

KNESKO SKIN THE SUNSCREEN- zinc oxide lotion
Knesko, LLC

Knesko Skin The Sunscreen

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Drug Facts

Active Ingredient	Purpose
Zinc Oxide 12%	Sunscreen

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 skin rash occurs.
- Keep out of reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Shake well.
- 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.



I HONOR
MY INTUITION

KNESKO
SKIN

THE SUNSCREEN
LA CRÈME SOLAIRE

BROAD SPECTRUM
SPF 50+

WATER RESISTANT
(80 MINUTES)

50 mL / 1.7 FL OZ



PRODUCT#1010

Drug Facts (continued)

Directions (continued)

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 - Limit your time in the sun, especially from 10 a.m. – 2 p.m.
 - Wear long-sleeved shirts, pants, hats, and sunglasses.

Other information

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

Inactive ingredients

Allantoin, Bisabolol, Butyloctyl Salicylate, C12-15 Alkyl Benzoate, C13-15 Alkene, Calcium Sodium Borosilicate, Caprylyl Glycol, Caprylyl Methicone, Dimethicone, Disteardimonium Hectorite, Ethylhexylglycerin, Hexylene Glycol, Isododecane, Lauryl PEG-8, Dimethicone, Lecithin, Niacinamide, Octyldodecyl Neopentanoate, Phenoxyethanol, Polyglyceryl-2 Dipolyhydroxystearate, Polyglyceryl-3 Polyricinoleate, Propanediol, Silica, Sodium Chloride, Sodium Hydroxide, Tetrasodium Glutamate Diacetate, Triethyl Salicylate, Water.

Question or Comments?

Call Toll-free 800-454-3091

DISTRIBUTED BY KNESKO, LLC,
107 N BIRNO RD #401
NEWBURY PARK, CA 91320

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zinc oxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:87407-236
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	132 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
POLYGLYCERYL-2 DIPOLYHYDROXYSTEARATE (UNII: 9229XJ4V12)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
DIMETHICONE 200 (UNII: RGS4T2AS00)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
.ALPHA.-BISABOLOL, (+)- (UNII: 105S6I733Z)	
LAURYL PEG-8 DIMETHICONE (300 CPS) (UNII: ELL2U7K8T8)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)	
ISODODECANE (UNII: A8289P68Y2)	
WATER (UNII: 059QF0KO0R)	
NIACINAMIDE (UNII: 25X51I8RD4)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
PROPANEDIOL (UNII: 5965N8W85T)	
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ALLANTOIN (UNII: 344S277G0Z)	
TRIDECYL SALICYLATE (UNII: AZQ08K38Z1)	
CAPRYLYL METHICONE (UNII: Q95M2P1KJL)	
C13-15 ALKANE (UNII: 114P5I43UJ)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CALCIUM SODIUM BOROSILICATE (UNII: 4MM76N4VMY)	
DISTEARDIMONIUM HECTORITE (UNII: X687XDK09L)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	

Contains**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87407-236-02	1 in 1 CARTON	04/20/2026	
1	NDC:87407-236-01	50 mL in 1 BOTTLE; 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	04/20/2026	

Labeler - Knesko, LLC (071524880)

Revised: 5/2026

Knesko, LLC