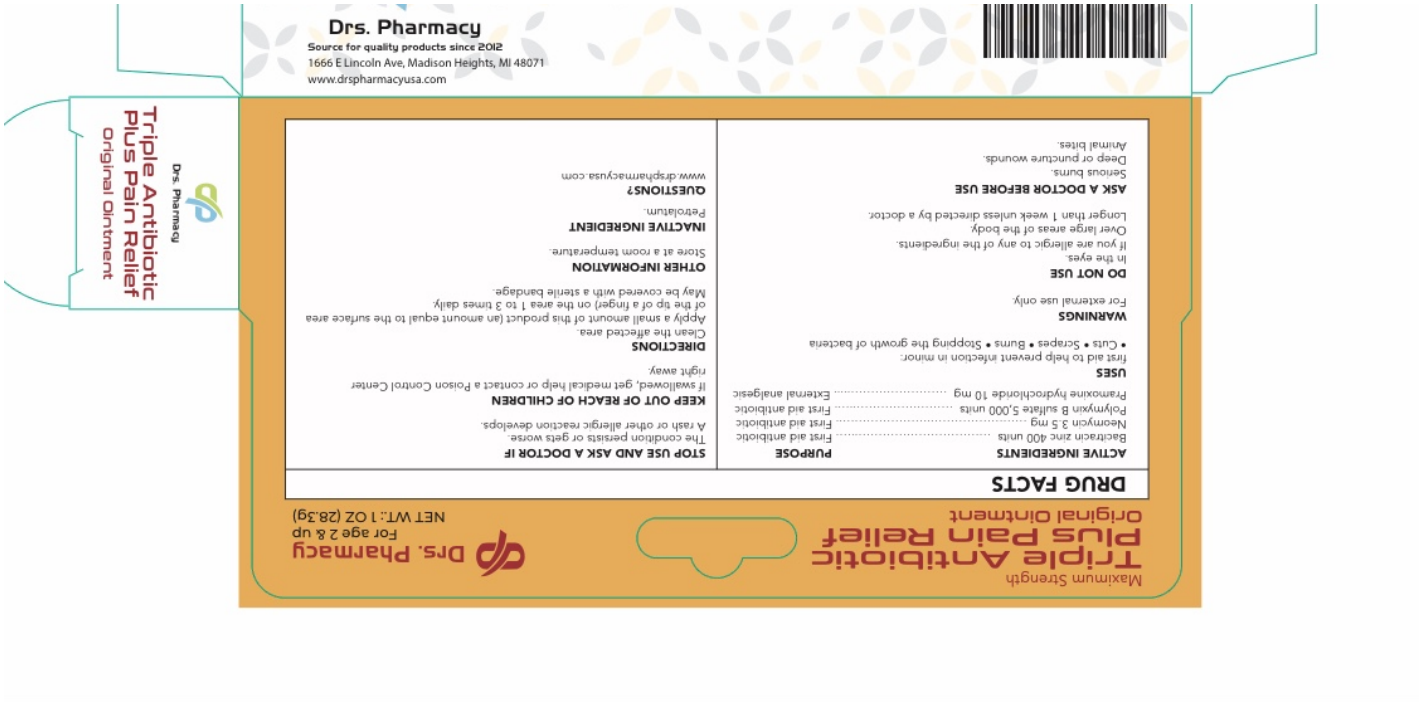
**OTHER INFORMATION**

Store at room temperature

**Inactive Ingredient**

Petrolatum

www.drspharmacyusa.com




# DRS. PHARMACY TRIPLE ANTIBIOTIC PLUS

bacitracin zinc, neomycin sulfate, polymyxin b sulfate, lidocaine ointment

## Product Information

| | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:80489-757 |
| **Route of Administration** | TOPICAL | | |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **BACITRACIN ZINC** (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I) | BACITRACIN | 400 [USP'U] in 1000 mg |
| **NEOMYCIN SULFATE** (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297) | NEOMYCIN | 3.5 mg in 1000 mg |
| **POLYMYXIN B SULFATE** (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K) | POLYMYXIN B | 5000 [USP'U] in 1000 mg |
| **PRAMOXINE HYDROCHLORIDE** (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056) | PRAMOXINE HYDROCHLORIDE | 10 mg in 1000 mg |

## Inactive Ingredients

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

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | white | **Score** | |

| | | | |
|---|---|---|---|
| **Shape** | | **Size** | |
| **Flavor** | | **Imprint Code** | |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:80489-757-01 | 1 in 1 CARTON | 09/05/2021 | |
| **1** | | 14000 mg in 1 TUBE; Type 0: Not a Combination Product | | |
| **2** | NDC:80489-757-02 | 1 in 1 CARTON | 09/05/2021 | |
| **2** | | 28300 mg 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 | 09/05/2021 | |

## Labeler - OL PHARMA TECH, LLC Drs PHARMACY (021170377)

## Registrant - OL PHARMA TECH, LLC Drs PHARMACY (021170377)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| OL PHARMA TECH, LLC Drs PHARMACY | | 021170377 | manufacture(80489-757) |