NUGLOW DAYTIME THERAPY WITH SPF 15- avobenzone, octinoxate, and zinc oxide cream
G.S. COSMECEUTICAL USA, 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.

NuGLOW®

DIRECTIONS

After cleansing, apply liberally to face daily. Apply evenly before sun exposure. Reapply as needed after towel drying, swimming or perspiring.

Children under 6 months: ask a doctor.

WARNINGS

For external use only.

When using this product keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor if rash or irritation develops and lasts.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

QUESTIONS?

1-866-605-9510 or visit www.nuglowskincare.com.

Daytime Therapy With SPF15

ACTIVE INGREDIENTS

Avobenzone 1.0%, Octinoxate 7.0%, Zinc Oxide 4.0%.

INACTIVE INGREDIENTS

Aqua (Water), Caprylic/Capric Triglyceride, Glyceryl Stearate SE, Butyrospermum Parkii (Shea Butter), Polysorbate 60, Cetyl Alcohol, Dimethicone, Cetyl Ricinoleate, Stearyl Alcohol, Persea Gratissima (Avocado) Oil, Glycerin, Sodium Hyaluronate, Camellia Oleifera (Green Tea) Leaf Extract, Chrondrus Crispus (Carrageenan), Sodium PCA, Xanthan Gum, Tocopheryl Acetate, Ascorbic Acid, Lactic Acid, Glucose, Triethoxycaprylylsilane, Phenoxyethanol, Caprylyl Glycol, Ethylhexylglycerin, Hexylene Glycol.

For expiration date, see bottom of jar.

Made in the U.S.A. for NuGlow, Los Angeles, CA 90049

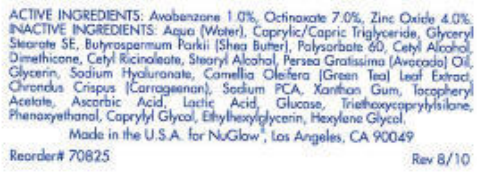
Reorder# 70825
Rev 8/10

PRINCIPAL DISPLAY PANEL - 34g Jar Label

NuGlow®

Daytime Therapy
With SPF15

Net Weight 1.2 Oz./34g e



NUGLOW DAYTIME THERAPY WITH SPF 15

avobenzone, octinoxate, and zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65113-8503
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	1 g in 100 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7 g in 100 g
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	4 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
AVOCADO OIL (UNII: 6VNO72PFC1)	
GLYCERIN (UNII: PDC6A3C0OX)	
CAMELLIA OLEIFERA LEAF (UNII: 5077EL0C60)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
LACTIC ACID (UNII: 33X04XA5AT)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	

Product Characteristics			
Color	WHITE	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65113-8503-3	34 g in 1 JAR		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part352	10/15/2010	

Labeler - G.S. COSMECEUTICAL USA, INC. (017014734)

Establishment

Name	Address	ID/FEI	Business Operations
G.S. COSMECEUTICAL USA, INC.		017014734	MANUFACTURE

Revised: 2/2011 G.S. COSMECEUTICAL USA, INC.