CLINIQUE ACNE SOLUTIONS CLEANSING FOAM- salicylic acid liquid
CLINIQUE LABORATORIES LLC

CLINIQUE ACNE SOLUTIONS CLEANSING FOAM

Active ingredient

Salicylic acid 1.5%

Purpose

Acne treatment

Use

- for the treatment of acne

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 this occurs, only use one topical acne medication at a time.

Keep out of reach of childeren.

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

Directions

- pump dispenser 2-3 times
- massage gently over wet skin, avoiding eye area
- rinse
- follow with Acne Solutions Clarifying Lotion
- if bothersome drying or peeling occurs, reduce usage to every other day
- because excessive drying of the skin may occur, start with one use daily, then gradually increase to twice a day if needed or as directed by a doctor
- after acne clears, continue using for preventative care

Inactive ingredients

water\aqua\eau,glycerin,sodium methyl cocoyl taurate,butylene glycol,sucrose,disodium phosphate,salicylic acid,sodium hyaluronate,phosphatidylcholine,caffeine,cola acuminata seed extract,algae extract,laminaria saccharina extract,lactobacillus ferment,stearamidopropyl dimethylamine,acetyl glucosamine,10-hydroxydecanoic

acid,arginine cocoate,peg/ppg-18/18 dimethicone,capryloyl glycine,ppg-6-decyltetradeceth-30,stearic acid,polyquaternium-7,disodium edta,phenoxyethanol,chloroxylenol,sodium benzoate <iln53446>

PRINCIPAL DISPLAY PANEL - 125 ml Bottle Label

CLINIQUE

acne solutions

cleansing

foam

STEP 1

SALICYLIC ACID ACNE MEDICATION

4.2 FL.OZ.LIQ./125 ml e

CLINIQUE ACNE SOLUTIONS CLEANSING FOAM

salicylic acid liquid

Product Information			

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49527-119
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	15 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
PHOSPHATIDYLCHOLINE, SOYBEAN (UNII: 1T6N4D9YV6)	
PORPHYRIDIUM PURPUREUM (UNII: K2P8K2558N)	
PEG/PPG-18/18 DIMETHICONE (UNII: 9H0AO7T794)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
SODIUM METHYL COCOYL TAURATE (UNII: JVL98CG53G)	
SUCROSE (UNII: C151H8M554)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
ARGININE COCOATE (UNII: 951Q8XZ62H)	
SACCHARINA LATISSIMA (UNII: 68CMP2MB55)	
CAFFEINE (UNII: 3G6A5W338E)	
COLA ACUMINATA SEED (UNII: 1F8VIW1479)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
STEARAMIDOPROPYL DIMETHYLAMINE (UNII: K7VEI00UFR)	
N-ACETYLGLUCOSAMINE (UNII: V956696549)	
CAPRYLOYL GLYCINE (UNII: 8TY5YO42NJ)	
10-HYDROXYDECANOIC ACID (UNII: NP03XO416B)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CHLOROXYLENOL (UNII: 0F32U78V2Q)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49527-119-01	125 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/15/2024	
2	NDC:49527-119-02	1 in 1 CARTON	04/15/2024	
2		50 mL in 1 BOTTLE, PUMP; 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	04/15/2024	

Labeler - CLINIQUE LABORATORIES LLC (044475127)

Registrant - Estee Lauder Companies Inc. (790802086)

Establishment

Name	Address	ID/FEI	Business Operations
The Estee Lauder Inc		802599436	manufacture(49527-119)

Establishment

Name	Address	ID/FEI	Business Operations
NORTHTEC KEYSTONE		949264774	label(49527-119) , pack(49527-119)

Establishment

Name	Address	ID/FEI	Business Operations
NORTHTEC LLC		943871157	label(49527-119) , pack(49527-119)

Revised: 4/2024 CLINIQUE LABORATORIES LLC