

IMAGE MD LIGHTENING RX - hydroquinone 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

INDICATIONS AND USAGE SECTION:

For Physician Dispensary Only

An effective lightening creme that diminish facial and body discolorations.

DIRECTIONS:

Apply to affected areas in the evening. Apply a sunblock during the day, or as directed by a physician.

Paraben-free

Contains Vitamin C and E

ACTIVE INGREDIENTS:

Hydroquinone 4%

INACTIVE INGREDIENTS:

Water, Glyceryl Monostearate, Polyoxyl 100 Stearate, Butylene Glycol, Stearyl Alcohol, Stearic Acid, Caprylic and Capric Triglyceride, Linoleic Acid, Soy Phospholipids, Dimethicone, C12-15 Alkyl Benzoate, Xanthan Gum, Magnesium Aluminum Silicate, Cetyl Alcohol, Stearyl Alcohol, Glycerin, Licorice, Phenoxyethanol, Ethylhexylglycerin, Hexylene Glycol, Lactic Acid, Disodium EDTA, Dipotassium Glycyrrizinate, Vitamin E, Sodium Metabisulfite, Fragrance-Orange.

WARNING:

For external use only.

Keep out of reach of children

DISTRIBUTOR:

Image International

Palm Beach, FL 33411 USA

IMAGE OF PRINCIPAL DISPLAY PANEL:



IMAGE MD LIGHTENING RX

hydroquinone cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62742-4046
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)	HYDROQUINONE	40 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62742-4046-1	29.6 mL in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358	01/01/2010	

Labeler - Allure Labs, Inc. (926831603)

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Allure Labs, Inc.