TROPICAL TOPICAL MATTE SPF 30- 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

Use: Apply to your cleansed and toned face and neck every morning as a daily moisturizer. Re-apply every 2 hours when exposed to sun or as needed.

How to use: Apply to your cleansed and toned face and neck every morning as a daily moisturizer. Re-apply every 2 hours when exposed to sun or as needed.

Other Ingredients: Water, Cyclopentasiloxane, Butylene Glycol, Glycerin, Cyclohexasiloxane, Glyceryl Stearate, PEG-100 Stearate, Polyglyceryl-3 Polydimethylsiloxyethyl, Dimethicone, Sorbitan Stearate, Cetyl Alcohol, Teprenone, Triethoxysilylethyl Polydimethylsiloxyethyl Hexyl Dimethicone, Caprylic/Capric Triglyceride, Imperata Cylendrica Root Extract, Arabidopsis Thaliana Extract, Plankton Extract, Micrococcus Lysate, Phoenix Dactylifera (Date) Fruit Extract, Polygonum Aviculare Extract, Sodium Lactate, Lecithin, Dipotassium Glycyrrhizate, Tocopheryl Linoleate/ Oleate, Citric Acid, Polyacrylamide, Phenoxyethanol, Caprylyl Glycol, Ethylhexylglycerine, Hexylene Glycol, Xanthan Gum, C13-14 Isoparaffin, Laureth-7, PEG-8, Carbomer, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Tocopherol, Asccorbyl Palmitate, Ascorbc Acid, Fragrance, Disodium EDTA.

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Sorella Apothecary, Reno, NV 89509.

SORELLA

APOTHECARY

DRUG FACTS

Active Ingredient:	Purpose:
Zinc Oxide 18.6 %	Sunscreen

Uses:

• Helps prevent sunburn. • If used as directed with other sun protection measures, this product decreases the risk of skin cancer and early skin aging caused by the sun.

Warnings:

• For external use only. • Do not use on damaged or broken skin. • When using this product keep out of eyes. Rinse with water to remove. • Stop use and ask a doctor if rash occurs. • Keep out of reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions:

• Apply liberally 15 minutes before sun exposure. • Children under 6 months of age: Ask a doctor. • Reapply at least every 2 hours.

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 a.m.-2 p.m.
• Wear long-sleeved shirts, pants, hats and sunglasses.

Other Information:

Protect the product in this container from excessive heat and direct sun.

Inactive Ingredients:

Water, Cyclopentasiloxane, Butylene Glycol, Glycerin, Cyclohexasiloxane, Glyceryl Stearate, PEG-100 Stearate, Polyglyceryl-3 Polydimethylsiloxyethyl Dimethicone, Sorbitan Stearate, Dimethicone, Cetyl Alcohol, Tapioca Starch, Triethoxycaprylylsilane, Polydimethylsiloxyethyl Hexyl Dimethicone, Caprylic/Capric Triglyceride, Imperata Cylindrica Root Extract, Arctostaphylos Uva-Ursi Leaf Extract, Plankton Extract, Micrococcus Lysate, Phoenix Dactylifera (Date) Fruit Extract, Polygonum Aviculare Extract, Sodium Lactate, Lecithin, Dipotassium Glycyrrhizate, Tocopheryl Linoleate/Oleate, Citric Acid, Polyacrylamide, Phenoxyethanol, Caprylyl Glycol, Ethylhexylglycerin, Hexylene Glycol, Xanthan Gum, C13-14 Isoparaffin, Laureth-7, PEG-8, Carbomer, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Tocopherol, Ascorbyl Palmitate, Ascorbic Acid, Fragrance, Disodium EDTA

VEGAN
GLUTEN FREE
cruelty free

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HOW TO USE

Apply to your cleansed and toned face and neck every morning as a daily moisturizer. Re-apply every 2 hours when exposed to sun or as needed.

