## DRS. ACNE CLEAR- salicylic acid gel OL PHARMA TECH, LLC Drs. PHARMACY

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## **ACTIVE INGREDIENT**

Salicylic Acid 0.5% w/w.

### PURPOSE

Acne treatment

#### USES

For the treatment of acne. Dries and clears acne pimples, blackheads and whiteheads and allows skin to heal.

## DO NOT USE

if you have sensitive skin or are sensitive to salicylic acid

### DIRECTIONS

Morning and evening, after a thorough cleansing of the skin, apply acne clear gel locally on cutanous imperfections. Then apply the usual day or night cream. Renew application 1 to 3 times daily.

#### WARNINGS

**For external use only**. Using other topical acne medications at the same time or immediately following use of this product may increase dryness or irritation of the skin. If this occurs, only one medication should be used unless directed by a doctor. Avoid direct contact with the eyes. If product gets into the eyes, rinse liberally with water. Discontinue use if skin irritation develops or increases. If irritation persists, consult a doctor.

## WHEN USING THIS PRODUCT

Avoid contact with eyes, lips, and mouth.

## **KEEP OUT OF REACH OF CHILDREN**

if swallowed, get medical help or contact poison control center right away

carbomer interpolymer type A, trolamine, vitamin E, propylene glycol, water, isopropyl alcohol, methyl paraben, EDTA, propyl paraben, DMDM hydantoin

## **OTHER INFORMATION**

- store at 15-30 C (59-86 F)
- close cap tightly after use
- keep away from heat

# QUESTIONS

www.drspharmacyusa.com

# PACKAGE LABEL





| DRS. ACNE<br>salicylic acid gel   |                  |                           |                              |                      |          |                             |  |  |  |
|---|------------------|---------------------------|------------------------------|----------------------|----------|-----------------------------|--|--|--|
|   |                  |                           |                              |                      |          |                             |  |  |  |
| <b>Product Info</b>   | rmation          |                           |                              |                      |          |                             |  |  |  |
| Product Type  |                  | HUMAN OTC DRUG            | NOTC DRUG Item Code (Source) |                      |          | NDC:80489-300               |  |  |  |
| Route of Admir  | nistration       | TOPICAL                   |                              |                      |          |                             |  |  |  |
| Activo Ingro  | liont/Activo     | Majaty                    |                              |                      |          |                             |  |  |  |
| Active Ingredient/Active Moiety   |                  |                           |                              |                      |          |                             |  |  |  |
| Ingredient Name<br>SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ) |                  |                           |                              | Basis of Strength    |          | <b>Strength</b> 5 mg in 1 g |  |  |  |
|   |                  |                           |                              |                      |          |                             |  |  |  |
| Inactive Ingredients  |                  |                           |                              |                      |          |                             |  |  |  |
|   |                  | Ingredient Name           |                              |                      |          | Strength                    |  |  |  |
|   |                  | E A (55000 CPS) (UNII: 59 | TL3WG5CO)                    |                      |          |                             |  |  |  |
| WATER (UNII: 059<br>PROPYLENE GLY   | . ,              | 167//3)                   |                              |                      |          |                             |  |  |  |
| .ALPHATOCOPH  |                  |                           |                              |                      |          |                             |  |  |  |
| METHYLPARABEN   |                  |                           |                              |                      |          |                             |  |  |  |
| EDETATE DISODI  | UM (UNII: 7FLD9: | 1C86K)                    |                              |                      |          |                             |  |  |  |
| TROLAMINE (UNII: 903K93S3TK)  |                  |                           |                              |                      |          |                             |  |  |  |
| ISOPROPYL ALCOHOL (UNII: ND2M416302)  |                  |                           |                              |                      |          |                             |  |  |  |
|   |                  |                           |                              |                      |          |                             |  |  |  |
| Packaging   |                  |                           |                              |                      |          |                             |  |  |  |
| # Item Code   | Pa               | ackage Description        |                              | Marketing St<br>Date | tart Mar | keting End<br>Date          |  |  |  |
| <b>1</b> NDC:80489-<br>300-01   | 1 in 1 CARTON    |                           |                              | 02/01/2022           |          |                             |  |  |  |

| M | arketing<br>Marketing<br>Category | Information<br>Application Number or Monograph<br>Citation         | Marketing Start<br>Date | Marketing End<br>Date |  |  |  |  |  |
|---|-----------------------------------|--|-------------------------|-----------------------|--|--|--|--|--|
| M | arketing                          | Information  |                         |                       |  |  |  |  |  |
|   | Marketing Information             |  |                         |                       |  |  |  |  |  |
|   |                                   |  |                         |                       |  |  |  |  |  |
| 2 |                                   | 30 g in 1 TUBE, WTH APPLICATOR; Type 0: Not a Combination Product  |                         |                       |  |  |  |  |  |
| 2 | NDC:80489-<br>300-02              | 1 in 1 CARTON  | 02/01/2022              |                       |  |  |  |  |  |
|   | (                                 | 20 g in 1 TUBE, WITH APPLICATOR; Type 0: Not a Combination Product |                         |                       |  |  |  |  |  |

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

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

| Establishment                    |         |           |                            |  |  |  |  |  |
|----------------------------------|---------|-----------|----------------------------|--|--|--|--|--|
| Name                             | Address | ID/FEI    | <b>Business Operations</b> |  |  |  |  |  |
| OL PHARMA TECH, LLC Drs PHARMACY |         | 021170377 | manufacture(80489-300)     |  |  |  |  |  |

Revised: 10/2023

OL PHARMA TECH, LLC Drs. PHARMACY