

ANEW REJUVENATE 24 HOUR EYE MOISTURIZER- octinoxate, octisalate, avobenzone, homosalate cream

Avon Products, Inc

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 ingredients

OCTINOXATE 7.50%, OCTISALATE 5.0%,
AVOBENZONE 2.0%, HOMOSALATE 2.0%.....

Purpose

.....Sunscreen

Uses

- helps prevent sunburn

Warnings

Skin Cancer/Skin Aging Alert: Spending time in the sun increases your risk of skin cancer and early skin aging. This product had been shown only to help prevent sunburn, **not** skin cancer or early skin aging.

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

Other Information

protect the product in this container from excessive heat and direct sun

Inactive ingredients:WATER/EAU, GLYCERIN, DIMETHICONE, BUTYLENE GLYCOL, PEG-8, POTASSIUM CETYL PHOSPHATE, PHENYL TRIMETHICONE, CETEARYL ALCOHOL, ISODECYL ISONONANOATE, PHYTOL, POLYMETHYL METHACRYLATE, TINOSPORA CORDIFOLIA ROOT/STEM EXTRACT, PINUS TAEDA BARK EXTRACT, RETINOL/SACCHAROMYCES POLYPEPTIDE, PHAEODACTYLUM TRICORNUTUM EXTRACT, GLYCINE SOJA (SOYBEAN) SEED EXTRACT, HELIANTHUS ANNUUS (SUNFLOWER) SEED EXTRACT, PLANKTON EXTRACT, HYDROLYZED MILK PROTEIN, SODIUM HYALURONATE, ANDROGRAPHOLIDE, PANTHENOL, TOCOPHEROL, KAEMPFERIA GALANGA ROOT EXTRACT, ORYZANOL, CETEARYL GLUCOSIDE, CAPRYLIC/CAPRIC TRIGLYCERIDE, CETYL ALCOHOL, CARBOMER, TROMETHAMINE, SODIUM DEHYDROACETATE, CHLORPHENESIN, PHENOXYETHANOL, DISODIUM EDTA.

Questions?1-800-FOR-AVON



Drug Facts For Eye Day Cream	
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Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10096-0149
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	75 mg in 1 mL	
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 mL	
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	20 mg in 1 mL	
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	20 mg in 1 mL	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10096-0149-2	1 in 1 CARTON		
1	NDC:10096-0149-1	10 mL in 1 JAR		
2	NDC:10096-0149-3	1.1 mL in 1 PACKET		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC monograph not final	part352	02/08/2013	
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Labeler - Avon Products, Inc (001468693)

Establishment

Name	Address	ID/FEI	Business Operations
Avon Products, Inc		005149471	manufacture(10096-0149)

Revised: 2/2013

Avon Products, Inc