I-MAX EXCELLENCE- avobenzone octinoxate oxybenzone lotion
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:

AVOBENZONE 2.5%

OCTINOXATE 7.5%

OXYBENZONE 5%

PURPOSE:

SUNSCREEN

USES:

HELPS PREVENT SUNBURNS. HIGHER SPF GIVES MORE SUNBURN PROTECTION.

WARNINGS

WARNINGS:

FOR EXTERNAL USE ONLY.

KEEP OUT OF REACH OF CHILDREN.

WHEN USING THIS PRODUCT

KEEP OUT OF EYES. RINSE WITH WATER TO REMOVE.

STOP USE AND ASK A DOCTOR IF

A RASH OR IRRITATION DEVELOPS AND LASTS. IF SWALLOWED, GET

MEDICAL HELP OR CONTACT A POISON CONTROL CENTER RIGHT AWAY.

DIRECTIONS

APPLY TO CLEAN SKIN 15 MINUTES BEFORE SUN EXPOSURE. REAPPLY

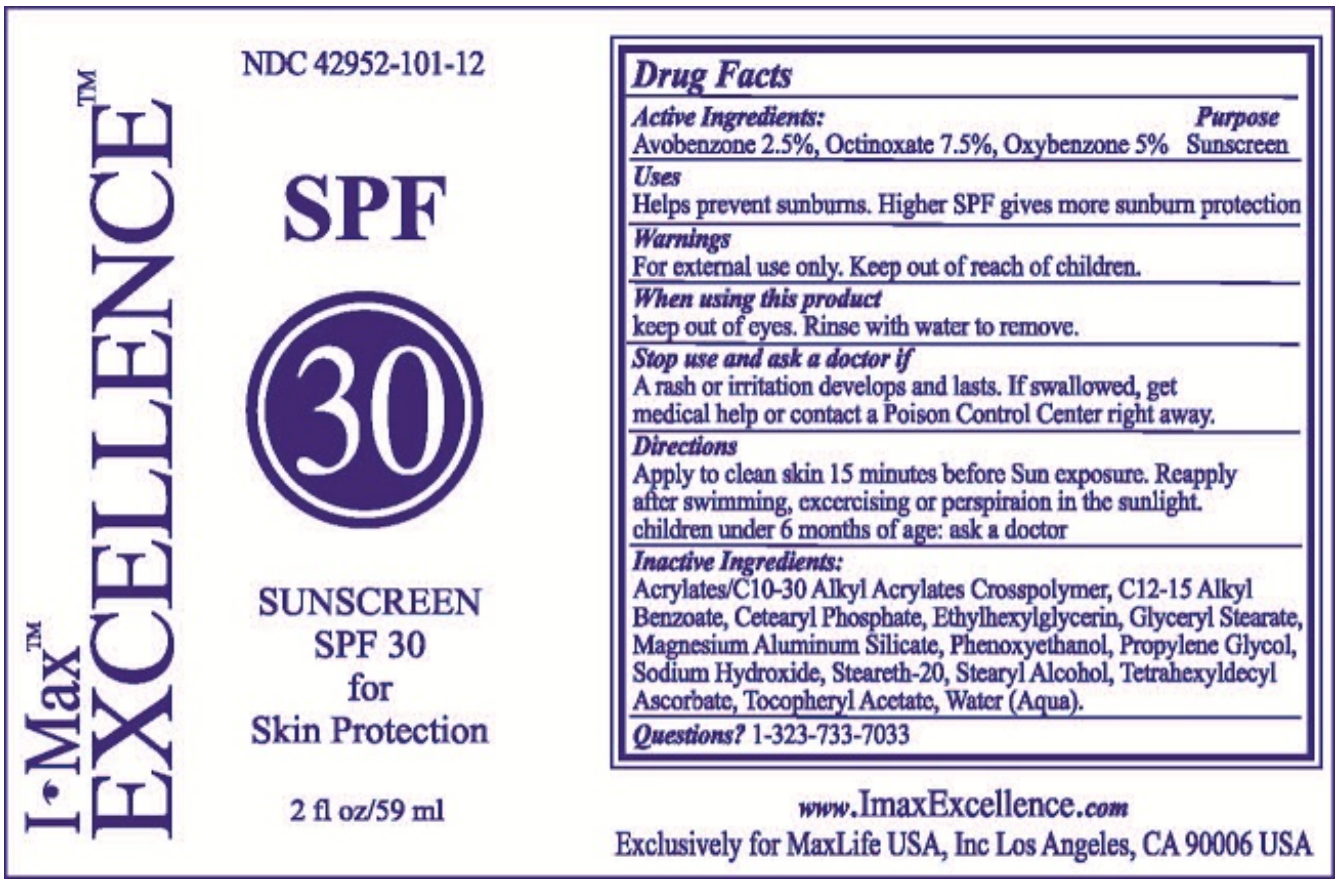
AFTER SWIMMING, EXCERCISING OR PERSPIRATION IN THE SUNLIGHT.

