FULL SPECTRUM MATTE AMBITION SKIN PRIMER SPF 20- ensulizole liquid
Noxell Corporation

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Covergirl Full Spectrum Matte Ambition Primer SPF 20

Active ingredient

Ensulizole 3%

Purpose

Sunscreen

Uses

helps prevent sunburn

Warnings

Skin Cancer/Skin Aging Alert: Spending time in the sun increases your risk of skin cancer and early skin aging. This product has been shown only to help prevent sunburn, not skin cancer or early skin aging.

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 product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

- apply liberally 15 minutes before sun exposure
- reapply at least every 2 hours
- use water resistant sunscreen if swimming or sweating
- children under 6 months: ask a doctor

Other information

Protect this product from excessive heat and direct sun

Inactive ingredients

Aqua/Water/Eau, Cyclopentasiloxane, Talc, Propylene Glycol, Dimethicone, Aluminum Starch Octenylsuccinate, PEG/PPG-18/18 Dimethicone, Sodium Chloride, PVP, Phenoxyethanol, Sodium Hydroxide, Acrylonitrile/Methyl Methacrylate/Vinylidene Chloride Copolymer, Synthetic Beeswax, Trihydroxystearin, Methicone, 1,2-Hexanediol, Caprylyl Glycol, Silica, Sodium Benzoate, Synthetic Wax, Polyglyceryl-4 Isostearate, Cetyl PEG/PPG-10/1 Dimethicone, Hexyl Laurate, Isopropyl Titanium Triisostearate, Ethylene Brassylate, Polyethylene.

Questions?

1-800-426-8374

full spectrum

COVERGIRL

MATTE AMBITION

MATTIFYING

SKIN PRIMER +

ENSULIZOLE

SUNSCREEN

SPF 20

30 mL (1.0 FL OZ)

FULL SPECTRUM MATTE AMBITION SKIN PRIMER SPF 20

ensulizole liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:22700-201
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ENSULIZOLE (UNII: 9YQ9DI1W42) (ENSULIZOLE - UNII:9YQ9DI1W42)	ENSULIZOLE	3 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALUMINUM STARCH OCTENYLSUCCINATE (UNII: I9PJ0O6294)	
PEG/PPG-18/18 DIMETHICONE (UNII: 9H0AO7T794)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
HEXYL LAURATE (UNII: 4CG9F9W01Q)	
ETHYLENE BRASSYLATE (UNII: 9A87HC7ROD)	
DIMETHICONE (UNII: 92RU3N3Y1O)	

SODIUM CHLORIDE (UNII: 451W47IQ8X)	
POLYGLYCERYL-4 ISOSTEARATE (UNII: 820DPX33S7)	
TALC (UNII: 7SEV7J4R1U)	
WATER (UNII: 059QF0KO0R)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 2) (UNII: V2W71V8T0X)	
ISOPROPYL TITANIUM TRIISOSTEARATE (UNII: 949E3KBJ1I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:22700-201-30	30 mL in 1 TUBE; Type 0: Not a Combination Product	02/28/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	02/28/2019	

Labeler - Noxell Corporation (003082997)

Establishment

Name	Address	ID/FEI	Business Operations
Noxell Corporation		003082997	manufacture(22700-201)

Revised: 1/2023 Noxell Corporation