

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

e.l.f Invisible Sunscreen SPF 35

Drug Facts

Active Ingredients

Avobenzene 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 generously and evenly 15 minutes before sun exposure.
- Reapply:

- after 80 minutes of swimming or sweating.
- immediately after towel drying.
- at least every 2 hours.
- Sun Protection Measures: 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: Ask a doctor.

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



Drug Facts

Active Ingredients	Purpose
Azobenzene 3.0%, Octisalate 5.0%, Octocrylene 8.0%, Homosalate 7.0%	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 generously and evenly 15 minutes before sun exposure • Reapply • after 80 minutes of swimming or sweating • immediately after towel drying • at least every 2 hours • **Sun Protection Measures:** 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 the sun, especially from 10 a.m. - 2 p.m. • Wear long-sleeved shirts, pants, hats, and sunglasses • Children under 6 months old ask a doctor.

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Inactive Ingredients butadiene, Dimethicone, Silica Dimethyl Silylate, Caprylyl Methacrylate, PEG-10 Dimethicone Vinyl Dimethicone Crosspolymer, Glycerin, PEG-12 Dimethicone/PF-20 Crosspolymer, Pargamita Prunella Seed Extract, Limnathes Alba (Meadowfoam) Seed Oil, Aloe Barbadensis Leaf Juice Powder, Polybutene-1, Neopentyl Glycol Diethoxysilane, Phenylbenzylcarbamate, Isopropyl Palmitate

Questions or comments? 1-888-315-8614



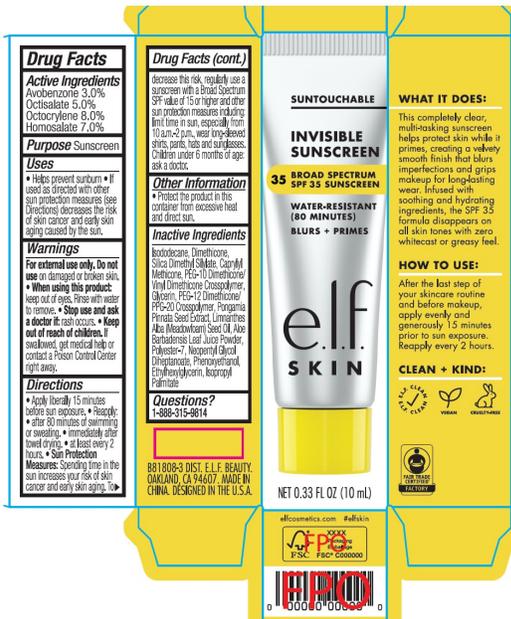
FRONT BACK

REFERENCE

Package Labeling:10ml



REFERENCE



FRONT BACK

REFERENCE

E.L.F INVISIBLE SUNSCREEN SPF 35
avobenzone, octisalate, octocrylene, homosalate cream

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, UNSPECIFIED (UNII: 92RU3N3Y1O)	
SILICA DIMETHYL SILYLATE (UNII: EU2PSP0G0W)	
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	

