

KIEHL'S SINCE 1851 BLUE HERBAL ACNE CLEANSER TREATMENT WITH FRANKINCENSE AND GINGER ROOT- salicylic acid lotion
L'Oreal USA Products Inc

Drug Facts

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

Keep out of reach of children.

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

Directions

- wet face, then work product into a lather
- massage onto face avoiding the eye area
- rinse well
- use twice daily

Inactive ingredients

water, coco-glucoside, propylene glycol, ammonium polyacryloyldimethyl taurate, glycerin, coco-betaine, sodium cocoyl isethionate, sodium methyl cocoyl taurate, dipropylene glycol, sodium chloride, butylene glycol, phenoxyethanol, hydrogenated coconut acid, capryloyl salicylic acid, coconut acid, t-butyl alcohol, sodium pca, camphor,

menthol, boswellia serrata extract, sodium isethionate, hamamelis virginiana (witch hazel) water, peumus boldus leaf extract, disodium EDTA, alcohol, zingiber officinale (ginger) root extract, sanguisorba officinalis root extract, cinnamomum cassia bark extract, xanthan gum, laminaria saccharina extract, zinc sulfate, pyridoxine HCL, sodium hydroxide, citric acid

Questions or comments?

Call toll free **1-800-946-4453**



SINCE **KIEHL'S** 1851

BLUE HERBAL ACNE CLEANSER TREATMENT

*Salicylic Acid Acne Treatment
with Frankincense & Ginger Root*

This cleanser, formulated with a combination of effective ingredients, helps purify pores, removing traces of dirt and oil. Our high-foaming, medicated cleanser helps clear acne blemishes, blackheads, and whiteheads and helps to prevent new ones from forming. Dermatologist-tested for safety. Inspired by the original Blue Astringent Herbal Lotion formulated in 1964.

5.0 fl. oz. - 150 ml

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Fm1a 685741 10 F.LL Code D193165/1

KIEHL'S SINCE 1851 LLC, NEW YORK, NY 10014
MADE IN U.S.A. • www.kiehls.com
Dist., Kiehl's Canada, Montreal H4T 1K5
106 rue Danton 92691 Levallois Perret Cedex
TSA 75000 93584 ST. OZEN CEDEX FR



2108601

OTHER

BARCODE



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C1 - Internal use

WITH FRANKINCENSE AND GINGER ROOT

salicylic acid lotion

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49967-086
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
WATER (UNII: 059QF0KO0R)			
COCO GLUCOSIDE (UNII: ICS790225B)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
AMMONIUM POLYACRYLOYLDIMETHYL TAURATE (55000 MPA.S) (UNII: F01RIY4371)			
GLYCERIN (UNII: PDC6A3C0OX)			
COCO-BETAINE (UNII: 03DH2IZ3FY)			
SODIUM COCOYL ISETHIONATE (UNII: 518XTE8493)			
SODIUM METHYL COCOYL TAURATE (UNII: JVL98CG53G)			
DIPROPYLENE GLYCOL (UNII: E107L85C40)			
SODIUM CHLORIDE (UNII: 451W47IQ8X)			
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)			
PHENOXYETHANOL (UNII: HIE492ZZ3T)			
HYDROGENATED COCONUT OIL (UNII: JY81OXM1OM)			
CAPRYLOYL SALICYLIC ACID (UNII: 5F7PJF6AA4)			
COCONUT ACID (UNII: 40U37V505D)			
TERT-BUTYL ALCOHOL (UNII: MD83SFE959)			
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)			
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)			
MENTHOL (UNII: L7T10EIP3A)			
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)			
SODIUM ISETHIONATE (UNII: 3R36J71C17)			
HAMAMELIS VIRGINIANA TOP WATER (UNII: NT00Y05A2V)			
PEUMUS BOLDUS LEAF (UNII: Q4EWM09M3O)			
EDETATE DISODIUM (UNII: 7FLD91C86K)			
ALCOHOL (UNII: 3K9958V90M)			
GINGER (UNII: C5529G5JPQ)			
SANGUISORBA OFFICINALIS ROOT (UNII: 4NYV2HT01X)			
CHINESE CINNAMON (UNII: WS4CQ062KM)			
XANTHAN GUM (UNII: TTV12P4NEE)			
SACCHARINA LATISSIMA (UNII: 68CMP2MB55)			
ZINC SULFATE (UNII: 89DS0H96TB)			
PYRIDOXINE HYDROCHLORIDE (UNII: 68Y4CF58BV)			
SODIUM HYDROXIDE (UNII: 55X04QC32I)			

CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-086-01	75 mL in 1 TUBE; Type 0: Not a Combination Product	04/26/2016	
2	NDC:49967-086-02	150 mL in 1 TUBE; Type 0: Not a Combination Product	04/26/2016	
3	NDC:49967-086-03	30 mL in 1 TUBE; Type 0: Not a Combination Product	04/26/2016	
4	NDC:49967-086-04	3 mL in 1 PACKET; Type 0: Not a Combination Product	04/26/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	04/26/2016	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L'Oreal USA, Inc.		185931458	manufacture(49967-086) , pack(49967-086)

Revised: 12/2023

L'Oreal USA Products Inc