ANEW ULTIMATE DAY MULTI-PERFORMANCE- 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 Ultimate Multi-Performance Day Cream SPF 25

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

IFor sunscreen use:

- apply liberally 15 minutes before sun exposure
- children under 6 month 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

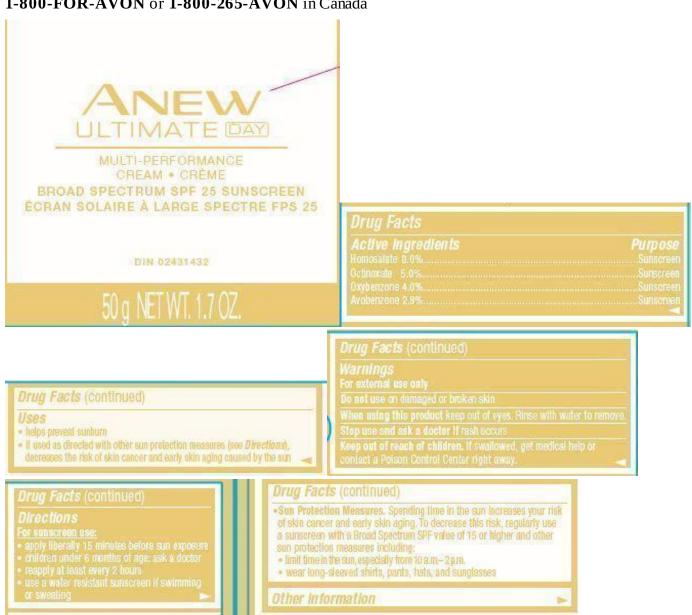
- protect the product in this container from excessive heat and direct sun
- product may stain certain types of fabrics

Inactive ingredients:

water/eau, glycerin, dimethicone, butylene glycol, glycolic acid, behenyl alcohol, hydrogenated polyisobutene, trisiloxane, ethylhexyl isononanoate, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, cetearyl alcohol, isohexadecane, phytol, thiodipropionic acid, PEG-100 stearate, ammonium hydroxide, parfum/fragrance, cetearyl glucoside, phenoxyethanol, dimethiconol, polysorbate 60, disodium EDTA, PEG-150 distearate, maltodextrin, tiliacora triandra leaf/vine extract, thiazolylalanine, saccharomyces ferment lysate filtrate, eclipta prostrata extract, saccharomyces/platinum ferment, isomalt, propylene glycol, SD alcohol 40-B, saccharomyces/copper ferment, saccharomyces/magnesium ferment, saccharomyces/malachite ferment, saccharomyces/tourmaline ferment, saccharomyces/zinc ferment, steareth-20, lecithin, vitis vinifera (grape) fruit cell extract, palmitoyl tetrapeptide-10, palmitoyl tetrapeptide-7, palmitoyl lysyl aminovaleroyl lysine, caramel, red 4/CI 14700.

Questions?

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



Drug Facts (continued)

- protect the product in this container from excessive heat and direct sun
- · product may stain certain types of fabrics

Inactive ingredients

water/eau, glycerin, dimethicone, butylene glycol, glycolic acid, behenyl alcohol, hydrogenated polyisobutene, trisiloxane, ethylhexyl isononanoate, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, cetearyl alcohol, isohexadecane, phytol, thiodipropionic acid, PEG-100 stearate, ammonium hydroxide, parfum/fragrance, cetearyl glucoside, phenoxyethanol, dimethiconol, polysorbate 60 disodium EDTA, PEG-150 distearate, maltodextrin, tiliacora triandra leat/vine extract, thiazolylatanine,

Drug Facts (continued)

saccharomyces/platinum ferment, isomalt, propylene glycol, SD alcohol 40-8, saccharomyces/capper ferment, saccharomyces/magnesium ferment, saccharomyces/magnesium ferment, saccharomyces/madachite ferment, saccharomyces/curmaline ferment, saccharomyces/curmaline ferment, saccharomyces/zinc ferment, steareth-20, lecithin, vits vinifera (grape) fruit cell extract, palmitoyl tetrapeptide-10, palmitoyl fetrapeptide-7, palmitoyl lysyly aminovaleroyl lysine, caramel, red 4/Cl 14700.

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Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:10096-0323

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S) HOMOSALATE OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51) OCTINOXATE OCTINOXATE OCTINOXATE OCTINOXATE OCTINOXATE OCTINOXATE OCTINOXATE

OXYBENZONE (UNII: 950OS7VE0 Y) (OXYBENZONE - UNII:95OOS7VE0 Y)

AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)

AVOBENZONE

AVOBENZONE

29 mg in 1 g

Inactive Ingredients

metre ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		

Packaging#Item CodePackage DescriptionMarketing Start DateMarketing End Date1NDC:10096-0323-21 in 1 CARTON12/19/2014

 1 NDC:10096-0323-1
 50 g in 1 JAR; Type 0: Not a Combination Product

 2 NDC:10096-0323-4
 1 in 1 CARTON
 12/19/2014

 2 NDC:10096-0323-3
 15 g in 1 JAR; Type 0: Not a Combination Product

 3 NDC:10096-0323-5
 1.1 g in 1 PACKET; Type 0: Not a Combination Product
 12/19/2014

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	12/19/2014	

Revised: 1/2019 New Avon LLC