AVON SUN SPORT REFRESH SUNSCREEN- homosalate, oxybenzone, octisalate, avobenzone, octocrylene lotion

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 ingredients HOMOSALATE 9.5%
OXYBENZONE 6.0%
OCTISALATE 4.75%
AVOBENZONE 3.0%
OCTOCRYLENE 2.8%
Purpose
Sunscreen

Uses

• helps prevent sunburn

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 generously and evenly 15 minutes before sun exposure
- children under 6 months of age: ask a doctor
- reapply after 80 minutes of swimming or sweating
- immediately after towel drying
- at least every 2 hours
- **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

• May stain some fabrics.

Inactive Ingredients:

water/eau, butyloctyl salicylate, dimethicone, PEG-8, styrene/acrylates copolymer, SD alcohol 40-B, silica, polyester-7, oleth-3 phosphate, neopentyl glycol diheptanoate, trisiloxane, dilauryl thiodipropionate, polyester-8, VP/eicosene copolymer, hydroxyethyl urea, boron nitride, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, tromethamine, dimethicone crosspolymer, menthyl PCA, parfum/fragrance,

phenoxyethanol, isohexadecane, PEG-8 laurate, hydrogenated lecithin, acrylates/C10-30 alkyl acrylate crosspolymer, chlorphenesin, disodium EDTA, polyglyceryl-3 diisostearate, menthol, oryzanol, panthenol, polysorbate 60, urea, kaempferia galanga root extract, phytol.

Questions? Call toll free 1-800-FOR-AVON



Drug Facts Active Ingredients **Purpose** Homosalate 9.5%.. Sunscreen Oxybenzone 6.0%.... Octisalate 4.75%.... Sunscreen .Sunscreen Avobenzone 3.0% .Sunscreen Octocrylene 2.8% helps prevent sunburn 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 dector if rash occurs Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Drug Facts (continued) Directions apply generously and evenly 15 minutes before sun exposure children under 6 months of age; ask a doctor reapply after 80 minutes of swimming or sweating · immediately after towel drying at least every 2 hours 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.

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Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10096-0313
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	95 mg in 1 mL	
OXYBENZONE (UNII: 950OS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	60 mg in 1 mL	
OCTISALATE (UNII: 4X49 Y0596W) (OCTISALATE - UNII:4X49 Y0596W)	OCTISALATE	47.5 mg in 1 mL	
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 mL	
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	28 mg in 1 mL	

l	Packaging				
ı	# Item Code Package Description		Marketing Start Date	Marketing End Date	
ı	1	NDC:10096-0313-1	236 mL 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	12/12/2013		

Labeler - New Avon LLC (080143520)

Establishment

Name	Address	ID/FEI	Business Operations
Avon Products, Inc		005149471	manufacture(10096-0313)

Revised: 2/2016 New Avon LLC