FLAWLESS FINISH PERFECTLY NUDE MAKEUP BROAD SPECTRUM SUNSCREEN SPF 15 SHADE CHESTNUT- octinoxate and titanium dioxide lotion Elizabeth Arden, 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.

250483

DESCRIPTION

Makeup that feels like your own skin, only better. Light, breathable formula smooths on seamlessly for a naturally luminous look that lasts all day. Minimizes the look of pores and imperfections immediately and over time with continued use. Hydro-pigment color technology, vitamins and antioxidants protect and perfect, so skin looks beautifully even-toned and flawless.

INDICATIONS & USAGE

smooth onto face.

WARNINGS For external use only

OTC - ACTIVE INGREDIENT

OCTINOXATE 4.00%.....Sunscreen

TITANIUM DIOXIDE 1.26%....Sunscreen

INACTIVE INGREDIENT

WATER/AQUA/EAU, ISOHEXADECANE, HEXYL LAURATE, MICA, GLYCERIN, POLYGLYCERYL-4 ISOSTEARATE, POTASSIUM CETYL PHOSPHATE, CETEARYL ALCOHOL, ASCORBYL PALMITATE, BETAINE, CARBOMER, GLYCERYL STEARATE, HYDROLYZED VEGETABLE PROTEIN, LECITHIN, MAGNESIUM ALUMINUM SILICATE, PEG-100 STEARATE, POLYGLYCERYL-10 MYRISTATE, SODIUM HYDROXIDE, TOCOPHERYL ACETATE, TREMELLA FUCIFORMIS SPOROCARP EXTRACT, CHLORPHENESIN, PHENOXYETHANOL, POTASSIUM SORBATE, SODIUM BENZOATE, SODIUM DEHYDROACETATE [MAY CONTAIN/PEUT CONTENIR (+/-) IRON OXIDES (CI 77491, CI 77492, CI 77499), TITANIUM DIOXIDE (CI 77891).

OTC - DO NOT USE

Do not

use on damaged or broken skin

OTC - KEEP OUT OF REACH OF CHILDREN Keep out of reach of children.

OTC - PURPOSE

OTC - STOP USE Stop use and ask a doctor if rash occurs

OTC - WHEN USING

keep out of eyes. Rinse with water to remove.



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octinoxate and titanium dioxide lotion

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Sour	rce)	NDC:679	38-2030
Route of Administration	TOPICAL				
A - 4' T 1' 4/A - 4' B.A					
Active Ingredient/Active Moiety					
Ingredient Name		1	Basis of Stren	gth	Strength

OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	60 mg in 1500 mg
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP) (TITANIUM DIO XIDE - UNII:15FIX9 V2JP)	TITANIUM DIO XIDE	18.9 mg in 1500 mg

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
ISOHEXADECANE (UNII: 918 X10 UF1E)		
HEXYL LAURATE (UNII: 4CG9F9W01Q)		
MICA (UNII: V8A1AW0880)		
GLYCERIN (UNII: PDC6A3C0OX)		
POLYGLYCERYL-4 ISOSTEARATE (UNII: 820 DPX33S7)		
PO TASSIUM CETYL PHO SPHATE (UNII: 03KCY6P7UT)		
ASCORBYL PALMITATE (UNII: QN83US2B0N)		
BETAINE (UNII: 3SCV180C9W)		
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)		
PEG-100 STEARATE (UNII: YD01N1999R)		
SO DIUM HYDRO XIDE (UNII: 55X04QC32I)		
CHLORPHENESIN (UNII: I670 DAL4SZ)		
PHENO XYETHANOL (UNII: HIE492ZZ3T)		
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SO DIUM DEHYDRO ACETATE (UNII: 8 W46 YN971G)		

Product Characteristics			
Color BR	ROWN (BROWN)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:67938-2030-1	1 in 1 BOX			
1 NDC:67938-2030-2	1500 mg in 1 BOTTLE			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	07/17/2013	

Labeler - Elizabeth Arden, Inc (849222187)

Establishment			
Name	Address	ID/FEI	Business Operations
Intercos		883457061	manufacture(67938-2030)

Revised: 8/2013 Elizabeth Arden, Inc