ACNE CLEANSING- salicylic acid gel
CLINIQUE LABORATORIES LLC

ACNE SOLUTIONS CLEANSING GEL

Drug Facts

Active Ingredient

Salicylic Acid 2%

Purpose

Acne Treatment

Uses

- treats acne
- clears acne blemishes
- helps prevent development of new acne blemishes

Warnings

For external use only

When using this product skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne medication at a time.

Keep out of reach of children.

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

Directions

- Wet hands and work into a lather
- Add water to increase lather
- Massage gently over wet skin, avoiding eye area
- Rinse
- Use A.M. and P.M.
- If bothersome drying or peeling occurs, reduce usage to every other day
- Follow with Acne Solutions Clarifying Lotion

Inactive ingredients

water\aqua\eau [] glycerin [] sodium laureth sulfate [] sodium chloride [] lauramidopropyl betaine [] butylene glycol [] sucrose [] gentiana lutea (gentian) root extract [] laminaria saccharina extract [] caffeine [] sodium hyaluronate [] acetyl glucosamine [] peg-120

methyl glucose dioleate [] benzophenone-4 [] sodium hydroxide [] tetrasodium edta [] disodium edta [] bht [] phenoxyethanol [] blue 1 (ci 42090) <iln50388>

Other information

Protect the product in this container from excessive heat and direct sun

PRINCIPAL DISPLAY PANEL - 125 mL Tube Carton

CLINIQUE

acne solutions

cleansing gel

SALICYLIC ACID

ACNE MEDICATION

ALL SKIN TYPES

4.2 FL.OZ.LIQ./125 ml e

ACNE CLEANSING

salicylic acid gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49527-045
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	20 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
LAURAMIDOPROPYL BETAINE (UNII: 23D6XVI233)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
SUCROSE (UNII: C151H8M554)	
GENTIANA LUTEA ROOT (UNII: S72O3284MS)	
SACCHARINA LATISSIMA (UNII: 68CMP2MB55)	
CAFFEINE (UNII: 3G6A5W338E)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
N-ACETYLGLUCOSAMINE (UNII: V956696549)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
SULISOBENZONE (UNII: 1W6L629B4K)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
EDETIC ACID (UNII: 9G34HU7RV0)	
EDETATE SODIUM (UNII: MP1J8420LU)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start**	**Marketing End**

#	Item Code	Package Description	Date	Date
1	NDC:49527-045-01	1 in 1 CARTON	10/01/2014	
1		125 mL 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	M006	10/01/2014	

Labeler - CLINIQUE LABORATORIES LLC (044475127)

Registrant - Estee Lauder Companies Inc. (790802086)

Establishment

Name	Address	ID/FEI	Business Operations
The Estee Lauder Inc		802599436	manufacture(49527-045) , pack(49527-045) , label(49527-045)

Revised: 8/2023 CLINIQUE LABORATORIES LLC