

**KIEHLS SINCE 1851 DERMATOLOGIST SOLUTIONS BREAKOUT CONTROL ACNE
TREATMENT FACIAL- salicylic acid lotion
L'Oreal USA Products Inc**

Drug Facts

Active ingredient

Salicylic acid 1.5%

Purpose

Acne treatment

Uses

- for the treatment of acne
- helps prevent 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 swallowed, get medical help or contact a Poison Control Center right away.

Directions

- clean the skin thoroughly before applying this product
- cover the entire affected area with a thin layer one to three times daily
- because excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a doctor
- if bothersome dryness or peeling occurs, reduce application to once a day or every other day

Inactive ingredients

water, aloe barbadensis leaf juice, di-C12-13 alkyl malate, niacinamide, cyclohexasiloxane, propylene glycol, aluminum starch octenylsuccinate, PEG-100

stearate, glyceryl stearate, cetyl alcohol, acrylates/C10-30 alkyl acrylate crosspolymer, sodium hydroxide, capryloyl salicylic acid, xanthan gum, zinc PCA, zingiber officinale (ginger) root extract, stearyl alcohol, myristyl alcohol, citric acid, potassium sorbate, octadecenedioic acid, sodium benzoate, citral, BHT

Questions or comments?

Call toll free 1-800-946-4453

This facial treatment lotion clears acne blemishes and helps prevent future breakouts, while reducing the appearance of dullness and rough texture. Formulated with Salicylic Acid and Vitamin B3, our acne-clearing facial treatment lotion calms and soothes skin, while evening skin tone appearance for a clear complexion. Over time, skin appears healthy and youthfully refined.

Kiehl's Dermatologist Solutions are highly advanced, targeted treatments developed by Kiehl's Since 1851. Relying on our extensive skincare expertise and botanical knowledge, Kiehl's chemists partner with an international team of leading dermatologists to deliver powerful, yet safe skincare solutions.



**Breakout Control
Acne Treatment
Facial Lotion**

Salicylic Acid Acne Treatment

Clinically Demonstrated* to
Diminish Acne Breakouts
while Reducing Rough Texture
and Dull Skin Tone

*with 1.5% Salicylic Acid
and Vitamin B3*

2.0 fl. oz. - 60 ml

*Tested in a dermatologist-controlled clinical study

**FSC LOGO
FPO**



KIEHL'S SINCE 1851 LLC
NEW YORK, NY 10014
MADE IN U.S.A.
Dist.: Kiehl's Canada, Montreal B4T 1K5
106 rue Denton 92691 Levallois Perreé Québec
TSA 10007 F 92667 ASNIERES CEDEX
www.kiehls.com
2023119



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Fmla 685689 38 F.I.L. Code D183906/1

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KIEHL'S SINCE 1851 DERMATOLOGIST SOLUTIONS BREAKOUT CONTROL ACNE TREATMENT FACIAL

salicylic acid lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49967-875
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)	
NIACINAMIDE (UNII: 25X51I8RD4)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ006294)	
PEG-100 STEARATE (UNII: YD01N1999R)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CAPRYLOYL SALICYLIC ACID (UNII: 5F7PJF6AA4)	
XANTHAN GUM (UNII: TTV12P4NEE)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
MYRISTYL ALCOHOL (UNII: V42034O9PU)	
OCTADECENEDIOIC ACID (UNII: 565ZMT5QRG)	
CITRAL (UNII: T7EU009VPP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-875-01	1 in 1 CARTON	11/01/2016	11/01/2016
1		60 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:49967-875-02	5 mL in 1 PACKET; Type 0: Not a Combination Product	11/01/2016	
3	NDC:49967-875-03	1.5 mL in 1 PACKET; Type 0: Not a Combination Product	11/01/2016	
4	NDC:49967-875-04	5 mL in 1 TUBE; Type 0: Not a Combination Product	11/01/2016	
5	NDC:49967-875-05	1 in 1 CARTON	11/01/2016	
5		60 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	11/01/2016	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L'Oreal USA, Inc.		185931458	manufacture(49967-875) , pack(49967-875)