EQUALINE ZINC OXIDE SUNSCREEN STICK SPF 50- zinc oxide stick United Natural Foods, Inc. dba UNFI

Equaline Zinc Oxide Sunscreen Stick SPF 50 CD2-449-161-3

Active Ingredients

Zinc Oxide 21.6%

Purpose

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

rash occurs.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- 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
- **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

Other Information

- protect the product in this container from excessive heat and direct sun
- may stain or damage some fabrics, materials or surfaces

∏Inactive Ingredient(s)

beeswax, butyloctyl salicylate, caprylyl/ capric triglyceride, caprylyl glycol, cetyl alcohol, euphorbia cerifera (candelilla) wax, fragrance, isopropyl palmitate, isostearic acid, neopentyl glycol diethylhexanoate, ozokerite, polyhydroxystearic acid, silica

Label





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 + apply liberally and everly 15 minutes before sun exposure + reapply: + after 80 minutes of swimming or swesting + immediately after tread down - at least ways.

Drug Facts (continued)

minutes of swimming or sweating - immediately after towel drying - at least every 2 hours - children under 6 months: Ask a doctor - Sun Protection Measures, Spending time in the sun increases your risk of skin carcer 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

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Drug Facts (continued)

including: • limit 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 - may stain some fabrics

Inactive ingredients beeswax, butyloctyl salicylate, caprylyl/capric triglyceride, caprylyl captorbia cerifera (candellis) wax, fragrance, isopropyl polymitate, isostearic acid, neopentyl glycol diethylhexanoste, ozokerite, polyhydroxystearic acid, silica,

EQUALINE ZINC OXIDE SUNSCREEN STICK SPF 50

zinc oxide stick

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:41163-969

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)

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216 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
POLYHYDROXYSTEARIC ACID (2300 MW) (UNII: YXH47AOU0F)	
CANDELILLA WAX (UNII: WL0328HX19)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
WHITE WAX (UNII: 7G1J5DA97F)	
NEOPENTYL GLYCOL DIETHYLHEXANOATE (UNII: U68ZV6W62C)	
ISOSTEARIC ACID (UNII: X33R8U0062)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
COCOA BUTTER (UNII: 5120YT1CRR)	
CERESIN (UNII: Q1LS2UJO3A)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	

l	P	Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
	1		42 g in 1 CONTAINER; Type 0: Not a Combination Product	09/14/2023			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M020	09/14/2023		

Labeler - United Natural Foods, Inc. dba UNFI (943556183)

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