ACNE CLARIFYING CLEANSER- salicylic acid liquid e.l.f. Cosmetics, Inc.

Acne Clarifying Cleanser

Drug Facts

Active ingredients

Salicylic Acid 1%

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

Cleanse the skin thoroughly before applying medication. 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. Sensitivity Test for a New User. Apply product sparingly to one or two small affected areas during the first 3 days. If no discomfort occurs, follow the directions stated eleswhere on this label.

Other Information

• Protect the product in this container from excessive heat and direct sun.

Inactive ingredients

Water (Aqua), Sodium C14-16 Olefin Sulfonate, Cocamidopropyl Betaine, Glycerin, Zinc PCA, Sodium Benzoate, Hydroxypropyl Methylcellulose, Sodium Hydroxide, Sodium Chloride, Citric Acid, Disodium EDTA, Niacinamide, Tranexamic Acid May Contain: Blue 1 (CI 42090), Yellow 5 (CI 19140)

Questions or comments

1-888-315-9814

Package Labeling: 76354-415-01





Package labeling: 76354-415-02

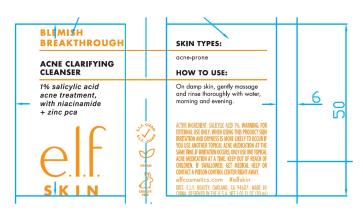












Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:76354-415

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII: O414PZ4LPZ)	SALICYLIC ACID	10 mg in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)		
COCAMIDOPROPYL BETAINE (UNII: 50CF3011KX)		
GLYCERIN (UNII: PDC6A3C0OX)		
ZINC PIDOLATE (UNII: C32PQ86DH4)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)		
SODIUM HYDROXIDE (UNII: 55X04QC32I)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)		
NIACINAMIDE (UNII: 25X5118RD4)		
TRANEXAMIC ACID (UNII: 6T84R30KC1)		

F	ackaging			
#	tem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76354-415- 01	1 in 1 CARTON	02/25/2022	
1		115 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:76354-415- 02	1 in 1 CARTON	02/25/2022	
2		30 mL in 1 BOTTLE; 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	02/25/2022	

Labeler - e.l.f. Cosmetics, Inc. (093902816)

Revised: 4/2024 e.l.f. Cosmetics, Inc.