NATURIUM SALICYLIC ACID BODY 2%- salicylic acid aerosol, spray
e.l.f. Cosmetics, Inc

Naturium Salicylic Acid Body Spray 2%

Drug Facts

Active Ingredient

Salicylic Acid 2.0%

Purpose

Acne Treatment

Uses

- For the treatment of acne.
- Clears acne blemishes and allows skin to heal.
- Helps prevent new acne blemishes from forming.

Warnings

For external use only.

Contents under pressure. Do not puncture or incinerate. Do not store at temperatures above 120F.

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.

Directions

Shake Well

- 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, NIACINAMIDE, ACETYL GLUCOSAMINE, HYDROLYZED OAT PROTEIN, GALACTOMYCES FERMENT FILTRATE, HYDROLYZED CICER SEED EXTRACT, MEDICAGO SATIVA (ALFALFA) EXTRACT, PUERARIA LOBATA ROOT EXTRACT, SILYBUM MARIANUM SEED EXTRACT, AMYLOPECTIN, DEXTRIN, 3-O-ETHYL ASCORBIC ACID, ALPHA-ARBUTIN, POLYDEXTROSE, HYDROXYACETOPHENONE, SODIUM GLUCONATE, POTASSIUM SORBATE, SODIUM BENZOATE, BUTYLENE GLYCOL

Questions?

Visit www.naturium.com

Product Packaging

NATURIUM SALICYLIC ACID BODY 2%

salicylic acid aerosol, spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76354-124
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

Ingredient Name	Strength
PUERARIA MONTANA VAR. LOBATA ROOT (UNII: PET93F4I3C)	
AMYLOPECTIN, UNSPECIFIED SOURCE (UNII: 4XO4QFV777)	
MEDICAGO SATIVA WHOLE (UNII: DJO934BRBD)	
SILYBUM MARIANUM SEED (UNII: U946SH95EE)	
3-O-ETHYL ASCORBIC ACID (UNII: 6MW60CB71P)	
ICODEXTRIN (UNII: 2NX48Z0A9G)	
SODIUM GLUCONATE (UNII: R6Q3791S76)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
N-ACETYLGLUCOSAMINE (UNII: V956696549)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
NIACINAMIDE (UNII: 25X51I8RD4)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
WATER (UNII: 059QF0KO0R)	
ALPHA-ARBUTIN (UNII: 72VUP07IT5)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76354-124-01	120 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	12/01/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	12/01/2022	

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

Revised: 11/2023 e.l.f. Cosmetics, Inc