**ANTI-ITCH- zinc acetate , diphenhydramine cream**
**OL PHARMA TECH, LLC**

---

**Active Ingredients**

Diphenhydramine hydrochloride 1%

Zinc Acetate 0.1%

**Purpose**

Topical analgesic

skin protectant

**uses**

temporarily relieves pain and itching associated with:

- insect bites
- minor burns
- sunburn
- minor skin irritations
- minor cuts
- scrapes
- rashes due to poison ivy, poison oak, and poison sumac
- dries the oozing and weeping of poison ivy, poison oak and poison sumac

**warnings**

For external use only

**Ask a doctor before use**

- on chicken pox
- on measles

**Do Not Use**

- on large areas of the body
- with any other product containing diphenhydramine, even one taken by mouth

**When using this product**

When using this product avoid contact with eyes

**Stop use and ask a doctor if**

- condition worsens or does not improve within 7 days
- symptoms persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

**Directions**

Do not use more than directed:
- adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 2 years of age: ask a doctor

protect from excessive heat (40°C/104°F)

**Inactive Ingredients**

cetostearyl alcohol, sodium cetostearyl sulfate, stearic acid, trolamine, mineral oil, propylene glycol, water, methyl paraben, propyl paraben, EDTA, vitamin E

**Questions**

drspharmacyusa.com

**Directions**

do not use more than directed:
- adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 2 years of age: ask a doctor


Itch stopping cream

# ANTI-ITCH

zinc acetate , diphenhydramine cream

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:80489-005 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **DIPHENHYDRAMINE HYDROCHLORIDE** (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M) | DIPHENHYDRAMINE HYDROCHLORIDE | 1 g in 100 g |
| **ZINC ACETATE** (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37) | ZINC CATION | 0.1 g in 100 g |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **DECYL OLEATE** (UNII: ZGR06DO97T) | |
| **TROLAMINE** (UNII: 9O3K93S3TK) | |

| | |
|---|---|
| **EDETATE DISODIUM** (UNII: 7FLD91C86K) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **PROPYLPARABEN** (UNII: Z8IX2SC1OH) | |
| **METHYLPARABEN** (UNII: A2I8C7HI9T) | |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |
| **MINERAL OIL** (UNII: T5L8T28FGP) | |
| **SODIUM CETOSTEARYL SULFATE** (UNII: 7ZBS06BH4B) | |
| **CETOSTEARYL ALCOHOL** (UNII: 2DMT128M1S) | |
| **STEARIC ACID** (UNII: 4ELV7Z65AP) | |
| **.ALPHA.-TOCOPHEROL** (UNII: H4N855PNZ1) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | white | **Score** | |
| **Shape** | | **Size** | |
| **Flavor** | | **Imprint Code** | |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:80489-005-01 | 1 in 1 CARTON | 01/01/2021 | |
| **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 | M016 | 01/01/2021 | |

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

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

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| OL PHARMA TECH, LLC Drs PHARMACY | | 021170377 | manufacture(80489-005) |

Revised: 1/2025 OL PHARMA TECH, LLC