IT COSMETICS BYE BYE BREAKOUT CONCENTRATED DERMA SERUM ACNE TREATMENT- salicylic acid lotion L'Oreal USA Products Inc

Drug Facts

Active ingredient

Salicylic acid 2%

Purpose

Acne treatment

Use

- for the treatment of acne
- helps prevent new acne blemishes from forming

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 ingrease to two or three times dialy if neded or as directed by a doctor
- if bothersome dryness or peeling occurs, reduce application to once a day or every other dayl

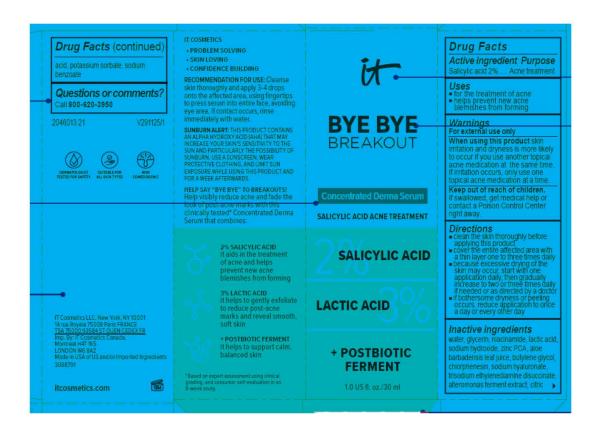
Inactive ingredients

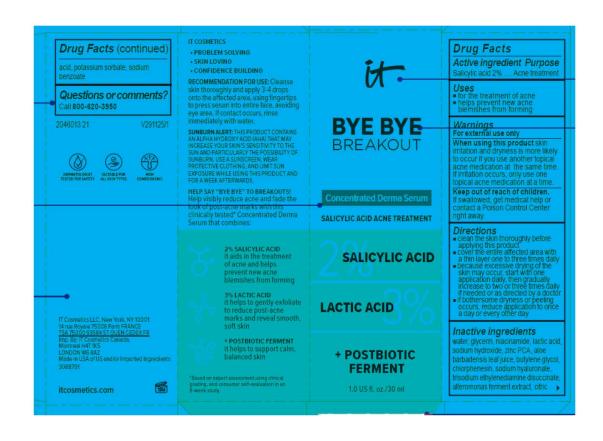
water, glycerin, niacinamide, lactic acid, sodium hydroxide, zinc PCA, aloe barbadensis

leaf juice, butylene glycol, chlorphenesin, sodium hyaluronate, trisodium ethylenediamine disuccinate, alteromonas ferment extract, citric acid, potassium sorbate, sodium benzoate

Questions or comments?

Call **800-620-3950**





ACNE TREATMENT

salicylic acid lotion

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Prod	uct	Intori	mation

HUMAN OTC DRUG **Item Code (Source)** NDC:49967-026 **Product Type**

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ) SALICYLIC ACID 20 mg in 1 mL

Inactive Ingredients

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Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
NIACINAMIDE (UNII: 25X5118RD4)	
LACTIC ACID (UNII: 33X04XA5AT)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ZINC PIDOLATE (UNII: C32PQ86DH4)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CHLORPHENESIN (UNII: 1670DAL4SZ)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
TRISODIUM ETHYLENEDIAMINE DISUCCINATE (UNII: YA22H34H9Q)	
ALTEROMONAS MACLEODII POLYSACCHARIDES (UNII: AP6XG2GR8Z)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

	Packaging				
4	t Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:49967- 026-01	1 in 1 CARTON	04/01/2022		
1		30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product			
2	NDC:49967- 026-02	4 mL in 1 PACKET; Type 0: Not a Combination Product	04/01/2022	04/01/2022	
3	NDC:49967- 026-03	4 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2022		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M006	04/01/2022		

Labeler - L'Oreal USA Products Inc (002136794)

EstablishmentNameAddressID/FEIBusiness OperationsL'Oreal USA, Inc.185931458manufacture(49967-026) , pack(49967-026)

Establishment				
Name	Address	ID/FEI	Business Operations	
Dimensional Merchandising Inc.		076693183	pack(49967-026)	

Revised: 12/2023 L'Oreal USA Products Inc