FLAWLESS FINISH PERFECTLY NUDE MAKEUP BROAD SPECTRUM SUNSCREEN SPF 15 SHADE LINEN- 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.

250369

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.

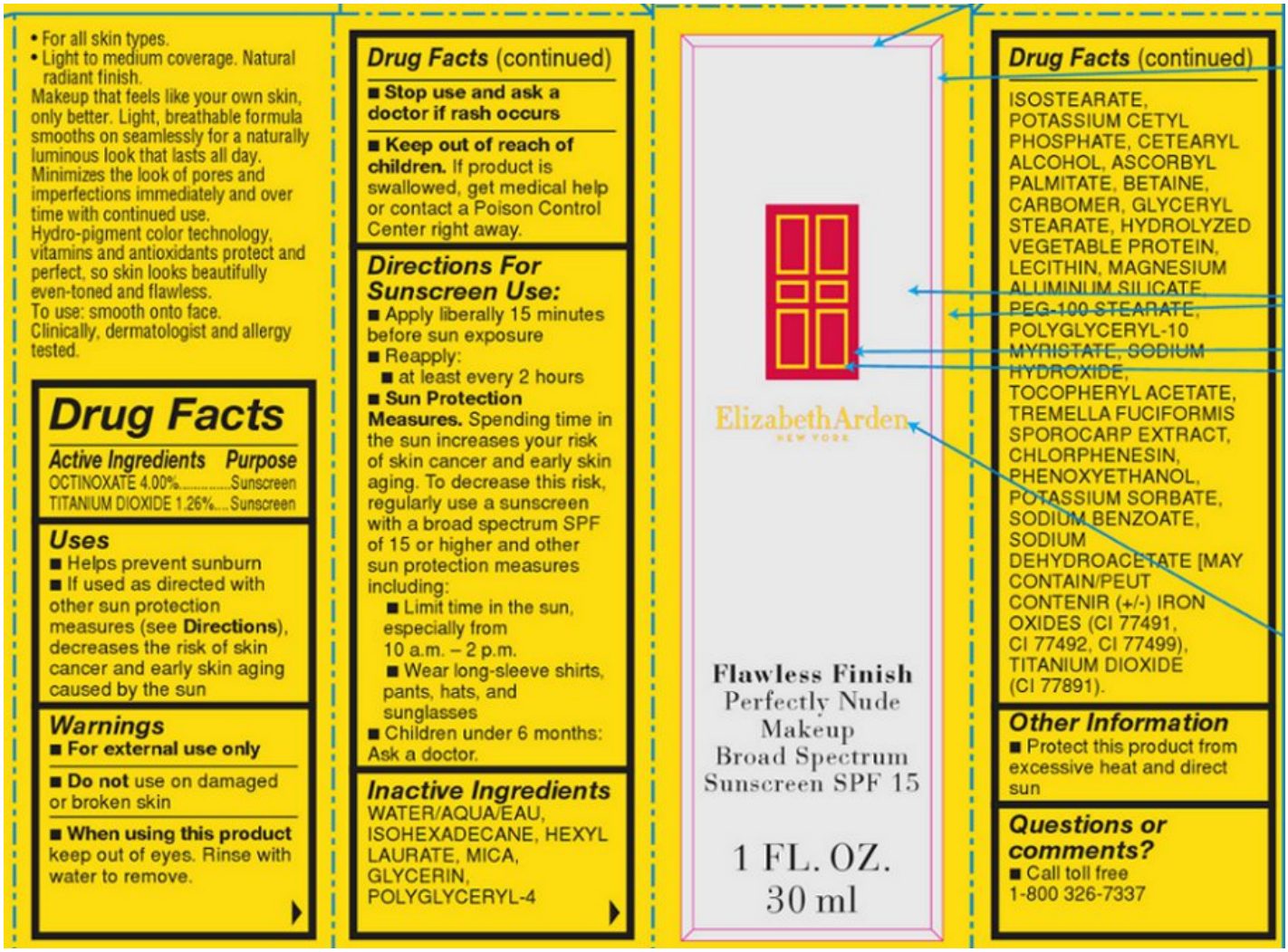
OTC - PURPOSE

Provides SPF 15 sun protection.

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 (Source)**	NDC:67938-2005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	60 mg in 1500 mg
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	18.9 mg in 1500 mg

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ISOHEXADECANE (UNII: 918X1OUF1E)	
HEXYL LAURATE (UNII: 4CG9F9W01Q)	
MICA (UNII: V8A1AW0880)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYGLYCERYL-4 ISOSTEARATE (UNII: 820DPX33S7)	
POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT)	
ASCORBYL PALMITATE (UNII: QN83US2B0N)	
BETAINE (UNII: 3SCV180C9W)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
PEG-100 STEARATE (UNII: YD01N1999R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CHLORPHENESIN (UNII: I670DAL4SZ)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM DEHYDROACETATE (UNII: 8W46YN971G)	

Product Characteristics

Color	BROWN (LIGHT BEIGE)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67938-2005-1	1 in 1 BOX		
1	NDC:67938-2005-2	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(67938-2005)