MAYBELLINE NEW YORK FIT ME MATTE PLUS PORELESS MATTIFYING PORELESS PRIMER BROAD SPECTRUM SPF 20 SUNSCREEN- avobenzone, homosalate, octisalate, and octocrylene liquid
L'Oreal USA Products Inc

Drug Facts

Active ingredients

Avobenzone 3%

Homosalate 10%

Octisalate 5%

Octocrylene 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

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

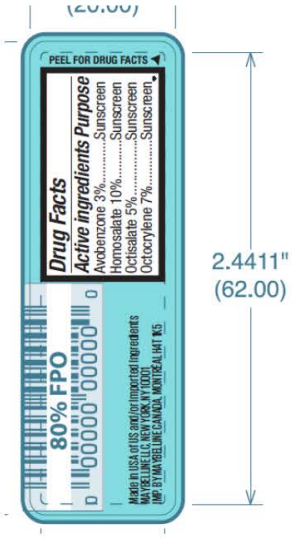
For sunscreen use:
● apply liberally 15 minutes before sun exposure
● reapply at least every 2 hours
● use a water resistant sunscreen if swimming or sweating
● **Sun Protection Measures**. Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly 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

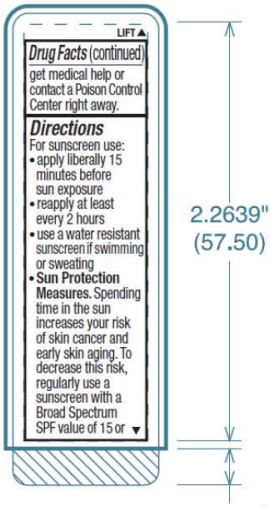
Other information

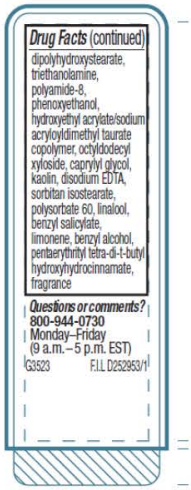
protect the product in this container from excessive heat and direct sun

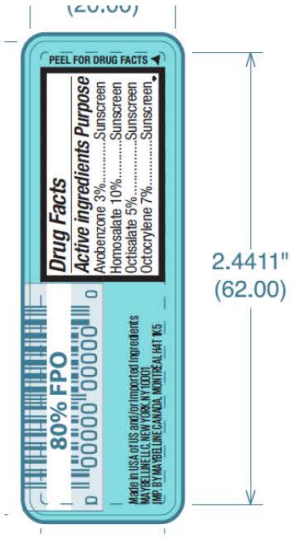
Inactive ingredients

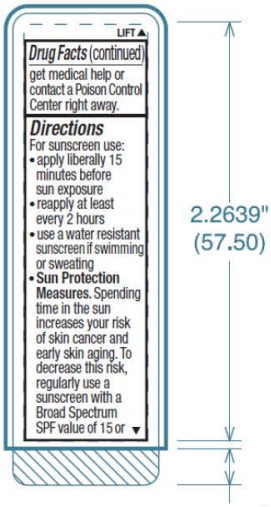
water, isododecane, isononyl isononanoate, silica, copernicia cerifera (carnauba) wax, silica silylate, glycerin, octyldodecanol, PEG-30 dipolyhydroxystearate, triethanolamine, polyamide-8, phenoxyethanol, hydroxyethyl acrylate/sodium acryloyldimethyl taurate, copolymer, octyldodecyl xyloside, caprylyl glycol, kaolin, disodium EDTA, sorbitan isostearate, polysorbate 60, linalool, benzyl salicylate, limonene, benzyl alcohol, pentaerythrityl tetra-di-t-butyl hydroxyhydrocinnamate, fragrance

