I-MAX LIGHTENING L- hydroquinone cream
MAXLIFE USA, 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:
HYDROQUINONE USP 2%

PURPOSE:

SKIN LIGHTENING

USES:

FOR THE GRADUAL FADING OF DARK AREAS OF THE SKIN.

WARNINGS:

AVOID CONTACT WITH EYES. SOME USERS MAY EXPERIENCE MILD SKIN IRRITATION.

IF IRRITATION BECOMES SEVERE, STOP USE AND CONSULT A DOCTOR.

THIS PRODUCT IS NOT INTENDED FOR USE IN THE PREVENTION OF SUNBURN AND CONTAINS AN ALPHA HYDROXY ACID (AHA) THAT MAY INCREASE YOUR SKIN'S SENSITIVITY TO THE SUN AND PARTICULARLY THE POSSIBILITY OF SUNBURN. SUN EXPOSURE SHOULD BE LIMITED BY USING A SUNSCREEN AGENT OR PROTECTIVE CLOTHING TO COVER BLEACHED SKIN AFTER TREATMENT IS COMPLETED TO PREVENT DARKENING FROM REOCCURING.

DIRECTIONS:

ADULTS: APPLY A SMALL AMOUNT AS A THIN LAYER ON THE AFFECTED AREA TWICE DAILY, OR USE AS DIRECTED BY A DOCTOR. IF NO IMPROVEMENT IS SEEN AFTER 3 MONTHS OF TREATMENT, USE OF THIS PRODUCT SHOULD BE DISCONTINUED. LIGHTENING EFFECT OF THIS PRODUCT MAY NOT BE NOTICEABLE WHEN USED ON VERY DARK SKIN.

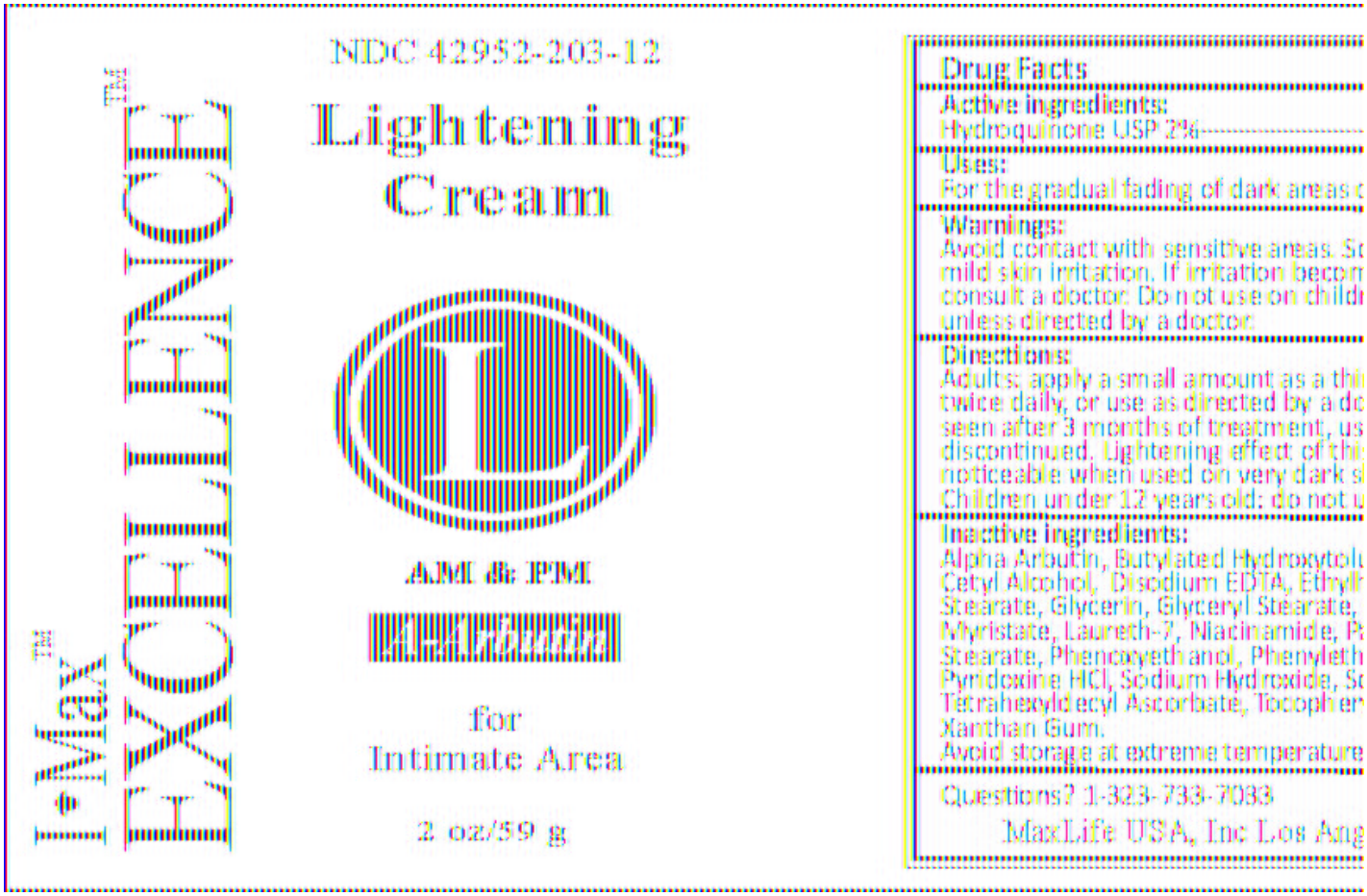
CHILDREN UNDER 12 YEARS OF AGE: DO NOT USE UNLESS DIRECTED BY A DOCTOR.

