DRS. ACNE CLEAR- salicylic acid gel
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 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

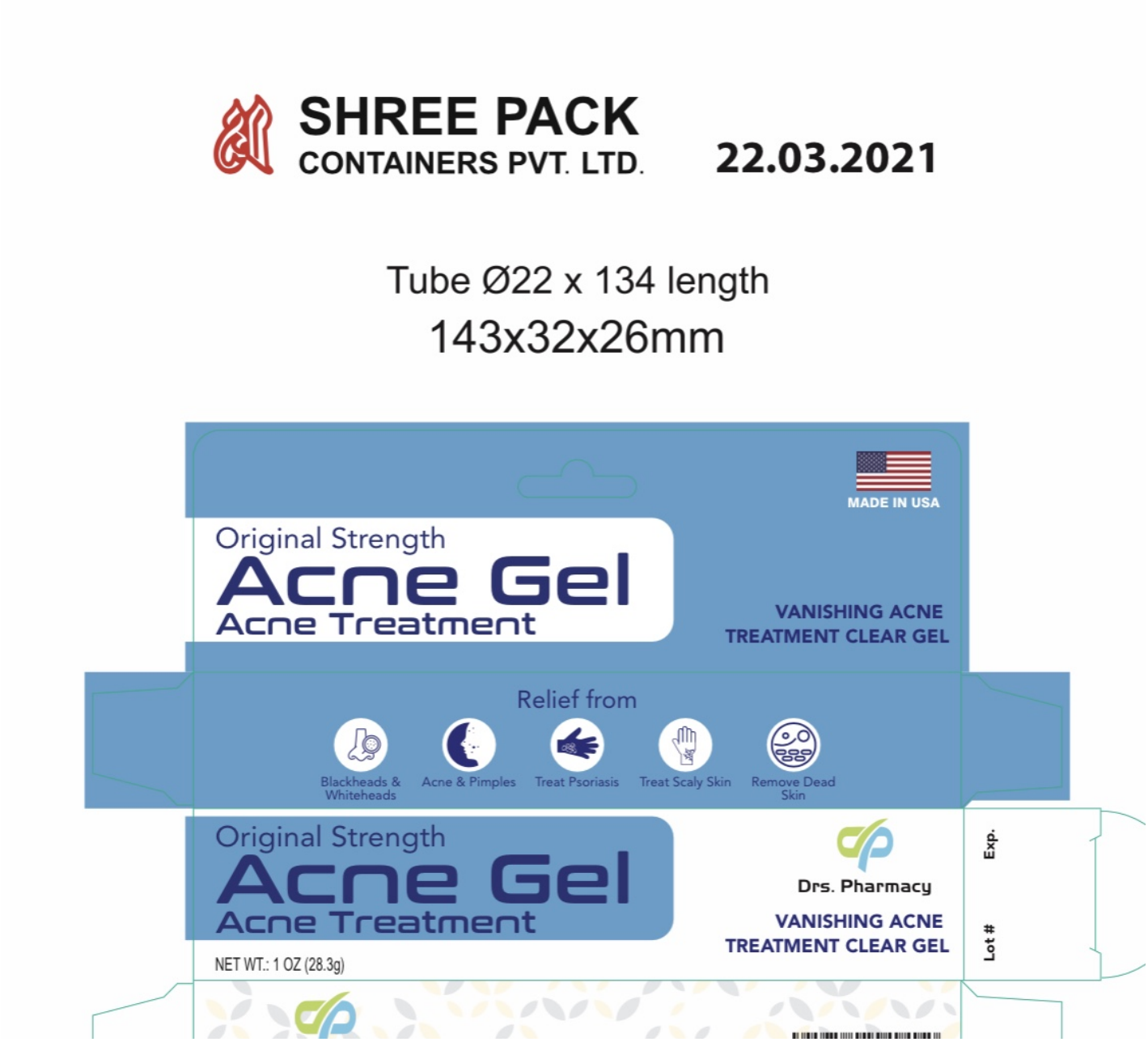
OTHER INFORMATION

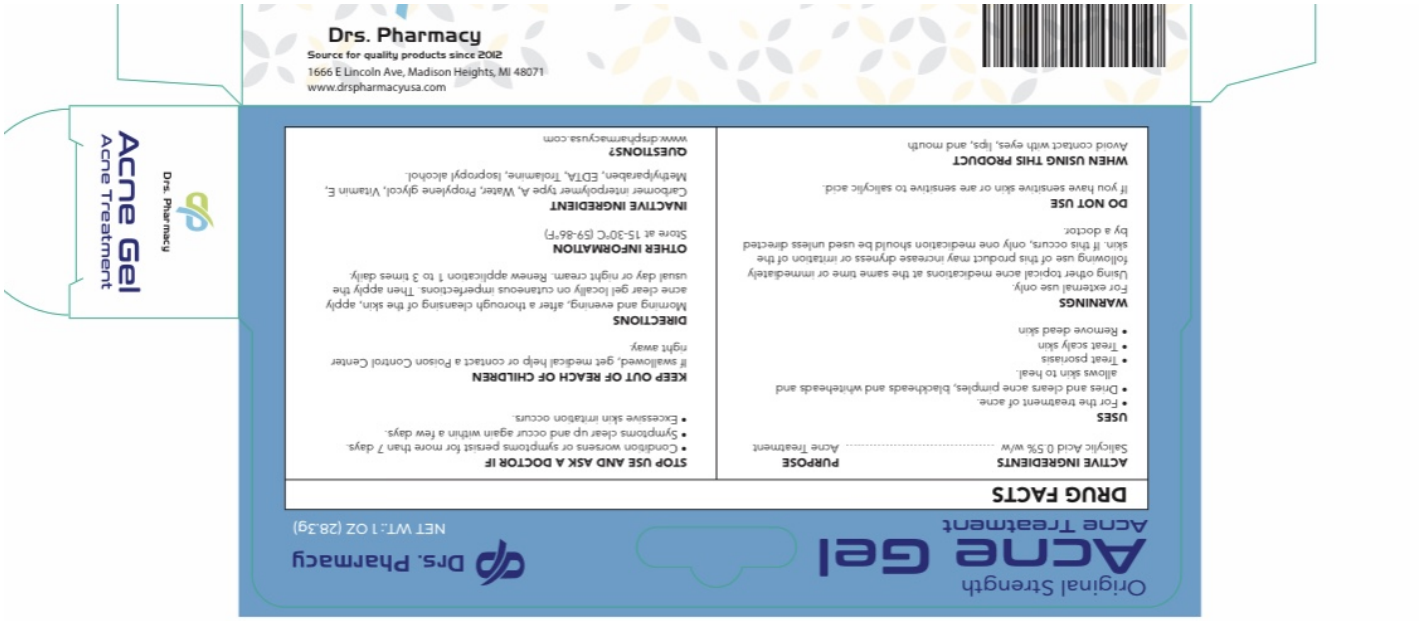
- store at 15-30 C (59-86 F)
- close cap tightly after use
- keep away from heat

QUESTIONS

www.drspharmacyusa.com

PACKAGE LABEL





DRS. ACNE CLEAR

salicylic acid gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80489-300
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	5 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
TROLAMINE (UNII: 9O3K93S3TK)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80489-300-01	1 in 1 CARTON	02/01/2022	
1		20 g in 1 TUBE, WITH APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	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
OL PHARMA TECH, LLC Drs PHARMACY		021170377	manufacture(80489-300)

Revised: 1/2023 OL PHARMA TECH, LLC Drs. PHARMACY