Tropical Topical

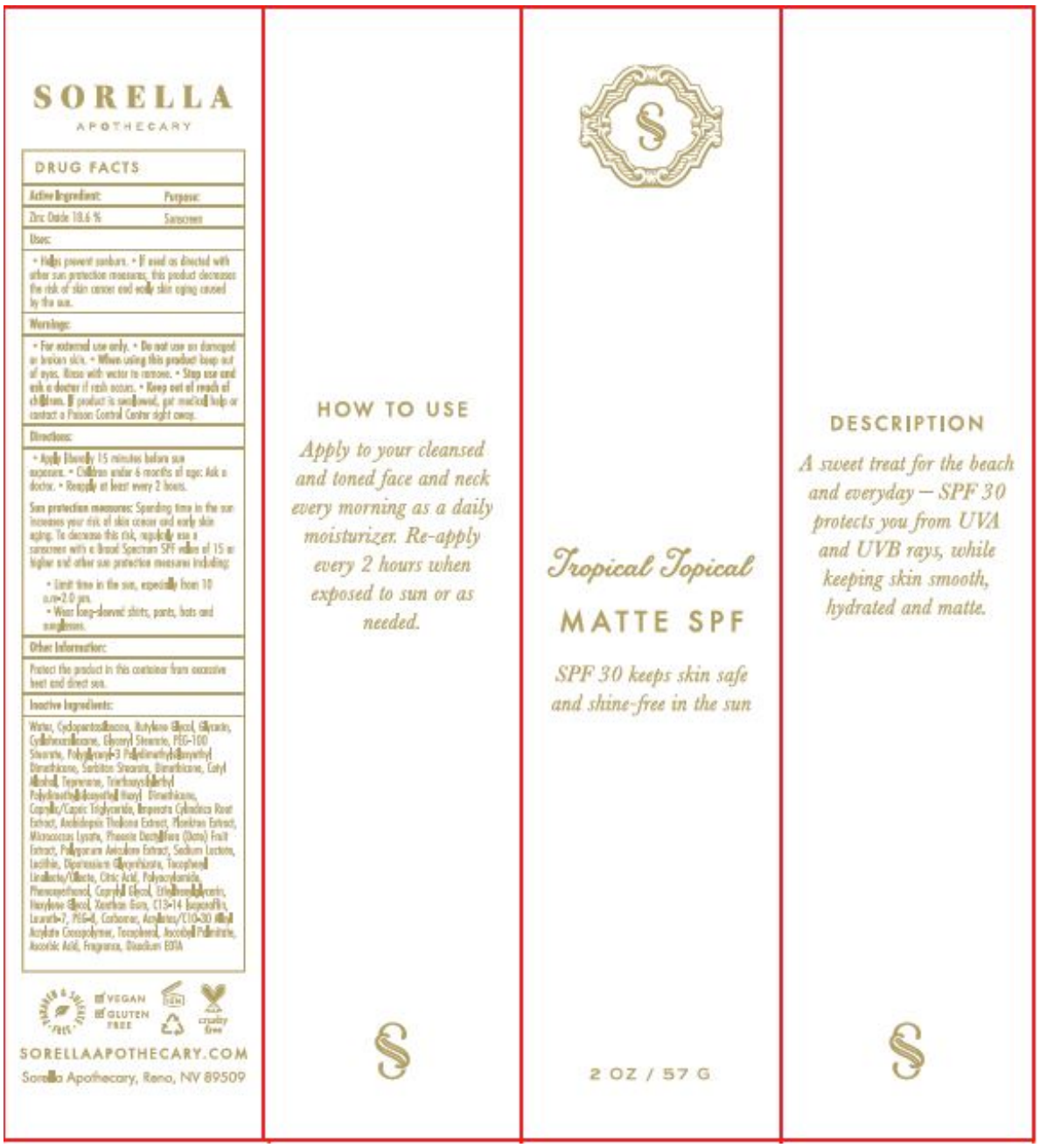
MATTE SPF

SPF 30 keeps skin safe and shine-free in the sun

2 OZ / 57 G

DESCRIPTION

A sweet treat for the beach and everyday — SPF 30 protects you from UVA and UVB rays, while keeping skin smooth, hydrated and matte.



TROPICAL TOPICAL MATTE SPF 30

zinc oxide cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62742-4157
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	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)	
TRIEPOXYCYCLOHEXASILANE (UNII: 066Q83563R)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
PEG-100 STEARATE (UNII: YD01N1999R)	
POLYGLYCERYL-3 POLYDIMETHYLSILOXYETHYL DIMETHICONE (4000 MPA.S) (UNII: RLA2U05Z4Q)	
SORBITAN MONOSTEARATE (UNII: NVZ4I0H58X)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
TEPRENONE (UNII: S8S8451A4O)	
CAPRYLIC/CAPRIC/LAURIC TRIGLYCERIDE (UNII: FJ1H6M2JG9)	
IMPERATA CYLINDRICA ROOT (UNII: VYT2JA85NH)	
ARABIDOPSIS THALIANA (UNII: AI3L60HQ81)	
MICROCOCCUS LUTEUS (UNII: LV6L29Z6AX)	
PHOENIX DACTYLIFERA WHOLE (UNII: 8QI9RWU9M1)	
POLYGONUM AVICULARE WHOLE (UNII: M990N03611)	
SODIUM LACTATE (UNII: TU7HW0W0QT)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
TOCOPHEROL (UNII: R0ZB2556P8)	
CITRIC ACID ACETATE (UNII: DSO12WL7AU)	
POLYACRYLAMIDE (10000 MW) (UNII: E2KR9C9V2I)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
XANTHAN GUM (UNII: TTV12P4NEE)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
LAURETH-7 (UNII: Z95S6G8201)	
PEG-8 DIMETHICONE (UNII: GIA7T764OD)	
CARBOMER 934 (UNII: Z135WT9208)	
(C10-C30)ALKYL METHACRYLATE ESTER (UNII: XH2FQZ38D8)	
ASCORBYL PALMITATE (UNII: QN83US2B0N)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62742-4157-2	1 in 1 CARTON	12/12/2017	
1	NDC:62742-4157-1	57 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	12/12/2017	

Labeler - Allure Labs Inc (926831603)

Revised: 12/2017 Allure Labs Inc