# FLAWLESS FINISH DUAL PERFECTION MAKEUP SPF 8 VANILLA- titanium dioxide powder

# 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.

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**BP1157** 

#### DESCRIPTION

For normal to oily skin. Adjustable coverage. Matte finish. Ultra fine powder and foundation in one. Wet it's a soft matte makeup. Dry it's a perfectly matte makeup or finishing powder. Ideal for touch-ups. SPF 8 provides sun protection. Clinically, dermatologist, and allergy tested.

### INDICATIONS AND USAGE

To Use: Smooth on to face before sun exposure.

# **WARNINGS**

Warning: For external use only. Keep out of eyes. Stop use and see a doctor is rash or irritation develops and lasts. Keep out of reach of children.

# **OTC - ACTIVE INGREDIENT**

Active Ingredients: Titanium Dioxide 7.0% w/w, Octinoxate 3.6% w/w.

#### INACTIVE INGREDIENT

Other Ingredients: Talc, Zinc Stearate, Nylon-12, Dimethicone, Ethylhexyl Palmitate, Pentaerythrityl Tetraethylhexanoate, Methicone, Aluminum Hydroxide, Butylparaben, Ethylparaben, Isobutylparaben, Methylparaben, Phenoxyethanol, Propylparaben, Iron Oxides, Mica, Titanium Dioxide.

# **DOSAGE & ADMINISTRATION**

Smooth on to face.

# OTC - KEEP OUT OF REACH OF CHILDREN

Keep out of reach of children.

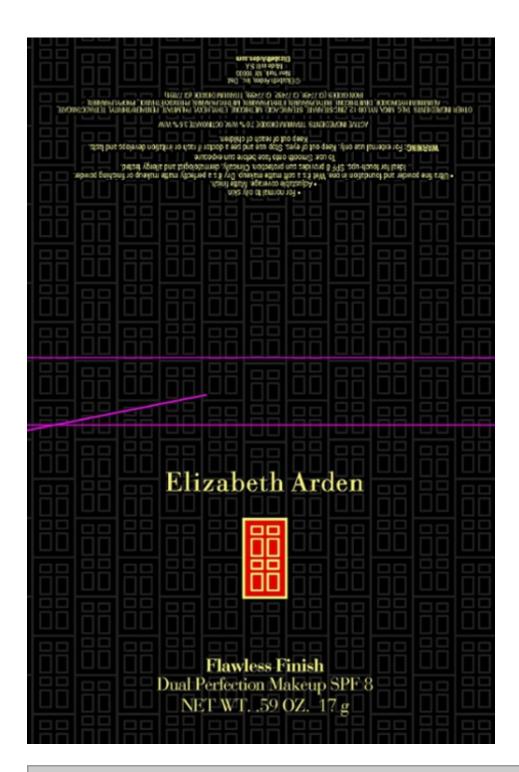
# **OTC - PURPOSE**

SPF 8 provides sun protection.

#### **OTC - WHEN USING**

Keep out of eyes.

#### PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



# FLAWLESS FINISH DUAL PERFECTION MAKEUP SPF 8 VANILLA

titanium dioxide powder

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67938-1157
Route of Administration	TOPICAL		
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Active Ingredient/Active Molety		
Ingredient Name	Basis of Strength	Strength
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP) (TITANIUM DIO XIDE - UNII:15FIX9 V2JP)	TITANIUM DIO XIDE	1.19 g in 17 g

Inactive Ingredients			
Ingredient Name	Strength		
TALC (UNII: 7SEV7J4R1U)			
MICA (UNII: V8A1AW0880)			
OCTINO XATE (UNII: 4Y5P7MUD51)			
ZINC STEARATE (UNII: H92E6QA4FV)			
STEARIC ACID (UNII: 4ELV7Z65AP)			
DIMETHICO NE (UNII: 92RU3N3Y1O)			
ETHYLHEXYL PALMITATE (UNII: 2865993309)			
ALUMINUM HYDRO XIDE (UNII: 5QB0T2IUN0)			
PHENOXYETHANOL (UNII: HIE492ZZ3T)			
METHYLPARABEN (UNII: A218 C7H19 T)			
BUTYLPARABEN (UNII: 3QPI1U3FV8)			
ETHYLPARABEN (UNII: 14255EXE39)			
PROPYLPARABEN (UNII: Z8IX2SC1OH)			

Product Characteristics			
Color	WHITE (Vanilla)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67938-1157-1	1 in 1 BOX		
1	NDC:67938-1157-2	17 g in 1 CONTAINER		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part352	05/24/2006		

# Labeler - Elizabeth Arden, Inc (849222187)

Revised: 1/2011 Elizabeth Arden, Inc