

CLEAR PATH TINTED FACIAL SUNSCREEN- titanium dioxide, zinc oxide cream
LULUS SPA BOUTIQUE

DRUG FACTS (CONT)
• SUN PROTECTION MEASURES: SPENDING TIME IN THE SUN INCREASES YOUR RISK OF SKIN CANCER 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 AM-2 PM
• WEAR LONG-SLEEVED SHIRTS, PANTS, HATS, AND SUNGLASSES

INACTIVE INGREDIENTS: ALUMINA, CYCLOHEXASILOXANE, CYCLOPENTASILOXANE, DIMETHICONE, DIMETHICONE CROSSPOLYMER, DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER, DIMETHICONOL, IRON OXIDE (CI 44791), LAURYL PEG/PPG-18/18 METHICONE, HYDROGEN DIMETHICONE, PEG-10 DIMETHICONE, TETRAHEXYLDECYL ASCORBATE, TOCOPHERYL ACETATE

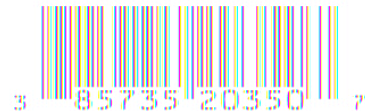
OTHER INFORMATION:
• PROTECT THIS PRODUCT FROM EXCESSIVE HEAT AND DIRECT SUN
• MAY STAIN SOME FABRICS
• PA+++



TINTED FACIAL SUNSCREEN

SPF 40 BROAD SPECTRUM
NON-COMEDOGENIC
WATER RESISTANT (80 MINUTES)

ABQ ACNE CLINIC | ALBUQUERQUE, NM



NET WT. 1.8 oz/ 53 g.

DRUG
ACTIVE
TITANIUM
ZINC
USE
DIRECTIONS
MEASUREMENTS
THE
AGING
WATER
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Table with 4 main sections: Product Information, Active Ingredient/Active Moiety, and Inactive Ingredients. Includes details on product type, route of administration, and specific ingredients like Zinc Oxide and Titanium Dioxide.

Ingredient Name	Strength
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE) (UNII: 9E4CO0W6C5)	
DIMETHICONOL (2000 CST) (UNII: T74O12AN6Y)	
CI 77491 (UNII: 1K09F3G675)	
CI 77492 (UNII: EX438O2MRT)	
PEG-10 DIMETHICONE (600 CST) (UNII: 8PR7V1SVM0)	
TETRAHEXYLDECYL ASCORBATE (UNII: 9LBV3F07AZ)	
ALUMINA (UNII: LMI26O6933)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
LAURYL PEG/PPG-18/18 METHICONE (UNII: ZJ5S27D9NX)	
CI 77499 (UNII: XM0M87F357)	
HYDROGEN DIMETHICONE (20 CST) (UNII: 12Z59IF64N)	
CYCLOHEXASILOXANE (UNII: XHK3U310BA)	
CYCLOPENTASILOXANE (UNII: 0THT5PCI0R)	
DIMETHICONE CROSSPOLYMER (UNII: UF7620L1W6)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85735-203-50	53 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/24/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	04/24/2018	

Labeler - LULUS SPA BOUTIQUE (043019360)

Establishment

Name	Address	ID/FEI	Business Operations
Custom Analytics LLC		144949372	analysis(85735-203)

Establishment

Name	Address	ID/FEI	Business Operations
Fragrance Manufacturing Inc		793406000	manufacture(85735-203)

Establishment

Name	Address	ID/FEI	Business Operations
Fallien Cosmeceuticals Ltd.		958388357	relabel(85735-203)