KERALYT 3 PERCENT- salicylic acid gel
Summers Laboratories Inc

SUMMERS LABS (as PLD) - KERALYT GEL 3% SALICYLIC ACID (11086-038)

ACTIVE INGREDIENT

SALICYLIC ACID 3%

PURPOSE

PSORIASIS, SEBORRHEIC DERMATITIS

USES

HELPS STOP THESE SYMPTOMS OF PSORIASIS, SEBORRHEIC DERMATITIS AND DANDRUFF

- FLAKING
- SCALING
- REDNESS
- IRRITATION
- ITCHING

WARNINGS

FOR EXTERNAL USE ONLY

ASK A DOCTOR BEFORE USE IF CONDITION COVERS A LARGE AREA OF THE BODY

WHEN USING THIS PRODUCT

- DO NOT GET INTO EYES. IF CONTACT OCCURS, RINSE EYES THOROUGHLY WITH WATER. IF IRRITATION PERSISTS, CONSULT A DOCTOR.

STOP USE AND CONSULT A DOCTOR IF CONDITION WORSENS OR DOES NOT IMPROVE AFTER REGULAR USE.

KEEP OUT OF REACH OF CHILDREN. IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER RIGHT AWAY.

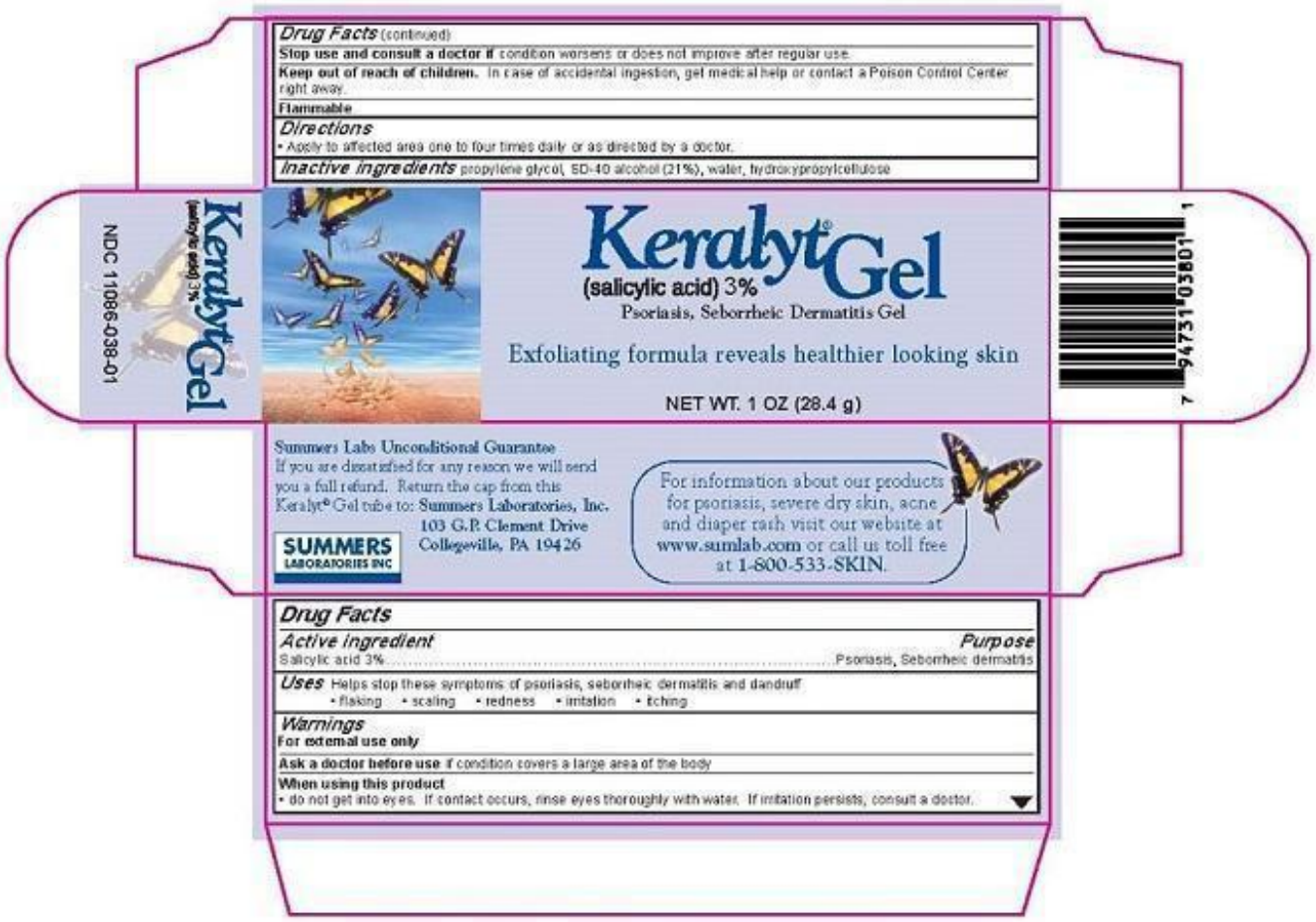
FLAMMABLE

DIRECTIONS

- APPLY TO AFFECTED AREA ONE TO FOUR TIMES DAILY OR AS DIRECTED BY A DOCTOR.

INACTIVE INGREDIENTS

PROPYLENE GLYCOL, SD-40 ALCOHOL (21%), WATER, HYDROXYPROPYLCELLULOSE



KERALYT 3 PERCENT

salicylic acid gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11086-038
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	3 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALCOHOL (UNII: 3K9958V90M)	
HYDROXYPROPYL CELLULOSE (UNII: RFW2ET671P)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11086-038-01	28.4 g in 1 TUBE; Type 0: Not a Combination Product	10/30/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	10/30/2013	

Labeler - Summers Laboratories Inc (002382612)

Revised: 10/2023 Summers Laboratories Inc