# MOISTURE THERAPY DAILY SKIN DEFENSE BODY- octinoxate, octis alate, oxybenzone lotion Avon Products, Inc

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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## **Active Ingredients**

...... Sunscreen ...... Sunscreen

Octinoxate, 7.5%
Octisalate, 5.0%
Oxybenzone, 1.0%
Purpose
Sunscreen

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

#### Directions

- apply generously and evenly 15 minutes before sun exposure
- children under 6 month of age:ask a doctor
- reapply at least every 2 hours
- use a water resistant sunscreen if swimming or sweating

## Other Information

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

## **Inactive ingredients:**

WATER/EAU, GLYCERIN, BUTYLENE GLYCOL, GLYCERYL STEARATE, PEG-8, SILICA, CETYL ALCOHOL, IMIDAZOLIDINYL UREA, METHYLPARABEN, CARBOMER, DIMETHICONE, ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER, DISODIUM EDTA, HYDROGENATED LECITHIN, PARFUM/FRAGRANCE, SODIUM HYDROXIDE, POLYGLYCERYL-3 DIISOSTEARATE, ORYZA SATIVA (RICE) BRAN OIL, DIMETHICONOL, HYDROGENATED POLYISOBUTENE, ASCORBYL PALMITATE, PANTHENOL, SIMMONDSIA CHINENSIS (JOJOBA) SEED OIL, TOCOPHERYL ACETATE, RETINYL PALMITATE, GLYCINE SOJA (SOYBEAN) STEROLS, LECITHIN, GLYCINE SOJA (SOYBEAN) OIL, TOCOPHEROL.

#### Questions or comments?



## Drug Facts (continued)

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Drug Facts

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#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:10096-0291

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			
OCTISALATE (UNII: 4X49 Y0596W) (OCTISALATE - UNII:4X49 Y0596W)	OCTISALATE	50 mg in 1 mL			
OXYBENZONE (UNII: 9500S7VE0Y) (OXYBENZONE - UNII:9500S7VE0Y)	OXYBENZONE	10 mg in 1 mL			

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:10096-0291-1	400 mL in 1 BOTTLE			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part352	02/27/2013				

## Labeler - Avon Products, Inc (001468693)

Establishment			
Name	Address	ID/FEI	Business Operations
Avon Products, Inc		005149471	manufacture(10096-0291)

Revised: 2/2013 Avon Products, Inc