

CVS PHARMACY SPF 50 BEACH GUARD SUN- avobenzone, homosalate, octisalate, octocrylene, oxybenzone stick
CVS Pharmacy

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.

CVS Pharmacy SPF 50 Beach Guard Sunstick

Active ingredients

Avobenzone 3.0%, Homosalate 15.0%, Octisalate 5.0%, Octocrylene 10%, Oxybenzone 3.0%

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 doctor if

- rash occurs.

Keep out of reach of children.

If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

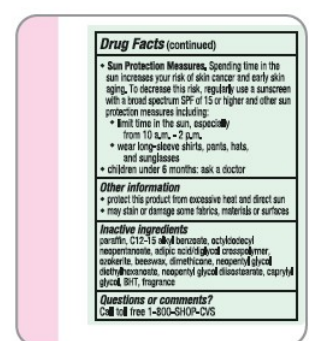
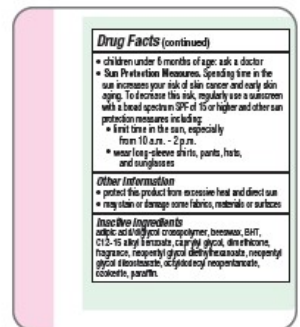
- apply liberally 15 minutes before sun exposure
- reapply:
 - after 80 minutes of swimming or sweating
 - immediately after towel drying
 - at least every 2 hours
- **Sun Protection Measures.** Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease this risk, regularly 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
- children under 6 months: Ask a doctor

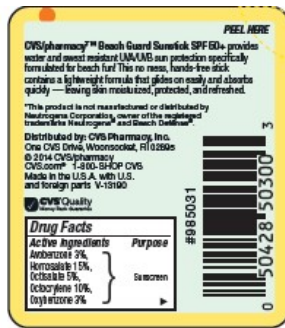
Other information

- protect the product in this container from excessive heat and direct sun
- may stain or damage some fabrics, materials or surfaces

Inactive ingredients

paraffin, C12-15 alkyl benzoate, octyldodecyl neopentanoate, adipic acid/diglycol crosspolymer, ozokerite, beeswax, dimethicone, neopentyl glycol diethylhexanoate, neopentyl glycol diisostearate, caprylyl glycol, BHT, fragrance





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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69842-149
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OXYBENZONE (UNII: 950OS7VE0Y) (OXYBENZ ONE - UNII:950OS7VE0Y)	OXYBENZ ONE	30 mg in 1 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	100 mg in 1 g
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	150 mg in 1 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZ ONE - UNII:G63QQF2NOX)	AVOBENZ ONE	30 mg in 1 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
CERESIN (UNII: Q1LS2UJO3A)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
PARAFFIN (UNII: I9O0E3H2ZE)	
ADIPIC ACID/DIGLYCOL CROSSPOLYMER (20000 MPA.S) (UNII: R9TPS68K19)	
WHITE WAX (UNII: 7G1J5DA97F)	
OCTYLDODECYL NEOPENTANOATE (UNII: X8725R883T)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
NEOPENTYL GLYCOL DIETHYLHEXANOATE (UNII: U68ZV6W62C)	
NEOPENTYL GLYCOL DIISOSTEARATE (UNII: 4M6OQ34JWW)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	

BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-149-28	42 g in 1 PACKAGE; Type 0: Not a Combination Product	10/30/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	10/30/2015	

Labeler - CVS Pharmacy (062312574)

Revised: 5/2022

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