

MOISTURE THERAPY INTENSIVE HEALING AND REPAIR BODY- octinoxate, octisalate, 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.

☐Active Ingredients

☐☐Octinoxate, 7.5%.....

Octisalate, 5.0%.....

Oxybenzone, 1.0%.....

☐Purpose

☐ ☐.....Sunscreen

.....Sunscreen

.....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 months 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, ORYZA SATIVA (RICE) BRAN OIL, SIMMONDSIA CHINENSIS (JOJOBA) SEED OIL, HYDROGENATED POLYISOBUTENE, LECITHIN, HYDROGENATED LECITHIN, GLYCINE SOJA (SOYBEAN) STEROLS, GLYCINE SOJA (SOYBEAN) OIL, CETYL ALCOHOL, DIMETHICONE, CARBOMER, ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER, POLYGLYCERYL-3 DIISOSTEARATE, DISODIUM EDTA, IMIDAZOLIDINYL UREA, DIMETHICONOL, METHYLPARABEN, SODIUM HYDROXIDE, PARFUM/FRAGRANCE.

Questions or comments?Call **1-800-FOR-AVON** or **1-800-265-AVON** in Canada.



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OXYBENZONE, 1.0%.....	Sunscreen
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Drug Facts (continued)	
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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10096-0249
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 Y0596 W) (OCTISALATE - UNII:4X49 Y0596 W)	OCTISALATE	50 mg in 1 mL
OXYBENZONE (UNII: 95OOS7VE0 Y) (OXYBENZONE - UNII:95OOS7VE0 Y)	OXYBENZONE	10 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10096-0249-1	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	04/11/2011	

Labeler - Avon Products, Inc. (001468693)

Establishment

Name	Address	ID/FEI	Business Operations
Avon Products, Inc.		005149471	manufacture(10096-0249)

Revised: 2/2013

Avon Products, Inc.