E.L.F INVISIBLE SUNSCREEN SPF 35- avobenzone, octisalate, octocrylene, homosalate cream
e.l.f. Cosmetics, Inc

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e.l.f Invisible Sunscreen SPF 35

Drug Facts

Active Ingredients

Avobenzone 3.0%

Octisalate 5.0%

Octocrylene 8.0%

Homosalate 7.0%

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.

Warnings

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 doctor if :

- rash occurs.

Keep out of reach of children.

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

Directions

- Apply liberally 15 minutes before sun exposure.
- After 80 minutes of swimming.

- Immediately after towel drying.
- At least every 2 hours.
- 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 Broad 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

Isododecane, Dimethicone, Silica, Dimethyl Silylate, Caprylyl Methicone, PEG-10 Dimethicone/Vinyl Dimethicone Crosspolymer, Glycerin, PEG-12 Dimethicone/PPG-20 Crosspolymer, Pongamia Pinnata Seed Extract, Limnanthes Alba (Meadowfoam) Seed Oil, Aloe Barbadensis Leaf Juice Powder, Polyester-7, Neopentyl Glycol Diheptanoate, Phenoxyethanol, Ethylhexylglycerin, Isopropyl Palmitate

Question or comments?

1-888-315-9814

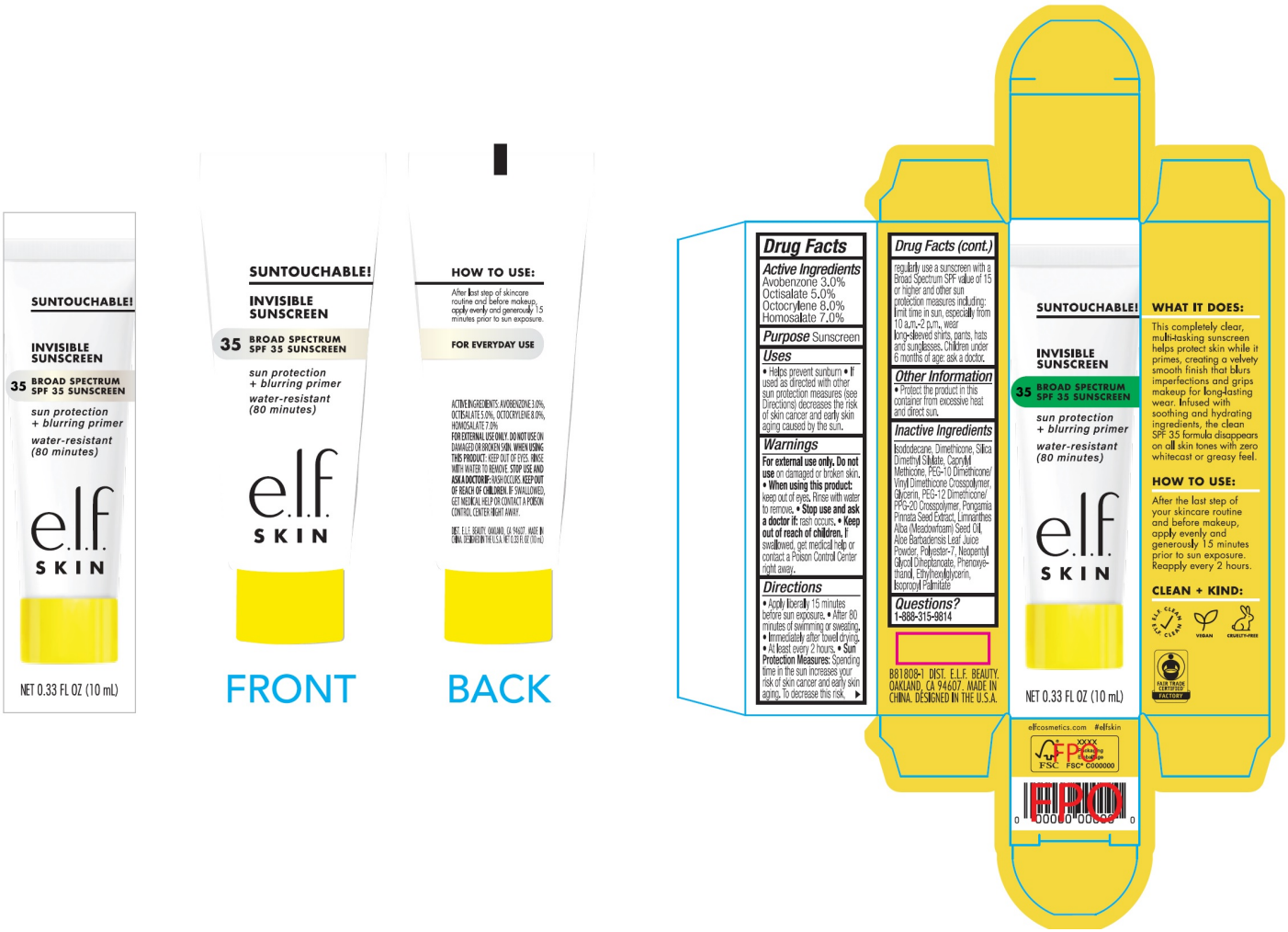
Package Labeling:50ml

REFERENCE

FRONT

BACK

Package Labeling:10ml



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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76354-444
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
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	80 mg in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	70 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ISODODECANE (UNII: A8289P68Y2)	
DIMETHICONE (UNII: 92RU3N3Y1O)	

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)	
GLYCERIN (UNII: PDC6A3C0OX)	
PEG-12 DIMETHICONE/PPG-20 CROSSPOLYMER (UNII: 965K72OQXO)	
PONGAMIA PINNATA SEED (UNII: C2BRV53B1V)	
MEADOWFOAM SEED OIL (UNII: 412ZHA4T4Y)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
POLYESTER-7 (UNII: 0841698D2F)	
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKW3C543X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76354-444-01	1 in 1 CARTON	02/15/2023	
1		50 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:76354-444-02	1 in 1 CARTON	02/15/2023	
2		10 mL in 1 TUBE; 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	02/15/2023	

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

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