

**CEILE SPF 30 PRIME PROTECT BROAD SPECTRUM SPF 30 SUNSCREEN-
sunscreen 30 mineral cream
LÚE COSMETICS INC**

DRUG FACTS

Active Ingredients

Zinc Oxide 10.00%

Uses

- helps prevent sunburn
- if used as directed with other sun protection measures (see Directions), 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 swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Shake Well •apply generously 15 minutes before sun exposure and as needed
- reapply at least every 2 hours.
- use a water resistant sunscreen if swimming or sweating
- 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
- children under 6 months of age: Ask a doctor

Inactive ingredients

Water (Aqua), Caprylic/Capric Triglyceride, Butylene Glycol, Polyhydroxystearic Acid, Hydroxyethyl Acrylate/Sodium Acryloyldimethyl Taurate Copolymer, Tocopheryl Acetate, Xylitol, Xylitylglucoside, Anhydroxylitol, Sodium Olivatate, Neopentyl Glycol Diethylhexanoate, Dimethicone, Triethoxycaprylsilane, Citric Acid, Phenoxyethanol, Ethylhexylglycerin, Iron Oxides.

Other Information

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

Questions?

1-866-860-0811

Purpose

Sunscreen

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Keep out of reach of children

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Principal Display Panel



SUNSCREEN

sunscreen 30 mineral cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83103-2000
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC CATION - UNII:13S1S8SF37)	ZINC CATION	10 mg in 100 mg

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CAPRYLIC/CAPRIC TRIGLYCERIDE (UNII: C9H2L21V7U)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (45000 MPA.S AT 1%) (UNII: 86FQE96TZ4)	
XYLITOL (UNII: VCQ006KQ1E)	
XYLITYLGLUCOSIDE (UNII: O0IEZ166FB)	
ANHYDROXYLITOL (UNII: 8XWR7NN42F)	
SODIUM OLIVATE (UNII: ND5Y5M6ZUT)	
NEOPENTYL GLYCOL DIETHYLHEXANOATE (UNII: U68ZV6W62C)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
TRIETHOXYCAPRYLYLSILANE (UNII: LDC331P08E)	
CITRIC ACID (UNII: 2968PHW8QP)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
BROWN IRON OXIDE (UNII: 1N032N7MFO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83103-2000-1	1 in 1 PACKAGE	01/30/2025	
1		50 mg 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 Drug	M020	01/30/2025	

