ANEW CLINICAL ABSOLUTE EVEN SPOT CORRECTING HAND- octinoxate, oxybenzone, avobenzone cream New Avon LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Active ingredientsOCTINOXATE 7.5%, OXYBENZONE 4.0%, AVOBENZONE 2.0%......

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
- children under 6 months of age: ask a doctor
- reapply at least every 2 hours
- use a water resistant sunscreen if swimming or sweating
- **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

Inactive ingredients WATER/EAU, GLYCERIN, PENTAERYTHRITYL
TETRAETHYLHEXANOATE, STEARETH-2, GLYCOLIC ACID, BEHENYL ALCOHOL,
DIMETHICONE, PEG-40 STEARATE, MYRISTYL MYRISTATE, TRISILOXANE, LAURYL
LACTATE, ASPARTIC ACID, ZEA MAYS (CORN) KERNEL EXTRACT, PHYTOL,
BUTYROSPERMUM PARKII (SHEA) BUTTER, COLA NITIDA SEED EXTRACT, GLYCINE
SOJA (SOYBEAN) SEED EXTRACT, PUNICA GRANATUM FRUIT JUICE, RETINYL
PALMITATE, DAUCUS CAROTA SATIVA (CARROT) ROOT EXTRACT, ASCORBYL
PALMITATE, HELIANTHUS ANNUUS (SUNFLOWER) SEED EXTRACT, RETINOL, GLYCINE
SOJA (SOYBEAN) OIL, SAXIFRAGA SARMENTOSA EXTRACT, VITIS VINIFERA (GRAPE)
FRUIT EXTRACT, TOCOPHEROL, BETA-CAROTENE, MORUS NIGRA ROOT EXTRACT,
SCUTELLARIA BAICALENSIS ROOT EXTRACT, GLYCERYL STEARATE, PEG-100

STEARATE, PVM/MA DECADIENE CROSSPOLYMER, XANTHAN GUM, SILICA, ACRYLATES/CARBAMATE COPOLYMER, MALTODEXTRIN, LECITHIN, TRIMETHYLSILOXYSILICATE, BUTYLENE GLYCOL, AMMONIUM HYDROXIDE, PHENOXYETHANOL, METHYLPARABEN, DISODIUM EDTA, BHT, PARFUM/FRAGRANCE.

Questions? 1-800-FOR-AVON



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Inactive Ingredients Waterfau, Glycerin, Pentaerythrityl Tetraethylhexanoate, Steareth-2, Glycolic acid, Behenyl Alcohol, Dimethicone, Peg-40 Stearate, Myristyl Myristate, Trisiloxane, Lauryl Lactate, Aspartic acid, Zea Mays (Corn) Kernel Extract, Phytol, Butyrospermum Parkii (Shea) Butter, Cola Nitida Seed Extract, Glycine Soja (Soybean) Seed Extract, Punica Granatum Fruit Juice, Retinyl Palmitate, Daucus Carota Sativa Carrot) Root Extract, Ascorbyl Palmitate, Helianthus Annius (Sunflower) Seed Extract, Tetinol, Glycine Soja (Soybean) Oil, Saxifraga Sarmentosa Extract, VITIS VINIFERA (GRAPE) FRUIT EXTRACT, TOCOPHEROL,

Drug Facts (continued)

BETA-CAROTENE, MORUS NIGRA ROOT EXTRACT, SCUTELLARIA BAICALENSIS ROOT EXTRACT, GLYCERYL STEARATE, PEG-100 STEARATE, PVM/MA DECADIENE CROSSPOLYMER, XANTHAN GUM, SILICA, ACRYLATES/CARBAMATE COPOLYMER, MALTODEXTRIN, LECTTHIN, TRIMETHYLSICATE, BUTYLENE GLYCOL, AMMONIUM HYDROXIDE, PHENOXYETHANOL, METHYLPARABEN, DISDOUM EDTA, BHT, PARFUM/ FRAGRANCE.

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Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10096-0290
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
octinoxate (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	o ctino xate	75 mg in 1 g	
oxybenzone (UNII: 9500S7VE0Y) (OXYBENZONE - UNII:950OS7VE0Y)	o xybe nz o ne	40 mg in 1 g	
avobenzone (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	avobenzone	20 mg in 1 g	

I	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10096-0290-2	1 in 1 CARTON		
1	NDC:10096-0290-1	75 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 not final	part352	11/11/2012	

Labeler - New Avon LLC (080143520)

Establishment			
Name	Address	ID/FEI	Business Operations
Avon Products, Inc		005149471	manufacture(10096-0290)

Revised: 3/2016 New Avon LLC