

ANEW REVERSALIST COMPLETE RENEWAL DAY- homosalate, 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.

Anew Reversalist Complete Renewal Day Cream Broad Spectrum SPF 25

Active ingredient

- Homosalate 7.0%.....
- Octinoxate 7.0%.....
- Oxybenzone 4.0%.....
- Avobenzone 2.85%.....

Purpose

-Sunscreen
-Sunscreen
-Sunscreen
-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, dimethicone, butylene glycol, isodecyl isononanoate, tromethamine, dilauryl thiodipropionate, phytol, ethylhexylglycerin, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, cetearyl alcohol, nylon-12, polymethylsilsesquioxane, isohexadecane, phenoxyethanol,

polymethyl methacrylate, thiodipropionic acid, parfum/fragrance, carbomer, xanthan gum, cetareth-20, sodium PCA, acrylates/C10-30 alkyl acrylate crosspolymer, disodium EDTA, hydrogenated lecithin, kaempferia galanga root extract, silica, polysorbate 60, trimethylsiloxysilicate, dimethicone crosspolymer, thiazolylalanine, amorphophallus campanulatus rhizome/root extract, punica granatum fruit juice, sesbania grandiflora flower extract, portulaca oleracea extract, tocopherol, palmitoyl lysyl aminovaleroyl lysine.

Questions?

1-800-FOR-AVON or 1-800-265-AVON in Canada



Drug Facts	
Active Ingredients	Purpose
Homosalate 7.0%.....	Sunscreen
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Drug Facts (continued)

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Drug Facts (continued)

polymethylsilsesquioxane, isohexadecane, phenoxyethanol, polymethyl methacrylate, thiodipropionic acid, parfum/fragrance, carbomer, xanthan gum, cetareth-20, sodium PCA, acrylates/C10-30 alkyl acrylate crosspolymer, disodium EDTA, hydrogenated lecithin, kaempferia galanga root extract, silica, polysorbate 60, trimethylsiloxysilicate, dimethicone crosspolymer, thiazolylalanine, amorphophallus campanulatus rhizome/root extract, punica granatum fruit juice, sesbania grandiflora flower extract, portulaca oleracea extract, tocopherol, palmitoyl lysyl aminovaleroyl lysine.

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10096-0312
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	70 mg in 1 g
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	70 mg in 1 g
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	40 mg in 1 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	28.5 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10096-0312-2	1 in 1 CARTON	12/12/2013	
1	NDC:10096-0312-1	50 g in 1 JAR; Type 0: Not a Combination Product		
2	NDC:10096-0312-4	1 in 1 CARTON	12/12/2013	
2	NDC:10096-0312-3	15 g in 1 JAR; Type 0: Not a Combination Product		
3	NDC:10096-0312-5	1.1 g in 1 PACKET; Type 0: Not a Combination Product	12/12/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	12/12/2013	

Labeler - New Avon LLC (080143520)