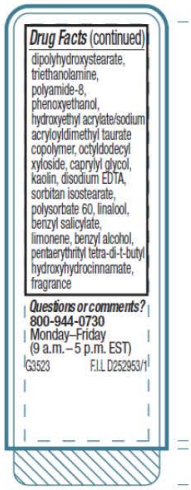


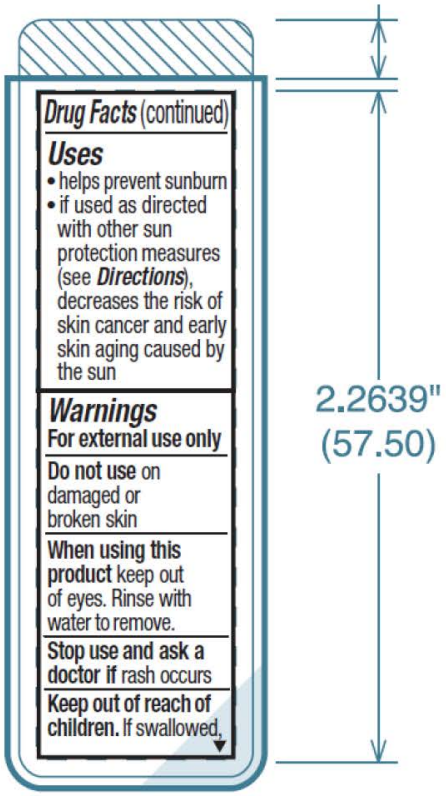


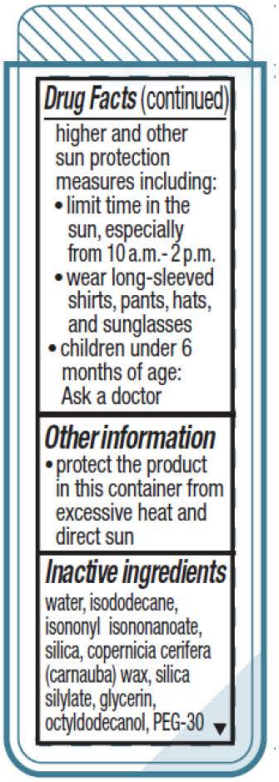


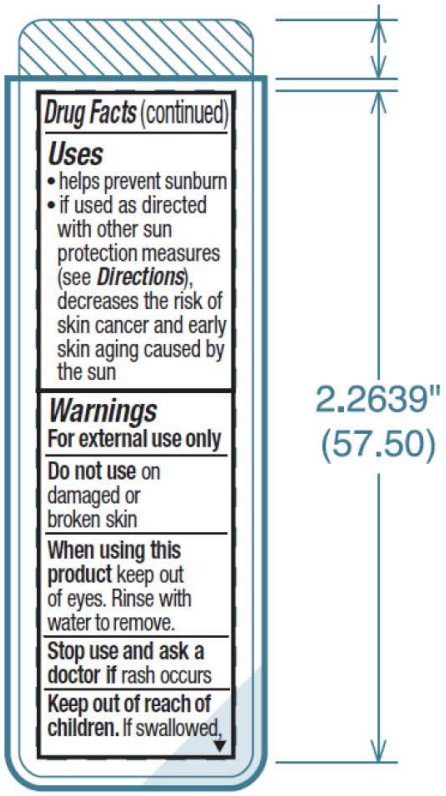


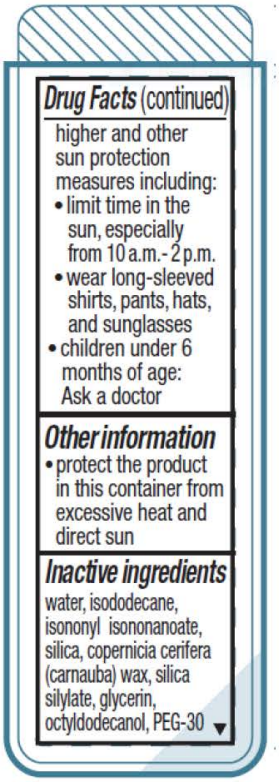












MATTIFYING PORELESS PRIMER BROAD SPECTRUM SPF 20 SUNSCREEN

avobenzone, homosalate, octisalate, and octocrylene liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49967-443
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
Homosalate (UNII: V06SV4M95S) (Homosalate - UNII:V06SV4M95S)	Homosalate	100 mg in 1 mL
Octisalate (UNII: 4X49Y0596W) (Octisalate - UNII:4X49Y0596W)	Octisalate	50 mg in 1 mL
Octocrylene (UNII: 5A68WGF6WM) (Octocrylene - UNII:5A68WGF6WM)	Octocrylene	70 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ISODODECANE (UNII: A8289P68Y2)	
ISONONYL ISONONANOATE (UNII: S4V5BS6GCX)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CARNAUBA WAX (UNII: R12CBM0EIZ)	
GLYCERIN (UNII: PDC6A3C0OX)	
OCTYLDODECANOL (UNII: 461N1O614Y)	
PEG-30 DIPOLYHYDROXYSTEARATE (UNII: 9713Q0S7FO)	
TROLAMINE (UNII: 9O3K93S3TK)	
POLYAMIDE-8 (4500 MW) (UNII: 77723GV81A)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (100000 MPA.S AT 1.5%) (UNII: 86FQE96TZ4)	
OCTYLDODECYL XYLOSIDE (UNII: 8Z6VNR46QM)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
KAOLIN (UNII: 24H4NWX5CO)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
BENZYL SALICYLATE (UNII: WAO5MNK9TU)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
PENTAERYTHRITOL TETRAKIS(3-(3,5-DI-TERT-BUTYL-4-HYDROXYPHENYL)PROPIONATE) (UNII: 255PIF62MS)	

Packaging

#	Item Code	Package Description	Marketing Start	Marketing End

#	Item Code	Package Description	Date	Date
1	NDC:49967-443-01	30 mL in 1 TUBE; Type 0: Not a Combination Product	07/01/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	07/01/2022	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment

Name	Address	ID/FEI	Business Operations
L'Oreal USA Products Inc		624244349	manufacture(49967-443) , pack(49967-443)

Revised: 12/2023 L'Oreal USA Products Inc