LANCOME PARIS TEINT IDOLE ULTRA CARE AND GLOW SKINCARE FOUNDATION BROAD SPECTRUM SPF 27 SUNSCREEN- ensulizole liquid
SICOS ET CIE

Drug Facts

Active ingredient

Ensulizole 3.6%

Purpose

Sunscreen

Uses

- 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

Flammable until dry.

Do not use near fire, flame, or heat.

Do not use

on damaged or broken skin

When using this product

keep out of eyes. Rinse with wate 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

For sunscreen use:

- shake well
- apply liberally 15 minutes before sun exposure
- reapply at least every 2 hours
- use a water resistant sunscreen if swimming or sweating
- **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 of age: Ask a doctor

Other information

protect the product in this container from excessive heat and direct sun

Inactive ingredients

dimethicone, water, isododecane, alcohol denat., trimethylsiloxysilicate, propylene glycol, glycerin, PEG/PPG-18/18 dimethicone, silica, synthetic fluorphlogopite, polysilicone-11, rosa gallica flower extract, moringa oleifera seed extract, polymnia sonchifolia root juice, sorbitol, mandelic acid, PEG-10 dimethicone, alpha-glucan oligosaccharide. sodium chloride, sodium hyaluronate, sodium hydroxide, silica silylate, cellulose acetate butyrate, aluminum hydroxide, ammonium polyacryloyldimethyl taurate, disodium phosphate, disodium stearoyl glutamate, hydroxyethylpiperazine ethane sulfonic acid, caprylyl glycol, citric acid, lactobacillus, maltodextrin, polyphosphorylcholine glycol acrylate, polyvinyl alcohol, butylene glycol, tocopherol, phenoxyethanol, fragrance, linalool, geraniol, limonene, citral, citronellol, benzyl alcohol; may contain: titanium dioxide, iron oxides

Questions or comments?

1-800-LANCOME

(1-800-526-2663)

Monday - Friday

(9 a.m. - 5 p.m. EST)

Drug Facts (continued)

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G993657 V289711/2

SCAN HERE FOR THE BEST MAKEUP TIPS.

Oil-free
Paraben-free
Allergy tested
Non-comedogenic
Tested under dermatological control

12M

CAUTION: FLAMMABLE UNTIL DRY. DO NOT USE NEAR FIRE, FLAME OR HEAT.

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TEINT IDOLE ULTRA

CARE & GLOW

ENSULIZOLE SUNSCREEN BROAD SPECTRUM SPF 27

SKINCARE FOUNDATION
UP TO 24H HEALTHY GLOW
TRANSFER-RESISTANT

COVERAGE
● ● ● ○
105W

30 ml - 1.0 FL. OZ.

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ensulizole liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51150-676
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ENSULIZOLE (UNII: 9YQ9DI1W42) (ENSULIZOLE - UNII:9YQ9DI1W42)	ENSULIZOLE	36 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
DIMETHICONE (UNII: 92RU3N3Y1O)	
WATER (UNII: 059QF0KO0R)	
ISODODECANE (UNII: A8289P68Y2)	
ALCOHOL (UNII: 3K9958V90M)	
TRIMETHYLSILOXYSILICATE (M/Q 0.6-0.8) (UNII: 5041RX63GN)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
PEG/PPG-18/18 DIMETHICONE (UNII: 9H0AO7T794)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MAGNESIUM POTASSIUM ALUMINOSILICATE FLUORIDE (UNII: YK3DC63Y5M)	
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE) (UNII: 9E4CO0W6C5)	
ROSA GALLICA FLOWER (UNII: X8W61WUV70)	
SORBITOL (UNII: 506T60A25R)	
MANDELIC ACID (UNII: NH496X0UJX)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
.ALPHA.-GLUCAN OLIGOSACCHARIDE (UNII: S95658MI3W)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
AMMONIUM POLYACRYLOYLDIMETHYL TAURATE (55000 MPA.S) (UNII: F01RIY4371)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
DISODIUM STEAROYL GLUTAMATE (UNII: 45ASM2L11M)	
HYDROXYETHYLPIPERAZINE ETHANE SULFONIC ACID (UNII: RWW266YE9I)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
TOCOPHEROL (UNII: R0ZB2556P8)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
GERANIOL (UNII: L837108USY)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
CITRAL (UNII: T7EU0O9VPP)	
.BETA.-CITRONELLOL, (R)- (UNII: P01OUT964K)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51150-676-01	1 in 1 CARTON	05/01/2022	05/01/2022
1		30 mL in 1 TUBE; Type 0: Not a Combination Product		

2	NDC:51150-676-02	1 mL in 1 PACKET; Type 0: Not a Combination Product	05/01/2022	12/08/2026
3	NDC:51150-676-03	1.6 mL in 1 PACKAGE; Type 0: Not a Combination Product	05/01/2022	01/19/2026
4	NDC:51150-676-04	1 in 1 CARTON	05/01/2022	05/16/2027
4		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	M020	05/01/2022	05/16/2027

Labeler - SICOS ET CIE (276993581)

Establishment

Name	Address	ID/FEI	Business Operations
SICOS ET CIE		276993581	manufacture(51150-676) , pack(51150-676)

Establishment

Name	Address	ID/FEI	Business Operations
SOCOPLAN		276221405	pack(51150-676)

Revised: 6/2024 SICOS ET CIE