FACIAL SUNSCREEN WITH ZINC SPF 50 DAYLOGIC- octinoxate 7.50% octisalate 5.00% oxybenzone 6.00% zinc oxide 7.00% cream Rite Aid

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.

Drug Facts

Active ingredients	Purpose
Octinoxate 7.5%	Sunscreen
Octisalate 5.0%	Sunscreen
Oxybenzone 6.0%	Sunscreen
Zinc Oxide 7.0%	

Uses • helps prevent sunburn

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

Directions

- apply liberally 15 minutes before sun exposure
- reapply: after 80 minutes of swimming or sweating
- immediately after towel drying at least every 2 hours
- **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 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-sleeve shirts, pants, hats, and sunglasses
- children under 6 months: Ask a doctor

Other information

• protect this product from excessive heat and direct sun.

Inactive ingredients

Aloe Barbadensis Leaf Extract,
Beeswax, Cetyl Dimethicone, Cetyl PEG/PPG-10/1 Dimethicone,
Cyclopentasiloxane, Cyclohexasiloxane, Diisopropyl Adipate, DMDM
Hydantoin, Ethylhexyl Palmitate, Ethylhexyl Stearate, Glycerin, Hydrated
Silica, Hydrogenated Castor Oil, Polyethylene, Sodium Chloride,
Theobroma Cacao (Cocoa) Seed Butter, Tocopheryl Acetate,



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Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-1214
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Octinoxate (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	Octinoxate	7.5 g in 100 g	
Octisalate (UNII: 4X49 Y0596W) (OCTISALATE - UNII:4X49 Y0596W)	Octisalate	5 g in 100 g	
Oxybenzone (UNII: 9500S7VE0Y) (OXYBENZONE - UNII:9500S7VE0Y)	Oxybenzone	6 g in 100 g	
Zinc Oxide (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	7 g in 100 g	

Inactive Ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
YELLOW WAX (UNII: 2ZA36H0S2V)		
CYCLOMETHICONE 6 (UNII: XHK3U310BA)		
CYCLOMETHICONE 5 (UNII: 0 THT5PCI0 R)		
Diisopropyl Adipate (UNII: P7E6 YFV72X)		
DMDM HYDANTO IN (UNII: BYR0546 TOW)		
Ethylhexyl Palmitate (UNII: 2865993309)		
Ethylhexyl Stearate (UNII: EG3PA2K3K5)		
Glycerin (UNII: PDC6A3C0OX)		
Hydrated Silica (UNII: Y6O7T4G8P9)		
Hydrogenated Castor Oil (UNII: ZF94AP8MEY)		
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)		
Sodium Chloride (UNII: 451W47IQ8X)		
CO CO A BUTTER (UNII: 5120 YT1CRR)		
ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)		
Triethoxycaprylylsilane (UNII: LDC331P08E)		
Water (UNII: 059QF0KO0R)		

Packaging				
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:11822-1214-1	28.3 g in 1 JAR; Type 0: Not a Combination Product	03/17/2017	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	03/17/2017	

Labeler - Rite Aid (014578892)

Registrant - Product Quest Mfg (927768135)

Establishment				
Name	Address	ID/FEI	Business Operations	
Product Quest Mfg		927768135	manufacture(11822-1214), label(11822-1214)	

Revised: 5/2018 Rite Aid