MAYBELLINE NEW YORK INSTANT AGE REWIND ERASER TREATMENT MAKEUP BROAD SPECTRUM SPF 18 SUNSCREEN- octinoxate lotion L'Oreal USA Products Inc

Drug Facts

Active ingredient

Octinoxate 7.5%

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

Questions or comments?

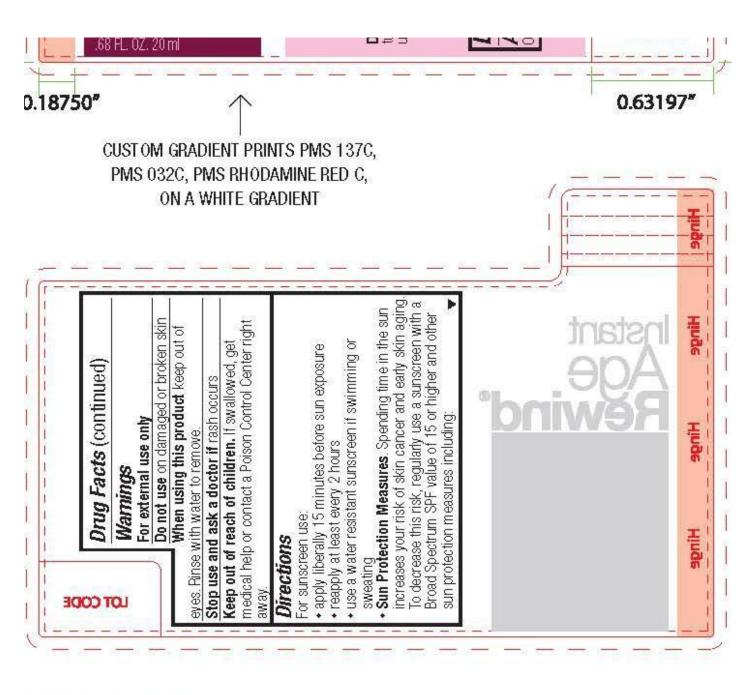
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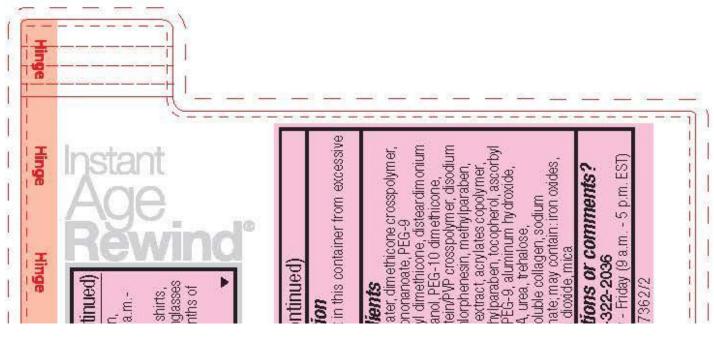
Monday - Friday (9 a.m. - 5 p.m. EST)

Inactive ingredients

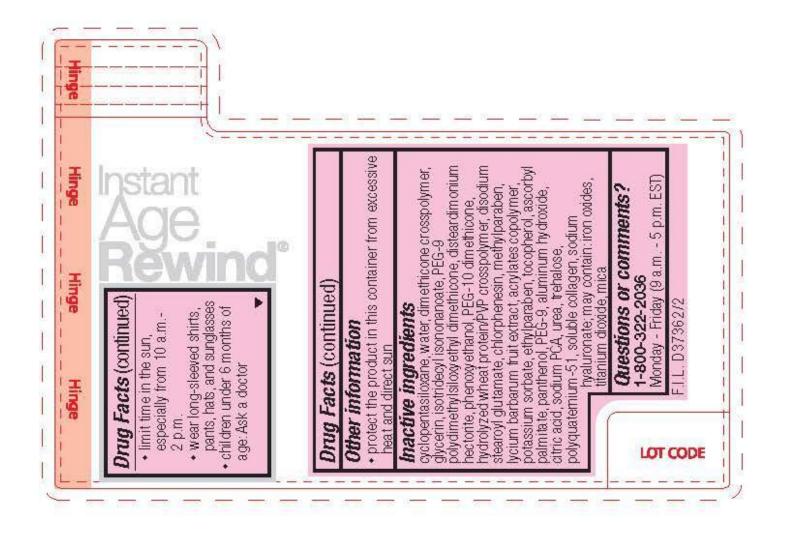
cyclopentasiloxane, water, dimethicone crosspolymer, glycerin, isotridecyl isononanoate, PEG-9 polydimethylsiloxyethyl dimethicone, disteardimonium hectorite, phenoxyethanol, PEG-10 dimethicone, hydrolyzed wheat protein/PVP crosspolymer, disodium stearoyl glutamate, chlorphenesin, methylparaben, lycium barbarum fruit extract, acrylates copolymer, potassium sorbate, ethylparaben, tocopherol, ascorbyl palmitate, panthenol, PEG-9, aluminum hydroxide, citric acid, sodium PCA, urea, trehalose, polyquaternium-51, soluble collagen, sodium hyaluronate; may contain: iron oxides, titanium dioxide, mica











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octinoxate lotion

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49967-007	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
GLYCERIN (UNII: PDC6A3C0OX)			
PHENOXYETHANOL (UNII: HIE492ZZ3T)			

Packaging					
	#	Item Code	tem Code Package Description		Marketing End Date
	1	NDC:49967-007- 01	20 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2010	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	01/01/2010	

Labeler - L'Oreal USA Products Inc (002136794)

Establishment				
Name	Address	ID/FEI	Business Operations	
L'Oreal USA, Inc.		624244349	manufacture(49967-007)	

Revised: 12/2023 L'Oreal USA Products Inc