

**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

**DIRECTIONS**

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.

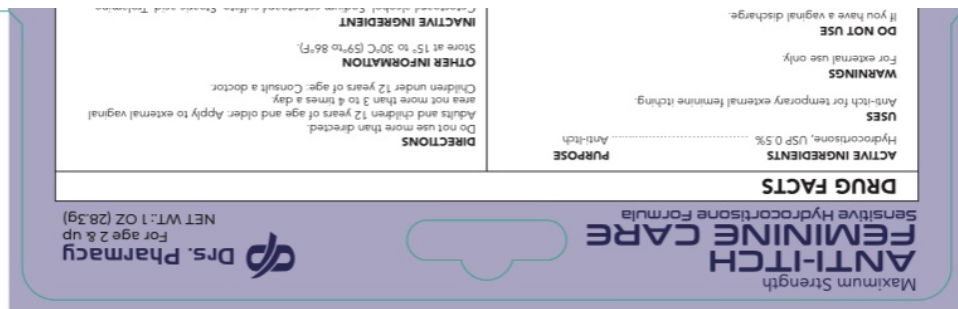
**OTHER INFORMATION**

- 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





## ANTI-ITCH FEMININE CARE

hydrocortisone 0.5% cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:80489-103
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>HYDROCORTISONE</b> (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ)	HYDROCORTISONE	5 mg in 1 g

### Inactive Ingredients

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

### 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>			

## 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: 10/2023

OL PHARMA TECH, LLC Drs PHARMACY