KEYS SOULCARE DAILY MOISTURIZER BROAD SPECTRUM SPF30 SUNSCREEN-
avobenzone, homosalate, octisalate, octocrylene cream
e.l.f. Cosmetics, Inc

KEYS SOULCARE DAILY MOISTURIZER BROAD SPECTRUM SPF30 SUNSCREEN

Drug Facts

Active ingredients

Avobenzone 3.0%

Homosalate 7.3%

Octisalate 4%

Octocrylene 8.5%

Purpose

Sunscreen

Use

- Helps prevent sunburn
- If used as directed with other sun protection measures (see Directions) decreases the risk of skin cancer and early skin aging caused by the sun.

Warning

For external use only.

Do not use

on damaged or broken skin.

When using this product:

keep out of eyes, Rinse with water to remove.

Stop use and ask a doctorif:

rash occurs.

Keep out of reach of children.

If swallowed, get medical help or conact a Poison Control Center right away.

Directions

- Apply libreally 15 minutes before sun exposure, Reapply at least every 2 hours.

- Use water resistant sunscreen if swimming or sweating.
- Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly use a sunscreen with a Braod Spectrum SPF value of 15 or higher and other sun protection measures including: limit time in sun, especially from 10 a.m.-2 p.m., wear long-sleeved shirts, pants, hats and sunglasses. Children under 6 months of age: ask a doctor. **Sun Protection Measures:**

