

AGELOC LUMISPA TREATMENT CLEANSER ACNE- salicylic acid lotion

NSE Products, Inc

ageLOC® LumiSpa™ Treatment Cleanser Acne

Drug Facts

Active Ingredient

Salicylic Acid (0.5%)

Purpose

Acne treatment

Uses

- For the treatment of acne.
- Penetrates pores to eliminate most 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 unless directed by a doctor. If swallowed, get medical help or contact a Poison Control Center right away.
- **Keep out of reach of children.**

Directions

- For face.
- Dampen skin with warm water.
- Apply ample treatment cleanser evenly over face, avoiding eye area.
- For optimal results, use with ageLOC® LumiSpa™ device as directed in ageLOC LumiSpa User's Manual.

Inactive Ingredients

Water (Aqua), Cetearyl Alcohol, Glycerin, PEG-8 Dimethicone, Cocamidopropyl Betaine, Caprylyl/Capryl Glucoside, Capryl/Capramidopropyl Betaine, Carbomer, Narcissus Tazetta Bulb Extract, Epilobium Angustifolium Flower/Leaf/Stem Extract, Schizandra Chinensis Fruit Extract, Xanthan Gum, Carnosine, Oryza Sativa (Rice) Extract, Disodium EDTA, Phenoxyethanol, Hydroxyacetophenone, Sodium Hydroxide, Sodium Chloride.

Questions?

1-888-742-7626

PRINCIPAL DISPLAY PANEL - 100 ml Tube Label

NU SKIN®

ageLOC®

LumiSpa™

TREATMENT CLEANSER

acne

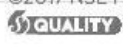
100 ml e (3.4 fl.oz.)



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SAFETY-ALLERGY-DERMATOLOGIST TESTED

SOLD EXCLUSIVELY BY NU SKIN ENTERPRISES
 AUTHORIZED DISTRIBUTORS
 MANUFACTURED IN THE U.S.A. EXCLUSIVELY FOR
 NSE PRODUCTS, INC., PROVO, UT, 84601
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AGELOC LUMISPA TREATMENT CLEANSER ACNE

salicylic acid lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62839-1495
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 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
Glycerin (UNII: PDC6A3C0OX)	
PEG-8 DIMETHICONE (600 CST) (UNII: GIA7T764OD)	
Cocamidopropyl Betaine (UNII: 5OCF3O11KX)	
CAPRYLYL/CAPRYL OLIGOGLUCOSIDE (UNII: E00JL9G9K0)	
Capryl/Capramidopropyl Betaine (UNII: 231H3ZT9NE)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
NARCISSUS TAZETTA BULB (UNII: K17762966S)	
EPILOBIUM ANGUSTIFOLIUM FLOWERING TOP (UNII: 08H094218D)	
SCHISANDRA CHINENSIS FRUIT (UNII: ABS794681C)	
Xanthan Gum (UNII: TTV12P4NEE)	
Carnosine (UNII: 8HO6PVN24W)	
RICE GERM (UNII: 7N2B70SFEZ)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
Phenoxyethanol (UNII: HIE492ZZ3T)	
Sodium Hydroxide (UNII: 55X04QC32I)	
Sodium Chloride (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62839-1495-1	100 mL in 1 TUBE; Type 0: Not a Combination Product	10/01/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	M006	10/01/2017	

Labeler - NSE Products, Inc (803486393)

Revised: 12/2024

NSE Products, Inc