KIEHLS SINCE 1851 DERMATOLOGIST SOLUTIONS BREAKOUT CONTROL TARGETED ACNE SPOT TREATMENT- sulfur lotion
L'Oreal USA Products Inc

Drug Facts

Active ingredient

Sulfur 10%

Purpose

Acne treatment

Use

for the treatment of acne

Warnings

For external use only

Do not use on

- broken skin
- large areas of the skin

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.
- apply only to areas with acne

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 ingredient

water, niacinamide, zea mays (corn) starch, glycolic acid, stearyl alcohol, glycerin, octyldodecyl myristate, cetearyl alcohol, glyceryl stearate, magnesium aluminum silicate, phenoxyethanol, polyacrylamide, ceteareth-20, sodium hydroxide, PEG-100 stearate, phenylethyl resorcinol, tocopheryl acetate, C13-14 isoparaffin, disodium EDTA, laureth-7, allantoin, aloe barbadensis leaf juice, hydrogenated lecithin, ethylhexylglycerin, glycyrrhiza glabra (licorice) root extract

Questions or comments?

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

Drug Facts (continued)

ceteareth-20, sodium hydroxide, PEG-100 stearate, phenylethyl resorcinol, tocopheryl acetate, C13-14 isoparaffin, disodium EDTA, laureth-7, allantoin, aloe barbadensis leaf juice, hydrogenated lecithin, ethylhexylglycerin, glycyrrhiza glabra (licorice) root extract

Questions or comments? Call toll free **1-800-948-4453**

Fmla 942010 1 F.I.L. Code D179591/1

Drug Facts (continued)

- 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, niacinamide, zea mays (corn) starch, glycolic acid, stearyl alcohol, glycerin, octyldodecyl myristate, cetearyl alcohol, glyceryl stearate, magnesium aluminum silicate, phenoxyethanol, polyacrylamide,

SINCE 1851

KIEHL'S

DERMATOLOGIST SOLUTIONS™

Breakout Control Targeted Acne Spot Treatment

Sulfur Acne Treatment

Clinically Demonstrated* to Rapidly Reduce the Appearance of Acne Breakouts

with 10% Sulfur and Vitamin B3

0.68 fl. oz. - 20 ml

*Tested in a dermatologist-controlled clinical study

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.

This targeted acne spot treatment helps to rapidly reduce the appearance of breakouts. Formulated with Sulfur and Vitamin B3, our blemish-clearing spot treatment blends evenly into skin and works to help reduce the appearance of acne.

SUNBURN ALERT: This product contains an alpha hydroxy acid (AHA) that may increase your skin's sensitivity to the sun and particularly the possibility of sunburn. Use a sunscreen, wear protective clothing, and limit sun exposure while using this product and for a week afterwards.

KIEHL'S SINCE 1851 LLC
NEW YORK, NY 10014
MADE IN U.S.A.
Dist., Kiehl's Canada,
Montreal H4T 1K5
106 rue Danton 92691
Levallois Perret Cedex
TSA 10007 F 92667
ASNIERES CEDEX
www.kiehls.com

PAP 12M

2001513

BAR CODE

Drug Facts (continued)

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.
- apply only to areas with acne

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Drug Facts

Active ingredient — **Purpose**
Sulfur 10% — Acne treatment

Uses
- for the treatment of acne

Warnings
For external use only

Do not use on
- broken skin
- large areas of the skin

Directions

- clean the skin thoroughly before applying this product
- cover the entire affected area with a thin layer one to three times daily

FSC LOGO (PMS 476)

Drug Facts (continued)

ceteareth-20, sodium hydroxide, PEG-100 stearate, phenylethyl resorcinol, tocopheryl acetate, C13-14 isoparaffin, disodium EDTA, laureth-7, allantoin, aloe barbadensis leaf juice, hydrogenated lecithin, ethylhexylglycerin, glycyrrhiza glabra (licorice) root extract

Questions or comments? Call toll free **1-800-948-4453**

Fmla 942010 1 F.I.L. Code D179591/1

Drug Facts (continued)

- 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, niacinamide, zea mays (corn) starch, glycolic acid, stearyl alcohol, glycerin, octyldodecyl myristate, cetearyl alcohol, glyceryl stearate, magnesium aluminum silicate, phenoxyethanol, polyacrylamide,

SINCE 1851

KIEHL'S

DERMATOLOGIST SOLUTIONS™

Breakout Control Targeted Acne Spot Treatment

Sulfur Acne Treatment

Clinically Demonstrated* to Rapidly Reduce the Appearance of Acne Breakouts

with 10% Sulfur and Vitamin B3

0.68 fl. oz. - 20 ml

*Tested in a dermatologist-controlled clinical study

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.

This targeted acne spot treatment helps to rapidly reduce the appearance of breakouts. Formulated with Sulfur and Vitamin B3, our blemish-clearing spot treatment blends evenly into skin and works to help reduce the appearance of acne.

SUNBURN ALERT: This product contains an alpha hydroxy acid (AHA) that may increase your skin's sensitivity to the sun and particularly the possibility of sunburn. Use a sunscreen, wear protective clothing, and limit sun exposure while using this product and for a week afterwards.

KIEHL'S SINCE 1851 LLC
NEW YORK, NY 10014
MADE IN U.S.A.
Dist., Kiehl's Canada,
Montreal H4T 1K5
106 rue Danton 92691
Levallois Perret Cedex
TSA 10007 F 92667
ASNIERES CEDEX
www.kiehls.com

PAP 12M

2001513

BAR CODE

Drug Facts (continued)

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.
- apply only to areas with acne

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Drug Facts

Active ingredient — **Purpose**
Sulfur 10% — Acne treatment

Uses
- for the treatment of acne

Warnings
For external use only

Do not use on
- broken skin
- large areas of the skin

Directions

- clean the skin thoroughly before applying this product
- cover the entire affected area with a thin layer one to three times daily

FSC LOGO (PMS 476)

KIEHLS SINCE 1851 DERMATOLOGIST SOLUTIONS BREAKOUT CONTROL TARGETED ACNE SPOT TREATMENT

sulfur lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49967-895
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)	SULFUR	100 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
NIACINAMIDE (UNII: 25X51I8RD4)	
GLYCOLIC ACID (UNII: 0WT12SX38S)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
PEG-100 STEARATE (UNII: YD01N1999R)	
PHENYLETHYL RESORCINOL (UNII: G37UFG162O)	
ALLANTOIN (UNII: 344S277G0Z)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:49967-895-01	1 in 1 CARTON	11/01/2016	
1		20 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:49967-895-02	3 mL in 1 PACKET; Type 0: Not a Combination Product	11/01/2016	
3	NDC:49967-895-03	1.5 mL in 1 PACKET; Type 0: Not a Combination Product	11/01/2016	
4	NDC:49967-895-04	3 mL in 1 TUBE; Type 0: Not a Combination Product	11/01/2016	

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	analysis(49967-895)

Establishment

Name	Address	ID/FEI	Business Operations
Englewood Lab, Inc.		172198223	manufacture(49967-895) , pack(49967-895)

Revised: 12/2023 L'Oreal USA Products Inc