EQUALINE ALL DAY MOISTURIZER- octis alate, avobenzone, homos alate, octocrylene lotion SUPERVALU 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

ACTIVE INGREDIENTS

OCTISALATE 5.0%, AVOBENZONE 3.0%, HOMOSALATE 3.0%, OCTOCRYLENE 2.6%

PURPOSE

SUNSCREEN

USE(S)

HELPS PREVENT SUNBURN. IF USED AS DIRECTED WITH OTHER SUN PROTECTION MEASURES (SEE DIRECTIONS), DECREASES THE RISK OF SKIN CANCER AND EARLY SKIN AGING CAUSED BY THE SUN.

WARNINGS

FOR EXTERNAL USE ONLY.

DO NOT USE

ON DAMAGED OR BROKEN SKIN.

WHEN USING THIS PRODUCT

KEEP OUT OF EYES. RINSE WITH WATER TO REMOVE.

STOP USING THIS PRODUCT AND ASK A DOCTOR IF

RASH OCCURS.

KEEP OUT OF REACH OF CHILDREN

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY.

DIRECTIONS

APPLY LIBERALLY 15 MINUTES BEFORE SUN EXPOSURE. USE A WATER RESISTANT SUNSCREEN IF SWIMMING OR SWEATING. REAPPLY AT LEAST EVERY 2 HOURS. CHILDREN UNDER 6 MONTHS: ASK A DOCTOR.

SUN PROTECTION MEASURES. SPENDING TIME IN THE SUN INCREASES YOUR RISK OF SKIN CANACER AND EARLY SKIN AGING. TO DECREASE THIS RISK, REGULARLY USE A SUNSCREEN WITH A BROAD SPECTRUM SPF VALUE OF 15 OR HIGHER AND OTHER SUN PROTECTION MEASURES INCLUDING: LIMIT TIME IN THE SUN, ESPECIALLY FROM 10 A.M. - 2 P.M., WEAR LONG-SLEEVED SHIRTS, PANTS, HATS, AND SUNGLASSES.

OTHER INFORMATION

PROTECT THIS PRODUCT FROM EXCESSIVE HEAT AND DIRECT SUN.

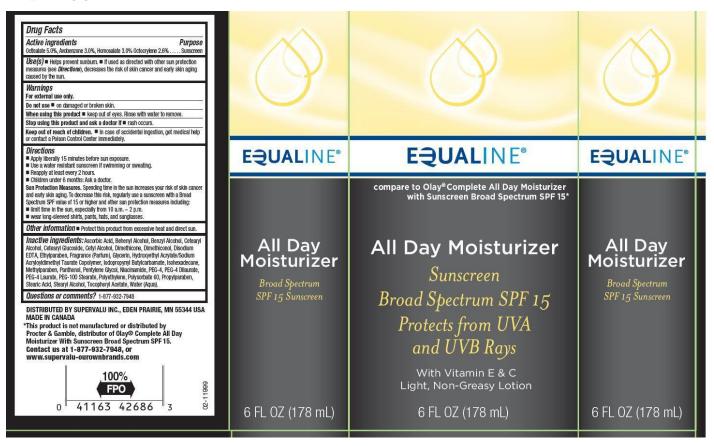
INACTIVE INGREDIENTS:

ASCORBIC ACID, BEHENYL ALCOHOL, BENZYL ALCOHOL, CETEARYL ALCOHOL, CETEARYL GLUCOSIDE, CETYL ALCOHOL, DIMETHICONE, DIMETHICONOL, DISODIUM EDTA, ETHYLPARABEN, FRAGRANCE (PARFUM), GLYCERIN, HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER, IODOPROPYNYL BUTYLCARBAMATE, ISOHEXADECANE, METHYLPARABEN, PANTHENOL, PENTYLENE GLYCOL, NIACINAMIDE, PEG-4, PEG-4 DILAURATE, PEG-4 LAURATE, PEG-100 STEARATE, POLYETHYLENE, POLYSORBATE 60, PROPYLPARABEN, STEARIC ACID, STEARYL ALCOHOL, TOCOPHERYL ACETATE, WATER (AQUA).

QUESTIONS OR COMMENTS?

1-877-932-7948

LABEL COPY



Product Information Product Type HUMAN OTC DRUG Route of Administration HUMAN OTC DRUG TOPICAL

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
OCTISALATE (UNII: 4X49 Y0 59 6 W) (OCTISALATE - UNII:4X49 Y0 59 6 W)	OCTISALATE	50 mg in 1 mL			
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL			
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	30 mg in 1 mL			
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	26 mg in 1 mL			

Inactive Ingredients	
Ingredient Name	Strength
ASCORBIC ACID (UNII: PQ6 CK8 PD0 R)	
DOCOSANOL (UNII: 9G10E216XY)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETEARYL GLUCO SIDE (UNII: 09 FUA47KNA)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIMETHICO NE (UNII: 92RU3N3Y1O)	
DIMETHICONOL (2000 CST) (UNII: T74O12AN6Y)	
EDETATE DISO DIUM (UNII: 7FLD91C86K)	
ETHYLPARABEN (UNII: 14255EXE39)	
GLYCERIN (UNII: PDC6 A3C0 OX)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86 FQE 96 TZ4)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
ISOHEXADECANE (UNII: 918 X10 UF1E)	
METHYLPARABEN (UNII: A2I8 C7HI9 T)	
PANTHENOL (UNII: WV9CM0O67Z)	
PENTYLENE GLYCOL (UNII: 50 C130 7 PZG)	
NIACINAMIDE (UNII: 25X51I8RD4)	
POLYETHYLENE GLYCOL 200 (UNII: R95B8J264J)	
PEG-4 DILAURATE (UNII: KCR71CW036)	
PEG-4 LAURATE (UNII: AYF4VM3N1Z)	
PEG-100 STEARATE (UNII: YD01N1999R)	
HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
.ALPHATO COPHERO L ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41163-356-06	178 mL in 1 BOTTLE, PLASTIC		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	03/15/2013	

Labeler - SUPERVALU INC. (006961411)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment				
Name	Address	ID/FEI	Business Operations	
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(41163-356)	

Revised: 3/2013 SUPERVALU INC.