

SUN SOLAR DEFENSE ORGANIC SPF30 - zinc oxide 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

Purpose:

Sunscreen

Uses:

- An organic 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
- Chemical free

Directions:

Apply liberally 15-30minute prior to sun exposure. Reapply after prolonged swimming or vigorous activity.

Indications:

Sensitive skin highly exposed to sun

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:

Water (Aqua), Cyclomethicone, Glycerin, Glyceryl Stearate (and) PEG 100 Stearate, Sorbitol, Imperata Cylindrica Root Extract, Polyacrylamide and C13-14 Isoparaffin and Laureth-7, Caprylyl Glycol and Phenoxyethanol and Hexylene Glycol, Lecithin, Ascorbyl Palmitate (Vitramin C), Xanthan Gum, Dipotassium Glyrrhizate, Olea Europea (Olive) Leaf Extract, Disodium EDTA.

Distributor:

Image International

Palm Beach, FL 33411 USA

www.imageskincare.com

Image of the product:



SUN SOLAR DEFENSE ORGANIC SPF30

zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62742-4038
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	190 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62742-4038-1	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)

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Allure Labs, Inc.