INACTIVE INGREDIENTS:

ALPHA ARBUTIN, C13-14 ISOPARAFFIN, CETYL ALCOHOL, ETHYLHEXYLGLYCERIN, ETHYLHEXYL STEARATE, GLYCERIN, GLYCERYL STEARATE, GLYCOLIC ACID, ISOPROPYL MYRISTATE, LAURETH-7, NIACINAMIDE, PANTOTHENIC ACID, PEG-100 STEARATE, PHENOXYETHANOL, PHENYLETHYL RESORCINOL, POLYACRYLAMIDE, PYRIDOXINE HCL, SODIUM HYDROXIDE, SODIUM METABISULFITE, TETRAHEXYLDECYL ASCORBATE, WATER (AQUA), XANTHAN GUM.

QUESTIONS? 1-323-733-7033

KEEP OUT OF REACH OF CHILDREN.



I-MAX LIGHTENING L

hydroquinone cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:42952-203
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE)	HYDROQUINONE	2 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
GLYCOLIC ACID (UNII: 0WT12SX38S)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

LAURETH-7 (UNII: Z95S6G8201)	
NIACINAMIDE (UNII: 25X51I8RD4)	
PANTOTHENIC ACID (UNII: 19F5HK2737)	
PEG-100 STEARATE (UNII: YD01N1999R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYACRYLAMIDE (10000 MW) (UNII: E2KR9C9V2I)	
PYRIDOXINE HYDROCHLORIDE (UNII: 68Y4CF58BV)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM METABISULFITE (UNII: 4VON5FNS3C)	
TETRAHEXYLDECYL ASCORBATE (UNII: 9LBV3F07AZ)	
WATER (UNII: 059QF0KO0R)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ALPHA-ARBUTIN (UNII: 72VUP07IT5)	
PHENYLETHYL RESORCINOL (UNII: G37UFG162O)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42952-203-12	59 g in 1 TUBE; Type 0: Not a Combination Product	09/18/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part358A	09/18/2012	

Labeler - MAXLIFE USA, INC. (785111431)

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