# TANNING SUNSCREEN SPF 4 DAYLOGIC- octinoxate 6% lotion 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.

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# **Drug Facts**

**Uses** • helps prevent sunburn

## Warnings

**Skin Cancer/Skin Aging Alert**: Spending time in the sun increases your risk of skin cancer and early skin aging. This product has been shown only to prevent sunburn, not skin cancer or early skin aging.

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
- children under 6 months: Ask a doctor

## Other information

• protect this product from excessive heat and direct sun.

# **Inactive ingredients**

Water, Mineral Oil, Polyethylene, Stearic Acid, Glyceryl Stearate SE, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadensis Leaf Juice, Ammonium Hydroxide, Ascorbyl Palmitate, Cetyl Alcohol, Diazolidinyl Urea, Disodium EDTA, Glycerin, Magnesium Aluminum Silicate, Methylparaben, PEG-40 Castor Oil, Phospholipids, Propylparaben, Retinyl Palmitate, Sorbitol, Theobroma Cacao (Cocoa) Seed Butter, Tocopheryl Acetate, Triethanolamine, Fragrance.



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# Active ingredients

Purpose

Octinoxate 6%....

.Sunscreen

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DISTRIBUTED BY: RITE AID 30 HUNTER LANE CAMP HILL, PA 17011 MADE IN THE USA WITH DOMESTIC AND FOREIGN COMPONENTS

look great feel great

## TANNING SUNSCREEN SPF 4 DAYLOGIC

octinoxate 6% lotion

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:11822-7788

Route of Administration TOPICAL

#### Active Ingredient/Active Moiety

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	Ingredient Name	Basis of Strength	Strength
	Octinoxate (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	Octinoxate	6 g in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
<b>AMMO NIA</b> (UNII: 5138 Q 19 F1X)	
Ascorbyl Palmitate (UNII: QN83US2B0N)	
Cetyl Alcohol (UNII: 936JST6JCN)	
Diazolidinyl Urea (UNII: H5RIZ3MPW4)	
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)	
Glycerin (UNII: PDC6A3C0OX)	
Glyceryl Stearate SE (UNII: FCZ5MH785I)	
Magnesium Aluminum Silicate (UNII: 6M3P64V0NC)	
Methylparaben (UNII: A2I8C7HI9T)	
Mineral Oil (UNII: T5L8T28FGP)	
PEG-40 Castor Oil (UNII: 4ERD2076EF)	
LECITHIN, SO YBEAN (UNII: 1DI56 QDM62)	
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	
Propylparaben (UNII: Z8IX2SC1OH)	
VITAMIN A PALMITATE (UNII: 1D1K0 N0 VVC)	
Sorbitol (UNII: 506T60A25R)	
Stearic Acid (UNII: 4ELV7Z65AP)	
COCOA BUTTER (UNII: 512OYT1CRR)	
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 903K93S3TK)	
WATER (UNII: 059QF0KO0R)	

# Item Code Package Description Marketing Start Date Marketing End Date	l	Packaging			
	l	# Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date
1 NDC:11822-7788-6 236 mL in 1 BOTTLE; Type 0: Not a Combination Product 11/10/2016	l	1 NDC:11822-7788-6	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/10/2016	

<b>Marketing Infor</b>			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	11/10/2016	

# **Labeler -** Rite Aid (014578892)

# **Registrant -** Product Quest Mfg (927768135)

Establishment				
Name	Address	ID/FEI	Business Operations	
Product Quest Mfg		927768135	manufacture(11822-7788), label(11822-7788)	

Revised: 5/2018 Rite Aid