BLING DAY EVERYDAY SUNSCREEN BROAD SPECTRUM SPF 50- ensulizole, octinoxate, octocrylene, titanium dioxide, zinc oxide cream Dong Sung Bio Pharm. Co., Ltd.

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.

42361-155 Drug Facts

Drug Facts

Ensulizole 3.9%

Octinoxate 7.4%

Octocrylene 3.0%

Titanium Dioxide 4.9%

Zinc Oxide 4.9%

Drug Facts

Sunscreen

Uses

Not only will it help prevent sunburn, it also reduces the risk of skin cancer and early skin aging if used as directed with other sun protection measures. (SPF value of 15 or higher)

Warnings

For external use only. Do not use on damaged or broken skin. Discontinue use if signs of irritation or rash appear. Avoid contact with eyes. Wash off immediately with cold water if product comes in contact with eyes. **Keep this and all drugs out of reach of children.** In case of accidental ingestion, seek professional assistance or contact a Poison Control Center right away. **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. and wear long-sleeved shirts, pants, hats and sunglasses. For use on skin only.

Warnings

Keep this and all drugs out of reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center right away.

Directions

At the end of basic skin care regimen, apply liberally on face to cover the entire facial area and lightly go over parts where extra cover-up is needed with detailed perfection. Apply 15 minutes before sun exposure. For added protection, reapply at least every 2 hours, after swimming, towel drying or extended sun exposure. Use a water resistant sunscreen if swimming or sweating. **Children under 6 months:** Ask a doctor.

Other Information

Protect this product from excessive heat and direct sun.

Inactive Ingredients

Water, Cocoglycerides, Polyglyceryl-3 Diisostearate, Pentylene Glycol, Glycerin, Cetearyl Alcohol, Triethanolamine, Dimethicone, Butylene Glycol, Pentaerythrityl Distearate, Disodium Cetearyl Sulfosuccinate, Beheneth-25, Aluminum Hydroxide, Stearic Acid, Dimethicone/PEG-10/15 Crosspolymer, Phenoxyethanol, Magnesium Aluminum Silicate, Fragrance, Xanthan Gum, Triethoxycaprylsilane

Everyday Sunscreen

Broad Spectrum SPF 50+

UVA/UVB Protection



BLING DAY EVERYDAY SUNSCREEN BROAD SPECTRUM SPF 50

ensulizole, octinoxate, octocrylene, titanium dioxide, zinc oxide cream

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:42361-155	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ENSULIZOLE (UNII: 9 YQ9 DI1W42) (ENSULIZOLE - UNII:9 YQ9 DI1W42)	ENSULIZOLE	39 mg in 1 g		
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP) (TITANIUM DIO XIDE - UNII:15FIX9 V2JP)	TITANIUM DIO XIDE	49 mg in 1 g		
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	49 mg in 1 g		
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	30 mg in 1 g		
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	74 mg in 1 g		

Inactive Ingredients	
Ingredient Name	Strength
POLYGLYCERYL-3 DIISOSTEARATE (UNII: 46 P231IQV8)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
TROLAMINE (UNII: 9O3K93S3TK)	
COCO-GLYCERIDES (UNII: ISE917DNUG)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALUMINUM HYDRO XIDE (UNII: 5QB0T2IUN0)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
BEHENETH-25 (UNII: 0G17KJ5M7P)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6 M3P6 4 V0 NC)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
XANTHAN GUM (UNII: TTV12P4NEE)	
TRIETHO XYCAPRYLYLSILANE (UNII: LDC331P08E)	
WATER (UNII: 059QF0KO0R)	
DIMETHICO NE (UNII: 92RU3N3Y10)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
PENTAERYTHRITYL DISTEARATE (UNII: 697WOT8HNB)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:42361-155-01	1 in 1 CARTON	12/0 1/20 18	
1	50 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 not final	part352	12/0 1/20 18	

Labeler - Dong Sung Bio Pharm. Co., Ltd. (687811661)

Registrant - Dong Sung World Wide USA, Inc. (784969219)

Establishment				
Name	Address	ID/FEI	Business Operations	
Dong Sung Bio Pharm. Co., Ltd.		687811661	label(42361-155)	

Establishment			
Name	Address	ID/FEI	Business Operations
Omar Sharif Cosmetic Co., Ltd.		689316318	manufacture(42361-155)

Revised: 10/2019 Dong Sung Bio Pharm. Co., Ltd.