ELTA MD- zinc oxide spray
CP Skin Health Group, Inc.

Elta MD UV Active Spray

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 physician

if rash occurs

Inactive Ingredients

Caprylic/Capric Triglyceride, Helianthys Annuus (Sunflower) Seed Wax, Cerearyl Nonanoate, Theobroma Cacao Cocoa) Seed Butter, Butyloctyl Salicylate, Lauryl Laurate, Polyhydroxystearic Acid, Euphorbia Cerifera (Candelilla) Was, Capryloyl Glycerin/Sebacic Acid Copolymer, Mica, Behenyl Behenate, Butyrospermum Parkii (Shea) Butter, Oryzanol, Ethyl Ferulate, Isostearic Acid, Lecithin, Polyglyceryl-3 Plyricinoleate, Bisabolol, Tocopherol, Silica

Uses

helps prevent sunburn
if used as directed with other sun protection measures (seeDirections), decreases the risk of UV damage and early skin aging caused by the sun

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

Sunscreen

Apply liberally 15 minutes before sun exposure
Reapply: after 80 minutes of swimming or sweating: immediately after towel drying: at lear every 2 hours
Sun Protection Measures

Spending time in the sun can increase your riske of skin cancer and early skin aging. To decrease this risk regularly use a sunscreen with a broad spectrum SPF 15 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-sleeve shirts, pants, hats, and sunglasses
Children under 6 months:Ask a physician

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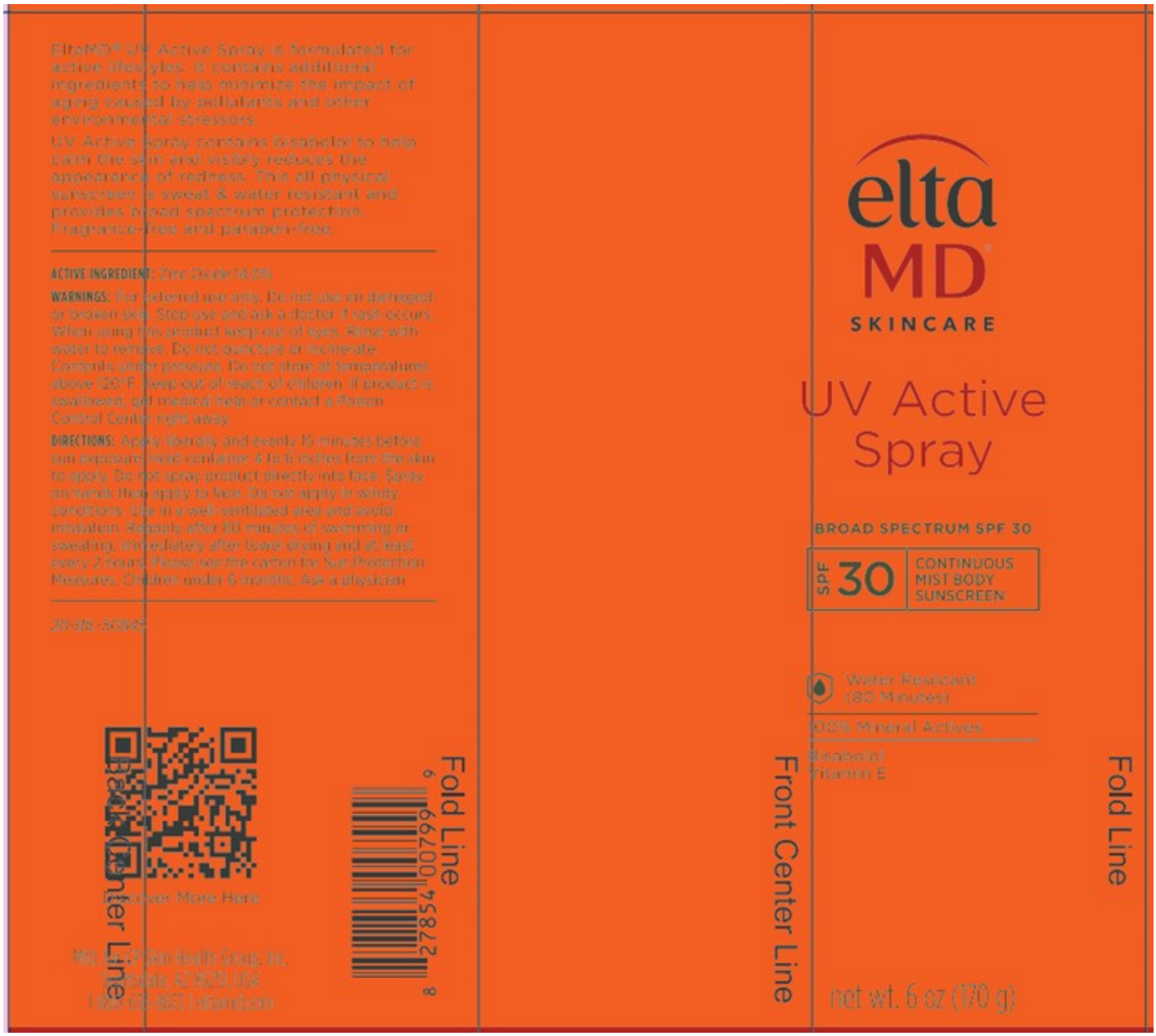

ELTA MD

zinc oxide spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72043-4341
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	14 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:72043-4341-1	1 in 1 CARTON	12/14/2023	
1		170 g in 1 BOTTLE, SPRAY; 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	12/14/2023	

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

Revised: 12/2023 CP Skin Health Group, Inc.