ALBA BOTANICA SENSITIVE MINERAL SPF 50- zinc oxide lotion The Hain Celestial Group, Inc.

AL00656 Alba Botanica Sensitive Mineral SPF 50 Formula PF163046-09

Zinc Oxide 24.1% (w/w)

Apply liberally 15 minutes before sun exposure. Reapply: \bullet after 80 minutes of swimming or sweating \bullet immediately after towel drying \bullet at least every 2 hours. Children under 6 months of age: ask a doctor.

Sunscreen

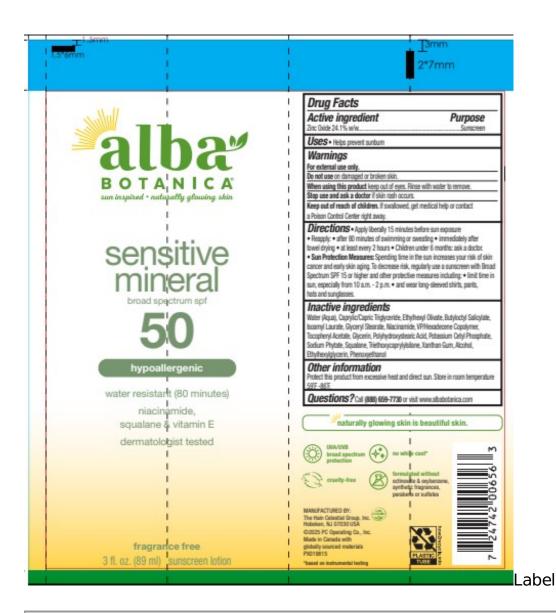
For external use only. Do not use on damaged or broken skin. When using this product keep out of eyes. Rinse with water to remove. Stop use and ask a doctor if skin rash occurs.

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

Water (Aqua), Caprylic/Capric Triglyceride, Ethylhexyl Olivate, Butyloctyl Salicylate, Isoamyl Laurate, Glyceryl Stearate, Niacinamide, VP/Hexadecene Copolymer, Tocopheryl Acetate, Glycerin, Polyhydroxystearic Acid, Potassium Cetyl Phosphate, Sodium Phytate, Squalane, Triethoxycaprylylsilane, Xanthan Gum, Alcohol, Ethylhexylglycerin, Phenoxyethanol

Helps prevents sunburns. If used as directed with other sun protection measures, decreases risk of skin cancer and early skin aging caused by sun exposure. **Skin Protection Measures**: Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease risk, regularly use sunscreen with Broad Spectrum SPF 15 or higher and other protective measures including: limit time in sun, especially from10am to 2pm, and wear long sleeved shirts, pants, hats and sunglases.

Alba Botanical Sensitive Mineral sunscreen SPF 50



ALBA BOTANICA SENSITIVE MINERAL SPF 50

zinc oxide lotion

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:61995-2036

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Basis of Strength	Strength
	Basis of Strength

ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z) ZINC OXIDE 24.1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ETHYLHEXYL OLIVATE (UNII: HTC7G3S2PV)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	

VINYLPYRROLIDONE/HEXADECENE COPOLYMER (UNII: KFR5QEN0N9)

GLYCERYL STEARATE (UNII: 2300U9XXE4)	
ISOAMYL LAURATE (UNII: M1SLX00M3M)	
NIACINAMIDE (UNII: 25X51I8RD4)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
SQUALANE (UNII: GW89575KF9)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT)	
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
GLYCERIN (UNII: PDC6A3C0OX)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ALCOHOL (UNII: 3K9958V90M)	
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	
SODIUM PHYTATE (UNII: 88496G1ERL)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:61995- 2036-1	59 g in 1 TUBE; Type 0: Not a Combination Product	11/26/2025		
2	NDC:61995- 2036-2	89 g in 1 TUBE; Type 0: Not a Combination Product	11/26/2025		
3	NDC:61995- 2036-3	113 g in 1 TUBE; Type 0: Not a Combination Product	11/26/2025		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M020	11/26/2025		

Labeler - The Hain Celestial Group, Inc. (117115556)

Registrant - The Hain Celestial Group, Inc. (081512382)

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