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.

M	NIDC 42952-203-12	Drug Facts
	Lightening	Active ingredients: Hydroquinone USP 2% Uses:
\bigcirc	Creann	For the gradual fading of dark areas of Warnings:
	and the second s	Avoid contact with sensitive areas. So mild skin initiation. If imitation become consult a doctor. Do not use on child
		unless directed by a doctor: Directions: Adults: apply a small amount as a thi twice daily, or use as directed by a do seen after 3 months of treatment, us discontinued. Lightening effect of thi noticeable when used on very dark s
		discontinued. Lightening effect of thi noticeable when used on very darks Children under 12 years old: do not t Inactive ingredients:
	AVDAL 999 DADAT	Alpha Arbutin, Butylated Hydroxytol Cetyl Alcohol, Disodium EDTA, Ethyl Stearate, Glycerin, Glyceryl Stearate,
	A-A ributin	Myristate, Laureth-7, Niacinamide, P Stearate, Phencxyeth anol, Phenyleth Pyridoxine HCl, Soldium Hydroxide, S Tetrahexyld ecyl Ascorbate, Tocopher
	for Intimate Area	Tecranexyld ecyl Ascorbate, Tocoph er Xanthan Gum. Avoid storage at extreme temperature
•	2 oz/59 g	Questions? 1-323-733-7033 MaxLife USA, Inc Los Ang

I-MAX LIGHTENING I	L				
hydroquinone cream					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source	2)	NDC:4295	52-203
Route of Administration	TOPICAL				
Active Ingredient/Active Mo	iety				
II	ıgredient Name		Basis of St	trength	Strength
HYDROQUINONE (UNII: XV74C1N1AE) (HYDROQUINONE - UNII:XV74C1N1AE) HYDROQUINONE				ONE	2 g in 100 g
Inactive Ingredients					
	Ingredient Name			S	Strength
C13-14 ISOPARAFFIN (UNII: E4F12R	OE70)				
CETYL ALCOHOL (UNII: 936JST6JC	CN)				
ETHYLHEXYLGLYCERIN (UNII: 147)	D247K3P)				
ETHYLHEXYL STEARATE (UNII: EG	3PA2K3K5)				
GLYCERIN (UNII: PDC6A3C0OX)					
GLYCERYL MONOSTEARATE (UNI	I: 230OU9XXE4)				
GLYCOLIC ACID (UNII: 0WT12SX38	S)				
ISOPROPYL MYRISTATE (UNII: 0 R	E8K4LNJS)				

L 1	AURETH-7 (UNII: Z95	S6G8201)						
NIACINAMIDE (UNII: 25X5118 RD4)								
PANTO THENIC ACID (UNII: 19F5HK2737)								
Pł	PEG-100 STEARATE (UNII: YD01N1999R)							
Pł	PHENOXYETHANOL (UNII: HIE492ZZ3T)							
PO	POLYACRYLAMIDE (10000 MW) (UNII: E2KR9C9V2I)							
PY	PYRIDO XINE HYDRO CHLO RIDE (UNII: 68 Y4CF58 BV)							
so	SODIUM HYDROXIDE (UNII: 55X04QC32I)							
so	DIUM METABISUL	F ITE (UNII: 4VON5FNS3C)						
TI	ETRAHEXYLDECYL	ASCORBATE (UNII: 9LBV3F07AZ)						
W	ATER (UNII: 059QF0	KO0R)						
X	ANTHAN GUM (UNII:	TTV12P4NEE)						
ALPHA-ARBUTIN (UNII: 72VUP07IT5)								
PHENYLETHYL RESORCINOL (UNII: G37UFG1620)								
Pł	IENYLETHYL RESO	RCINOL (UNII: G37UFG162O)						
PI	IENYLETHYL RESO	RCINOL (UNII: G37UFG162O)						
PI	IENYLETHYL RESO	RCINOL (UNII: G37UFG162O)						
	IENYLETHYL RESO ackaging	RCINOL (UNII: G37UFG162O)						
		RCINOL (UNII: G37UFG162O) Package Description		Marketing Start Date	Marketing End Date			
P #	ackaging			Marketing Start Date 09/18/2012	Marketing End Date			
P #	ackaging Item Code	Package Description		0	Marketing End Date			
P #	ackaging Item Code	Package Description		0	Marketing End Date			
P # 1	ackaging Item Code NDC:42952-203-12	Package Description 59 g in 1 TUBE; Type 0: Not a Combinat		0	Marketing End Date			
P # 1	ackaging Item Code NDC:42952-203-12 Iarketing Info	Package Description 59 g in 1 TUBE; Type 0: Not a Combinat prmation	ion Product	09/18/2012				
P # 1	ackaging Item Code NDC:42952-203-12	Package Description 59 g in 1 TUBE; Type 0: Not a Combinat prmation	ion Product	09/18/2012	Marketing End Date			
P # 1	ackaging Item Code NDC:42952-203-12 Iarketing Info	Package Description 59 g in 1 TUBE; Type 0: Not a Combinat 59 mation 9 rmation 9 Application Number or Monog	ion Product	09/18/2012				

Labeler - MAXLIFE USA, INC. (785111431)

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MAXLIFE USA, INC.