SCARLIGHT MD - hydroquinone liquid
Scarguard Labs, LLC

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

Hydroquinone 2%

Purpose

Skin Lightener

Uses

- lightens dark (brownish) discoloration in the skin such as age and liver spots

Warnings

For external use only

Do not use

- on children under 12 years of age. Consult a doctor.
- on mucous membranes

When using this product

- mild irritation may occur
- avoid contact with eyes. If contact occurs, rinse with water.

Stop use and ask a doctor if

- irritation becomes severe
- condition worsens

Keep out of reach of children.

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

Directions

- adults: brush a small amount twice daily. Rub in.
- limit sun exposure and use a sunscreen, a sun blocking agent or protective clothing to cover bleached skin when using and after using this product in order to prevent darkening from reoccurring
- discontinue if symptoms persist for more than 3 months
- children under 12 years of age, consult a doctor before use

Other Information

- store at 15º to 30°C (59 to 86°F)
- keep bottle tightly closed or product will evaporate

Inactive Ingredients

retinoic acid, melatonin, MSM, BHT, na metabisulfite, arbutin, cystamine, licorice root, dandelion root, hydroxyanisole, ascorbic acid, hydroxypropylcellulose, kojic acid, azelaic acid, acetone, propylene glycol, ethyl alcohol (SDA), distilled water q.s.

Questions

call
1-877-566-5935

Carton 15mL

Carton 30mL

SCARLIGHT MD

hydroquinone liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:64269-9902
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYDROQUINONE (UNII: XV74C1N1AE) (Hydroquinone - UNII:XV74C1N1AE)	HYDROQUINONE	20 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Propylene Glycol (UNII: 6DC9Q167V3)	
Alcohol (UNII: 3K9958V90M)	
Acetone (UNII: 1364PS73AF)	
Azelaic Acid (UNII: F2VW3D43YT)	
Hydroxypropyl Cellulose (UNII: RFW2ET671P)	
Ascorbic Acid (UNII: PQ6CK8PD0R)	
Butylated Hydroxyanisole (UNII: REK4960K2U)	
TARAXACUM OFFICINALE ROOT (UNII: 9DE5YCO0RU)	
Licorice (UNII: 61ZBX54883)	
Arbutin (UNII: C5INA23HXF)	
Sodium Metabisulfite (UNII: 4VON5FNS3C)	

Butylated Hydroxytoluene (UNII: 1P9D0Z171K)	
Dimethyl Sulfone (UNII: 9H4PO4Z4FT)	
Melatonin (UNII: JL5DK93RCL)	
Tretinoin (UNII: 5688UTC01R)	
Water (UNII: 059QF0KO0R)	
Kojic Acid (UNII: 6K23F1TT52)	
Cystamine (UNII: R110LV8L02)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64269-9902-8	30 mL in 1 BOTTLE, WITH APPLICATOR		
2	NDC:64269-9902-7	15 mL in 1 BOTTLE, WITH APPLICATOR		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part358	11/01/2002	

Labeler - Scarguard Labs, LLC (842204575)

Registrant - Scarguard Labs, LLC (842204575)

Revised: 12/2009 Scarguard Labs, LLC