

**PRADA REVEAL SKIN OPTIMIZING FOUNDATION BROAD SPECTRUM SPF 17
SUNSCREEN- ensulizole liquid
SICOS ET CIE**

Drug Facts

Active ingredient

Ensulizole 2.7%

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 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

For sunscreen use:

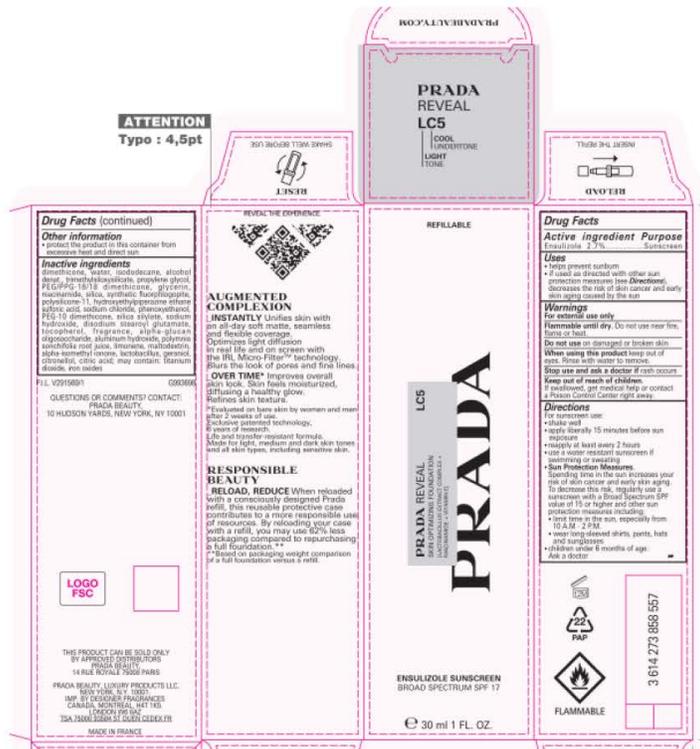
- shake well before use
- apply liberally 15 minutes before sun exposure
- reapply at least every 2 hours
- use a water resistant sunscreen if swimming or sweating
- **Sun Protection Measures.** Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly 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, PEG/PPG-18/18 dimethicone, glycerin, niacinamide, silica, synthetic fluorophlogopite, polysilicone-11, hydroxyethylpiperazine ethane sulfonic acid, sodium chloride, phenoxyethanol, PEG-10 dimethicone, silica silylate, sodium hydroxide, disodium stearyl glutamate, tocopherol, fragrance, alpha-glucan oligosaccharide, aluminum hydroxide, polymnia sonchifolia root juice, limonene, maltodextrin, alpha-isomethyl ionone, lactobacillus, geraniol, citronellol, citric acid; may contain: titanium dioxide, iron oxides



C1 - Internal use

PRADA REVEAL SKIN OPTIMIZING FOUNDATION BROAD SPECTRUM

SPF 17 SUNSCREEN

ensulizole liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51150-380
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ENSULIZOLE (UNII: 9YQ9DI1W42) (ENSULIZOLE - UNII:9YQ9DI1W42)	ENSULIZOLE	27 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)	
PEG/PPG-18/18 DIMETHICONE (UNII: 9H0AO7T794)	
GLYCERIN (UNII: PDC6A3C0OX)	
NIACINAMIDE (UNII: 25X51I8RD4)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE) (UNII: 9E4CO0W6C5)	
HYDROXYETHYLPIPERAZINE ETHANE SULFONIC ACID (UNII: RWW266YE9I)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
DISODIUM STEAROYL GLUTAMATE (UNII: 45ASM2L11M)	
TOCOPHEROL (UNII: R0ZB2556P8)	
.ALPHA.-GLUCAN OLIGOSACCHARIDE (UNII: S95658MI3W)	
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
ISOMETHYL-.ALPHA.-IONONE (UNII: 9XP4LC555B)	
LACTOBACILLUS ACIDOPHILUS BCMC 12130 (UNII: MVC2243UZ X)	
GERANIOL (UNII: L837108USY)	
CITRONELLOL ACETATE, (S)- (UNII: 78RZL4H51H)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

Packaging

Marketing Start Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51150-380-01	1 in 1 CARTON	06/01/2023	05/14/2027
1		30 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:51150-380-02	3 mL in 1 PACKAGE; Type 0: Not a Combination Product	06/01/2023	02/28/2027
3	NDC:51150-380-03	1 mL in 1 PACKET; Type 0: Not a Combination Product	06/01/2023	
4	NDC:51150-380-04	3 mL in 1 PACKET; Type 0: Not a Combination Product	01/02/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	06/01/2023	

Labeler - SICOS ET CIE (276993581)

Establishment

Name	Address	ID/FEI	Business Operations
SICOS ET CIE		276993581	manufacture(51150-380)

Establishment

Name	Address	ID/FEI	Business Operations
BPS 60		272259304	pack(51150-380)

Establishment

Name	Address	ID/FEI	Business Operations
Socoplan		276221405	pack(51150-380)

Revised: 7/2025

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