CREAL ZINK SUNSCREEN SPF 50 WELL AT WALGREENS- octinoxate 7.5% octisalate 5.0% zinc oxide 14.5% stick

Walgreens

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
Octinoxate 7.5%
Octisalate 5.0%
Zinc Oxide 14.5%

Purpose Sunscreen Sunscreen Sunscreen

Uses • helps prevent sunburn

Warnings

For external use only

Flammable: Do not use near heat, flame, or while smoking.

Do not use on damaged or broken skin

When using this product • Keep out of eyes. Rinse eyes with water to remove. • Keep away from face to avoid breathing it • Do not puncture or incinerate. Contents under pressure. Do not store at temperatures above 120°F.

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.

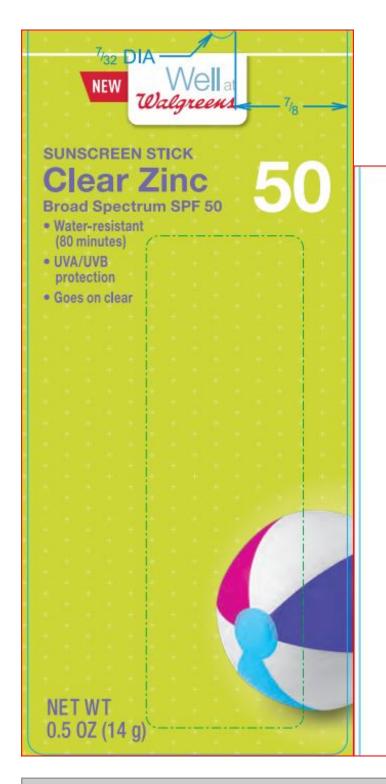
Directions

- spray liberally and spread evenly by hand 15 minutes before sun exposure
- reapply: after 80 minutes of swimming or sweating
- immediately after towel drying at least every 2 hours
- hold container 4 to 6 inches from the skin to apply
- do not spray directly into face. Spray on hands then apply to face.
- do not apply in windy conditions use in a well-ventilated area
- **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

Inactive ingredients

Aloe Barbadensis Leaf Extract Ascorbic Acid C12-15 Alkyl Benzoate Camellia Sinensis Leaf Extract Copernicia Cerifera (Carnauba) Wax
Dimethicone
Helianthus Annuus (Sunflower) Seed Oil
Hydrogenated Microcrystalline Wax
Hydrogenated Polyisobutene
Octyldodecanol
Paraffin
PEG-8 Dimethicone
Polyethylene
Retinyl Palmitate
Silica
Sodium Propoxyhydroxypropyl Thiosulfate Silica
Synthetic Wax
Tocopheryl Acetate

Triethoxycaprylylsilane



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Inactive ingredients Aloe Barbadensis Leaf Extract, Ascorbic Acid, C12 – 15 Alkyl Benzoate, Camellia Sinensis Leaf Extract, Copernicia Cerifera (Carnauba) Wax, Dimethicone, Helianthus Annus (Sunflower) Seed Oil, Hydrogenated Microcrystalline Wax, Hydrogenated Polyisobutene, Octyldodecanol, Paraffin, Peg-8 Dimethicone, Polyethylene, Retinyl Palmitate, Silica, Sodium Propoxyhydroxypropyl Thiosulfate Silica, Synthetic Wax, Tocopheryl Acetate, Triethoxycaprylysilane.

Other information • protect this product from excessive heat and direct sun. • may stain fabrics.

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Droduct	Information
Product	Intormation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0363-1117

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
Zinc Oxide (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	14.5 g in 100 g

Inactive Ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)		
GREEN TEA LEAF (UNII: W2ZU1RY8B0)		
CARNAUBA WAX (UNII: R12CBM0EIZ)		
Dimethicone (UNII: 92RU3N3Y1O)		
SUNFLOWER OIL (UNII: 3W1JG795YI)		
Octyldodecanol (UNII: 461N1O614Y)		
PARAFFIN (UNII: 19 O0 E3H2ZE)		
PEG-8 Dimethicone (UNII: GIA7T764OD)		
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)		
VITAMIN A PALMITATE (UNII: 1D1K0 N0 VVC)		
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)		
SODIUM PROPOXYHYDROXYPROPYL THIOSULFATE SILICA (UNII: 208G222332)		
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)		
Triethoxycaprylylsilane (UNII: LDC331P08E)		
ASCORBIC ACID (UNII: PQ6CK8PD0R)		

l	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:0363-1117-01	14 g in 1 TUBE; Type 0: Not a Combination Product	12/07/2015	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	12/07/2015	

Labeler - Walgreens (008965063)

Registrant - Product Quest Mfg (927768135)

Establishment			
Name	Address	ID/FEI	Business Operations
Product Quest Mfg		927768135	manufacture(0363-1117), label(0363-1117)

Revised: 2/2018 Walgreens