

NO MAKEUP CONCEALER MEDIUM BROAD SPECTRUM SPF 35- zinc oxide and titanium dioxide cream

Allure Labs, 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:

Zinc Oxide - 5.98%

Titanium Dioxide - 4.80%

Sunscreen

Helps prevent sunburn.

if used as directed with other sun protection measures (see directions), decreases the chance of skin cancer and early skin aging caused by the sun.

For external use only.

Do not use:

on damaged or broken skin.

Stop use and ask a doctor if rash occurs.

keep out of eyes. rinse with water to remove.

if swallowed, get medical help or contact a Poison Control Center right away.

Directions: Apply along the lower eye contours. gently blend with fingertips and smooth outward. Any excess can also be applied on targeted areas of the face to cover imperfections.

For sunscreen use: apply liberally 15 minutes before sun exposure. use a water resistant sunscreen if swimming or sweating. reapply at least every 2 hours. not for use on children.

Sun protection measures: Spending time in the sun increases your risk of skin cancer and early skin aging. to decrease the 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.

Inactive Ingredients: Cyclopentasiloxane, Water/Aqua/Eau, Dimethicone/PEG-10/15 Crosspolymer, Cyclotetrasiloxane, Dimethicone, Sodium PCA, Polyglyceryl-3 polydimethylsiloxylethyl Dimethicone, glycerin, PPG-12/SMDI Copolymer, Nylon-12, Mica (CI 77019), C12-15 Alkyl Benzoate, Stearic Acid, Tocopheryl Acetate, Glucosyl Hesperidin, Linoleamidopropyl PG-Dimonium Chloride Phosphate, tetrahexyldecyl Ascorbate, Dimethyl MEA, Palmitoyl Hexapeptide, Palmitoyl Tripeptide-1, Palmitoyl

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62742-4106
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	59.8 mg in 1 g
TITANIUM DIOXIDE (UNII: 15FIX9V2JP) (TITANIUM DIOXIDE - UNII:15FIX9V2JP)	TITANIUM DIOXIDE	48.0 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
WATER (UNII: 059QF0K00R)	
CYCLOMETHICONE 4 (UNII: CZ227117JE)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	
POLYGLYCERYL-3 POLYDIMETHYLSILOXYETHYL DIMETHICONE (4000 MP.A.S) (UNII: RLA2U05Z4Q)	
GLYCERIN (UNII: PDC6A3C0OX)	
PPG-12/SMDI COPOLYMER (UNII: 1BK9DDD24E)	
NYLON-12 (UNII: 446U8J075B)	
MICA (UNII: V8A1AW0880)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
GLUCOSYL HESPERIDIN (UNII: 432C95B6YE)	
LINOLEAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: 5Q87K461JO)	
TETRAHEXYLDECYL ASCORBATE (UNII: 9LBV3F07AZ)	
DEANOL (UNII: 2N6K9DRA24)	
PALMITOYL HEXAPEPTIDE-12 (UNII: HO4ZT5S86C)	
PALMITOYL LYSYLDIOXYMETHIONYLLYSINE (UNII: T7A529FB8O)	
PALMITOYL TETRAPEPTIDE-7 (UNII: Q41S464P1R)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
CERAMIDE 2 (UNII: C04977SRJ5)	
N-HYDROXYSUCCINIMIDE (UNII: MJE3791M4T)	
HYALURONIC ACID (UNII: S270N0TRQY)	
CHRYSIN (UNII: 3CN01F5ZJ5)	
POLYGLYCERYL-4 ISOSTEARATE (UNII: 820DPX33S7)	
CETYL PEG/PPG-10/1 DIMETHICONE (HLB 1.5) (UNII: V2W71V8T0X)	
PEG/PPG-18/18 DIMETHICONE (UNII: 9H0AO7T794)	
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE) (UNII: 9E4CO0W6C5)	
HEXYL LAURATE (UNII: 4CG9F9W01Q)	
TRIBEHENIN (UNII: 8OC9U7TQZ0)	
CETEARYL OLIVATE (UNII: 58B69Q84JO)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
SORBITAN OLIVATE (UNII: MDL271E3GR)	

HEXYLENE GLYCOL (UNII: KEH0A3F75J)
STEARETH-20 (UNII: L0Q8IK9E08)
PEG-10 RAPESEED STEROL (UNII: 258O76T85M)
TRIETHOXYCAPRYLSILANE (UNII: LDC331P08E)
SILICA DIMETHYL SILYLATE (UNII: EU2PSP0G0W)
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)
HYDROXYPROPYL .ALPHA.-CYCLODEXTRIN (UNII: ZFR0T80O4Y)
Phenoxyethanol (UNII: HIE492ZZ3T)
ALUMINUM HYDROXIDE (UNII: 5QB0T2IUN0)
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)
FERRIC OXIDE RED (UNII: 1K09F3G675)
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)
FERROSFERRIC OXIDE (UNII: XM0M87F357)
PALMITOYL TRIPEPTIDE-1 (UNII: RV743D216M)

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62742-4106-2	1 in 1 CARTON	12/01/2017	
1	NDC:62742-4106-1	9 g in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part352	12/01/2017	

Labeler - Allure Labs, Inc (926831603)