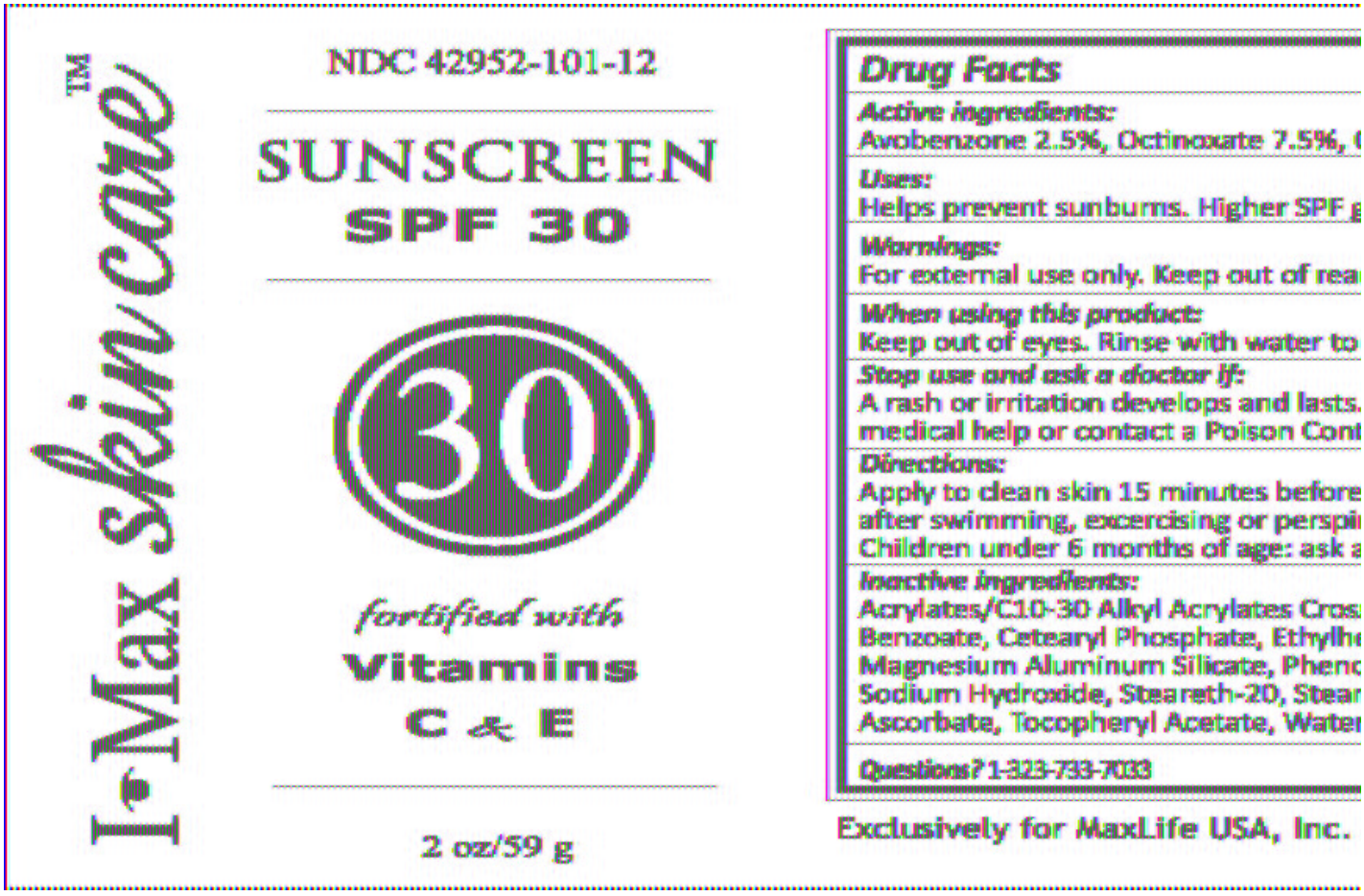
CHILDREN UNDER 6 MONTHS OF AGE: ASK A DOCTOR.

INACTIVE INGREDIENTS:

ALKYL BENZOATE, ACRYLATES/C10-30 ALKYL ACRYLATES CROSSPOLYMER, C12-15 GLYCERYL STEARATE, CETEARYL PHOSPHATE, ETHYLHEXYLGLYCERIN, MAGNESIUM ALUMINUM SILICATE, PHENOXYETHANOL, PROPYLENE GLYCOL, SODIUM HYDROXIDE, STEARETH-20, STEARYL ALCOHOL, TETRAHEXYLDECYL ASCORBATE, TOCOPHERYL ACETATE, WATER (AQUA).

QUESTIONS? 1-323-733-7033





I-MAX EXCELLENCE

avobenzone octinoxate oxybenzone lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:42952-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	2.5 g in 100 mL
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7.5 g in 100 mL
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	5 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
CETYL PHOSPHATE (UNII: VT07D6X67O)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
STEARETH-20 (UNII: L0Q8IK9E08)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
TETRAHEXYLDECYL ASCORBATE (UNII: 9LBV3F07AZ)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:42952-101-12	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/15/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	03/15/2012	

Labeler - MAXLIFE USA, INC. (785111431)

Registrant - MAXLIFE USA, INC. (785111431)

Establishment

Name	Address	ID/FEI	Business Operations
CUSTOM RESEARCH LABS INC		028611598	manufacture(42952-101)

Revised: 11/2018

MAXLIFE USA, INC.