**ANTI-ITCH FEMININE CARE- hydrocortisone 0.5% cream**
**OL PHARMA TECH, LLC Drs PHARMACY**

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## ACTIVE INGREDIENT

Hydrocortisone, USP 0.5%

## PURPOSE

Anti-itch

## USES

For temporary external feminine itching

## WARNINGS

For External Use Only

Avoid Contact with eyes

## DIRETIONS

Adults and children 12 years and older Apply to external vaginal area not more than 3 to 4 times a day
Children under 12 years Consult a doctor

## Stop use and ask a doctor if

- you have a vaginal discharge. Consult a physician.
- condition worsens, or if symptoms persist for more than 7 days, or clear up and occur again in a few days.

## KEEP OUT OF REACH OF CHILDREN

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

## OTHER INFORMATION

- store at 20°-25°C (68°-77°F)

## INACTIVE INGREDIENT

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

**QUESTIONS**

www.drspharmacyusa.com

**PACKAGE LABEL**


# ANTI-ITCH FEMININE CARE

hydrocortisone 0.5% cream

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:80489-103 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **HYDROCORTISONE** (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ) | HYDROCORTISONE | 5 mg in 1 g |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |
| **PROPYLPARABEN** (UNII: Z8IX2SC1OH) | |
| **.ALPHA.-TOCOPHEROL** (UNII: H4N855PNZ1) | |
| **METHYLPARABEN** (UNII: A2I8C7HI9T) | |
| **EDETATE DISODIUM** (UNII: 7FLD91C86K) | |
| **CETEARETH-12** (UNII: 7V4MR24V5P) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **STEARIC ACID** (UNII: 4ELV7Z65AP) | |
| **SODIUM CETOSTEARYL SULFATE** (UNII: 7ZBS06BH4B) | |
| **CETOSTEARYL ALCOHOL** (UNII: 2DMT128M1S) | |
| **OCTYLDODECANOL** (UNII: 461N1O614Y) | |
| **TROLAMINE** (UNII: 9O3K93S3TK) | |

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

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:80489-103-01 | 1 in 1 CARTON | 02/01/2022 | |
| **1** | | 20 g in 1 TUBE; Type 0: Not a Combination Product | | |
| **2** | NDC:80489-103-02 | 1 in 1 CARTON | 02/01/2022 | |
| **2** | | 30 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 | M005 | 02/01/2022 | |

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

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

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| OHIO LAB PHARMA | | 021170377 | manufacture(80489-103) |

Revised: 1/2025 OL PHARMA TECH, LLC Drs PHARMACY