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 resistent 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, Polyhydrocxystearic Acid, Silica, Sodium Hyaluronate, Pentylene, Glycol, Sodium Benzoate, Carprylyl 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 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.