GARNIER OMBRELLE ULTRA LIGHT ADVANCED 60 FACE SUNSCREEN- drometrizole trisiloxane, octocrylene, bemotrizinol, octisalate, avobenzone, homosalate and ensulizole lotion
L'OREAL USA PRODUCTS INC

Drug Facts

Active ingredients

DROMETRIZOLE TRISILOXANE 7%
OCTOCRYLENE 5%
BEMOTRIZINOL 5%
OCTISALATE 5%
AVOBENZONE 3%
HOMOSALATE 2%
ENSULIZOLE 0.50%

Warnings

For external use only. Do not use on broken skin. When using this product avoid contact witheyes. If contact occurs, rinse thoroughlly with water. 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

Adults and adolescents 12 to 17 years.
Apply generously and evenly 15 minutes before sun exposure. Reapply at least every 2 hours. Reapply after 80 minutes of swimming or sweating. Reapply immediately after towel drying. 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 Broad Spectrum SPF 15 or higher sunscreen. Limit time in the sun, especially from 10 a.m. – 2 p.m. Wear long-sleeve shirts, pants, hats and sunglasses.

Inactive ingredients

wwater, alcohol denat., silica, isopropyl myristate, glycerin, diisopropyl sebacate, propanediol, c12-22 alkyl acrylate/hydroxyethylacrylate copolymer, perlite, tocopherol, triethanolamine, caprylyl glycol, hydroxyethylcellulose, acrylates/c10-30 alkyl acrylate crosspolymer, pentylene glycol, butylene glycol, citric acid

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SUNSCREEN

drometrizole trisiloxane, octocrylene, bemotrizinol, octisalate, avobenzone, homosalate and ensulizole lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49967-046
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
DROMETRIZOLE TRISILOXANE (UNII: HC22845I1X) (DROMETRIZOLE TRISILOXANE - UNII:HC22845I1X)	DROMETRIZOLE TRISILOXANE	70 mg in 1 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	20 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	50 mg in 1 mL
BEMOTRIZINOL (UNII: PWZ1720CBH) (BEMOTRIZINOL - UNII:PWZ1720CBH)	BEMOTRIZINOL	50 mg in 1 mL
ENSULIZOLE (UNII: 9YQ9DI1W42) (ENSULIZOLE - UNII:9YQ9DI1W42)	ENSULIZOLE	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIISOPROPYL SEBACATE (UNII: J8T3X564IH)	
PROPANEDIOL (UNII: 5965N8W85T)	
PERLITE (UNII: 0SG101ZGK9)	
TOCOPHEROL (UNII: R0ZB2556P8)	
TROLAMINE (UNII: 9O3K93S3TK)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49967-046-	1 in 1 CARTON	01/01/2023	

1	01	1 in 1 CARTON	01/01/2023	
1		50 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
export only		01/01/2023	

Labeler - L'OREAL USA PRODUCTS INC (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
Dimensional Merchandising Inc.		076693183	manufacture(49967-046) , pack(49967-046)

Revised: 1/2026 L'OREAL USA PRODUCTS INC