Other Information

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

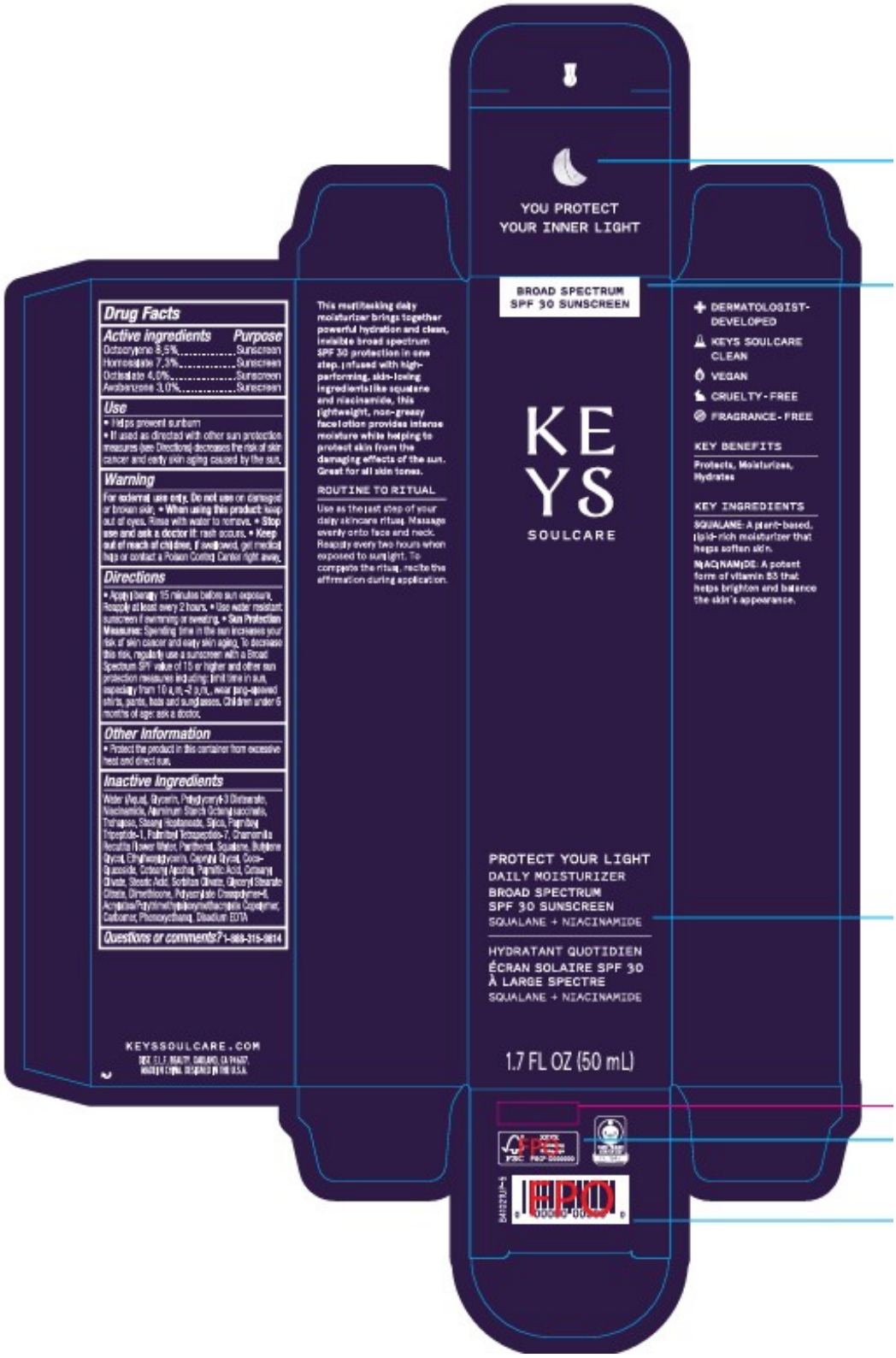
Inactive Ingredients

Water (Aqua), Glycerin, Polyglyceryl-3 Distearate, Niacinamide, Aluminum Starch Octenylsuccinate, Trehalose, Stearyl Heptanoate, Silica, Palmitoyl Tripeptide-1, Palmitoyl Tetrapeptide-7, Chamomilla Recutita Flower Water, Panthenol, Squalane, Butylene Glycol, Ethylhexylglycerin, Caprylyl Glycol, Coco-Glucoside, Cetearyl Alcohol, Palmitic Acid, Cetearyl Olivate, Stearic Acid, Sorbitan Olivate, Glyceryl Stearate Citrate, Dimethicone, Polyacrylate Crosspolymer-6, Acrylates/Polytrimethylsiloxymethacrylate Copolymer, Carbomer, Phenoxyethanol, Disodium EDTA

Questions or comments?

1-888-315-9814

Package Labeling:50 mL



Package Labeling:15 mL

Package Labeling:5 mL

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76354-454
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	73 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	40 mg in 1 mL

OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	85 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86FQE96TZ4)	
POLYACRYLATE CROSSPOLYMER-6 (UNII: Q7UI015FF9)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYGLYCERYL-3 DISTEARATE (UNII: ZI1LK470XV)	
NIACINAMIDE (UNII: 25X51I8RD4)	
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ0O6294)	
TREHALOSE (UNII: B8WCK70T7I)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARYL HEPTANOATE (UNII: 2M4UGL1NCN)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PALMITOYL TRIPEPTIDE-1 (UNII: RV743D216M)	
PALMITOYL TETRAPEPTIDE-7 (UNII: Q41S464P1R)	
CHAMOMILE FLOWER OIL (UNII: 60F80Z61A9)	
PANTHENOL (UNII: WV9CM0O67Z)	
SQUALANE (UNII: GW89575KF9)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
PALMITIC ACID (UNII: 2V16EO95H1)	
CETEARYL OLIVATE (UNII: 58B69Q84JO)	
SORBITAN OLIVATE (UNII: MDL271E3GR)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
GLYCERYL STEARATE CITRATE (UNII: WH8T92A065)	
STEARYL CAPRYLATE (UNII: 06TS6O9194)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
COCO GLUCOSIDE (UNII: ICS790225B)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76354-454-01	1 in 1 CARTON	02/17/2025	
1		50 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:76354-454-02	1 in 1 CARTON	02/17/2025	
2		15 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:76354-454-03	1 in 1 CARTON	02/17/2025	

3		5 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	M020	02/17/2025	

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

Revised: 2/2025 e.l.f. Cosmetics, Inc