SUN SOLAR DEFENSE HYDRATING - zinc oxide, ethylhexyl methoxycinnamate, benzophenone, ethylhexyl salicylate cream
Allure Labs, 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.

Drug Facts

Active Ingredients:

Zinc Oxide 5%

Ethylhexyl Methoxycinnamate (Octinoxate) 7.5%

Benzophenone-3 3%

Ethylhexyl Salicylate (Octisalate) 5%

Uses:

A hydrating UVA/UVB broad spectrum daily moisturizer. Provides ultimate protection against the aging effects of the sun and other environmental exposures. Enriched with essential vitamins and anti-oxidants that prevent free radical damage and preserving skin hydration for the entire day.

Water-resistant

Paraben-free

Directions:

Apply to cleaned skin every morning for ultimate protection.

Indications: Dry, dehydrated, rosacea, sensitive.

Warnings:

For External use only

When using this product-

Keep out of eyes. If contact occurs rinse with water.

Discontinue use if irritation or redness occurs

Keep out of reach of children

Inactive Ingredients:

Deionized water (Aqua), C12-15 Alkyl Benzoate, Glyceryl Stearate (and) PEG-100 Stearate, Glycerin, Sodium Hyaluronate, Imperata Cylindrica Root Extract, Octyl Dodecyl Neopentanoate, Sorbitol, Caprylic/Capric Truglyceride, Camellia sinensis (Green Tea) Leaf Extract, Biosaccharide Gum-1, Algae Extract and Mugwort Extract, Caprylyl Glycol and Phnoxyethanol and Hexylene Glycol, Sodium PCA, Lecithin, Sorbitan Stearate, Symphytum Officinale Extract and Plantago Ovata Seed Extract, Yeast Extract, Aloe Barbadensis Leaf Extract, Spiraea Ulmaria Flower Extract (and) Centella Asiatica Extract, Magnesium Aluminum Silicate, Tocopheryl Acetate, Ascorbyl Palmitate, Disodium EDTA, Dipotassium Glycyrrizate, Amino Acids.

Image of the Product:

Distributor:

Image International

Palm Beach, FL 33411 USA

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62742-4037

Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	50 mg in 1 mL
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 mL
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	30 mg in 1 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:62742-4037-1	118 mL in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	01/01/2010	

Labeler - Allure Labs, Inc. (926831603)

Revised: 6/2010 Allure Labs, Inc.