

ALBA BOTANICA SENSITIVE MINERAL SPF 30- zinc oxide lotion
The Hain Celestial Group, Inc.

AL00441 Alba Botanica Sensitive Mineral SPF 30
Formula PF163046-09

Zinc Oxide 24.1% (w/w)

Apply liberally 15 minutes before sun exposure. Reapply: ● after 80 minutes of swimming or sweating ● immediately after towel drying ● 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 Olivatate, 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 prevent 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 from 10am to 2pm, and wear long sleeved shirts, pants, hats and sunglasses.

Alba Botanical Sensitive Mineral sunscreen SPF 50



ALBA BOTANICA SENSITIVE MINERAL SPF 30

zinc oxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:61995-2034
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	24.1 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
VINYLPYRROLIDONE/HEXADECENE COPOLYMER (UNII: KFR5QEN0N9)	
GLYCERYL STEARATE (UNII: 230OU9XXE4)	
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)
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)
XANTHAN GUM (UNII: TTV12P4NEE)
ETHYLHEXYL OLIVATE (UNII: HTC7G3S2PV)
ALCOHOL (UNII: 3K9958V90M)
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)
SODIUM PHYTATE (UNII: 88496G1ERL)
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61995-2034-1	59 g in 1 TUBE; Type 0: Not a Combination Product	11/26/2025	
2	NDC:61995-2034-2	89 g in 1 TUBE; Type 0: Not a Combination Product	11/26/2025	
3	NDC:61995-2034-3	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)

Revised: 11/2025

The Hain Celestial Group, Inc.