SHEERZINC SPF 30 TINTED - TAN- zinc oxide 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 Ingredient: Zinc oxide - 18.6%

Purpose: Sunscreen

Uses:

- To help protect the skin from harmful UVA and UVB rays

Warnings: For external use only

When using this product:

- Keep out of eyes. If contact occurs, rinse with water.
- Discontinue use if irritation or redness occurs.

Stop use and ask a doctor

- If severe skin irritation developes

Keep out of reach of children:

- If swallowed, get medical help or contact a Poison Cpntrol center right away.

Directions:

- Apply sunscreen in the AM to finger tips and gently massage into the skin. Reapply as needed.

Other Ingredients: Water (Aqua), Cyclopentasiloxane, Butylene glycol, glycerin, Caprylic/Capric triglyceride, Glycerin Stearate, PEG-100 Stearate, Polyglyceryl-3 Polymethylsiloxyeethyl Dimethicon, Cyclohexasiloxane, Sorbitan Stearate, Dimethicone, Polyacrylamide, C13-14 Isoparaffin, Laureth-7,phenoxyethanol, Caprylyl Glycol, Ethylhexyglycerin, Hexylene glycol, Imperata Cylindrica Root Extract, PEG-8, Carbomer, triethoxysilylethyl Polydimethylsiloxyethyl Hexyl Dimethicone, Cetyl Alcohol, Lecithin, Tocopherol, Ascorbyl Palmitate, Ascorbic Acid, Citric Acid, Xanthan gum, Polyester-7, Neopentyl Glycol Diheptanoate, tocopherol Linoleate/Oleate, Teprenone, Phoenix Dactylifera (Date) Fruit Extract, Polygonum Aviculare Extract, Sodium Lactate, Disodium EDTA, Dipotassium Glycyrrhizate, Arabidopsis Thaliana Extract, Plankton Extract, Micrococcus Lysate, Triethoxycaprylysilane, Iron Oxides (CI77491, CI77492, CI77499)

Manufactured for DermaQuest, Inc.

Hayward, CA 94544

1272 GK, NL Made in USA

dermaquestinc.com

SHEERZINC SPF 30 TINTED - TAN

zinc oxide cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62742-4095
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	186 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
DIMETHICONE CROSSPOLYMER (450000 MPA.S AT 12% IN CYCLOPENTASILOXANE) (UNII: UF7620L1W6)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
GLYCERIN (UNII: PDC6A3C0OX)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
PEG-100 STEARATE (UNII: YD01N1999R)	
POLYGLYCERYL-3 POLYDIMETHYLSILOXYETHYL DIMETHICONE (4000 MPA.S) (UNII: RLA2U05Z4Q)	
CYCLOMETHICONE 6 (UNII: XHK3U310BA)	
SORBITAN MONOSTEARATE (UNII: NVZ4I0H58X)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
LAURETH-7 (UNII: Z95S6G8201)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
IMPERATA CYLINDRICA ROOT (UNII: VYT2JA85NH)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
CARBOMER 1342 (UNII: 809Y72KV36)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
TOCOPHEROL (UNII: R0ZB2556P8)	
ASCORBYL PALMITATE (UNII: QN83US2B0N)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
XANTHAN GUM (UNII: TTV12P4NEE)	
POLYESTER-7 (UNII: 0841698D2F)	
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKW3C543X)	
.ALPHA.-TOCOPHEROL LINOLEATE, D- (UNII: G0N132Q0ED)	
TEPRENONE (UNII: S8S8451A4O)	
DATE (UNII: H3O7QI5HY7)	
POLYGONUM AVICULARE TOP (UNII: ZCD6009IUF)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
GLYCYRRHIZINATE DIPOTASSIUM (UNII: CA2Y0FE3FX)	
ARABIDOPSIS THALIANA (UNII: AI3L60HQ81)	
MICROCOCCUS LUTEUS (UNII: LV6L29Z6AX)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	

FERROSOFERRIC OXIDE (UNII: XM0M87F357)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62742-4095-2	1 in 1 CARTON	12/01/2017	
1	NDC:62742-4095-1	56.7 g in 1 TUBE; 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	11/28/2017	

Labeler - Allure Labs Inc (926831603)

Registrant - Allure Labs Inc (926831603)

Establishment

Name	Address	ID/FEI	Business Operations
Allure Labs Inc		926831603	manufacture(62742-4095)

Revised: 11/2017 Allure Labs Inc