

ELTA MD UV AOX EYE- zinc oxide cream
CP Skin Health Group, Inc.

Elta MD UV AOX Eye

Directions

Apply liberally 15 minutes before sun exposure

Use a water resistant sunscreen if swimming or sweating

Reapply at least every 2 hours

Warning

For external use only

Do not use on damaged or broken skin

When using this product keep out of eye. Rinse with water to remove.

Stop use and ask a physician if rash occurs

Water, Diheptul Succinate, Butyloctyl Salicylate, Cocos Nucifera (Coconut) Oil, Dimethicone Crosspolymer, Capryloyl Glycerin/Sebacic Acid Copolymer, Glycerin, Dimethicone, Cetearyl Alcohol, Sodium Stearoyl Glutamate, Arachidyl Alcohol, Microcrystalline Cellulose, Phenoxyethanol, Albizia Julibrissin Bark Extract, Cetyl Alcohol, Tocopherol, Coco-Glucoside, Behenyl Alcohol, Butylene Glycol, Octyldodecyl Oleate, Iron Oxides, Arachidyl Glucoside, Bisabolol, Butyrospermum Parkaii (Shea) Butter, Sodium Gluconate, Theobroma Cacao (Cocoa) Seed Butter, Cellulose Gum, Ethylhexylglycerin, Isoceteth-10, Polyhydroxystearic Acid, Silica, Sodium Hyaluronate, Pentylene Glycol, Sodium Benzoate, Carpryl Glycol, Darutoside, Acetyl Tetrapeptide-5, N-Prolyl Palmitoyl Tripeptide-56 Acetate

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

Keep out of reach of children

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

Purpose

Sunscreen

Elta MD UV AOX Eye

elta MD
SKINCARE
UV AOX Eye
BROAD SPECTRUM SPF 30
TINTED



UV AOX Eye

BROAD SPECTRUM SPF 30

SPF **30** TINTED EYE SUNSCREEN

100% Mineral Active (Zinc Oxide)

Protects Skin From Photoaging

Antioxidants & Peptides

Transparent Zinc Oxide Finish

Ophthalmologist Tested

net wt. 0.4 oz (11 g)

Drug Facts (continued)

Gluconate, Theobroma Cacao (Cocoa) Seed Butter, Cellulose Gum, Ethylhexylglycerin, Isoceteth-10, Polyhydroxystearic Acid, Silica, Sodium Hyaluronate, Pentylene Glycol, Sodium Benzoate, Caprylyl Glycol, Darutoside, Acetyl Tetrapeptide-5, N-Prolyl Palmitoyl Tripeptide-56 Acetate

Questions or comments?

Call toll free
1-800-633-8872

20-BIL-55046



Drug Facts

Active ingredient	Purpose
Zinc Oxide 15.0%	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
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When using this product keep out of eyes. Rinse with water to remove.

Stop use and ask a physician if rash occurs

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Directions • apply liberally 15 minutes before sun exposure • use a water-resistant sunscreen if swimming or sweating • reapply at least every 2 hours • **Sun Protection Measures.** Spending time in the sun increases your

Drug Facts (continued)

risk of skin cancer and early skin aging. To decrease this risk, regularly use a sunscreen with a Broad Spectrum SPF 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 this product from excessive heat and direct sun

Inactive ingredients

Water, Diheptyl Succinate, Butyloctyl Salicylate, Cocos Nucifera (Coconut) Oil, Dimethicone/Vinyl Dimethicone Crosspolymer, Capryloyl Glycerin/Sebacic Acid Copolymer, Glycerin, Dimethicone, Cetearyl Alcohol, Sodium Stearoyl Glutamate, Arachidyl Alcohol, Microcrystalline Cellulose, Phenoxyethanol, Albizia Julibrissin Bark Extract, Cetyl Alcohol, Tocopherol, Coco-Glucoside, Behenyl Alcohol, Butylene Glycol, Octyldodecyl Oleate, Iron Oxides, Arachidyl Glucoside, Bisabolol, Butyrospermum Parkii (Shea) Butter, Sodium

ELTA MD UV AOX EYE

zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72043-4342
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	15 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72043-4342-1	1 in 1 CARTON	01/09/2024	
1		11 g 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	01/09/2024	

Labeler - CP Skin Health Group, Inc. (611921669)

Revised: 1/2024

CP Skin Health Group, Inc.