GOODSENSE SUNSCREEN SPF 30- avobenzone, homosalate, octisalate, octocrylene lotion
Solskyn Personal Care 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.

GoodSense Sunscreen SPF 30

Active Ingredients

Avebenzone 3%,

Homosalate 10%

Octisalate 5%,

Octocrylene 10%

Purpose

Sunscreen

Warnings

For external use only.

Do not use

Do not use on damaged or broken skin.

When using this product

Stop use and ask a doctor

Stop use and ask a doctor if rash occurs.

Keep out of reach of children

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Directions

- Apply liberally 15 minutes before sun exposure
- Re-apply
  - after 80 minutes of swimming or sweating
  - Immediately after towel drying
  - at least every 2 hours
- DFD

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 broad spectrum SPF of 15 or higher and

other sun protection measures including:

- limit time in the sun, especially from 10a.m.-2p.m.
- war long-sleeved shirt, pants, hats, and sunglasses
- Children under 6 months: ask a doctor

Other information

Other information

- Protect the product in this container from excessive heat and direct sun.
- May stain fabric

Inactive ingredients

Inactive ingredients: Water, Glycerin, C28-52 Olefin/Undecylenic Acid Copolymer, Neopentyl Glycol, Diheptanoate, Stearic Acid, Polyglyceryl-3, Methylglucose, Distearate, Tapioca Starch, Glyceryl Stearate, PEG-100 Stearate, Microcrystalline Cellulose, Phenethyl Benzoate, Cellulose Gum, Tocopheryl Acetate, Fragrance, Polymethylsilsesquioxane, Caprylyl Glycol, Aloe Barbadensis Leaf Juice, Stearyl Alcohol, Dimethicone, Disodium EDTA, Sodium Hydroxide, Phenoxyethanol

Distributed by

Distributed by: Geiss, Destin & Dunn, Inc, Peachtree City, GA 30269
www.valuelabels.com/800-715-3485
GoodSense® is a registered trademark of L. Perrigo Company
*This product is not manufactured or distributed by Bayer, the owner of the registered trademark Coppertone®

Principal Display Panel

Principal Display Panel

GOODSENSE

SUNSCREEN

LOTION

BROAD SPECTRUM

SPF 30

- Water Resistant

(80 Minutes)

- UVA/UVB Protection
- Hypoallergenic
- Oxybenzone Free

Compare to active ingredients of

Coppertone

100%

Satisfaction

Guaranteed

8 FL OZ (237 mL)

PKLA-41423-GSF1

GOODSENSE SUNSCREEN SPF 30

avobenzone, homosalate, octisalate, octocrylene lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70281-601
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 g
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	10 g in 100 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	10 g in 100 g

Inactive Ingredients	
Ingredient Name	Strength
POLYMETHYLSILSESQUIOXANE (4.5 MICRONS) (UNII: 59Z907ZB69)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
NEOPENTYL GLYCOL DIHEPTANOATE (UNII: 5LKW3C543X)	
POLYGLYCERIN-3 (UNII: 4A0NCJ6RD6)	
METHYL GLUCOSE (UNII: QCF122NF3R)	
STARCH, TAPIOCA (UNII: 24SC3U704I)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
PEG-100 STEARATE (UNII: YD01N1999R)	
PHENETHYL BENZOATE (UNII: 0C143929GK)	
CARBOXYMETHYLCELLULOSE (UNII: 05JZI7B19X)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70281-601-24	237 g in 1 CONTAINER; Type 0: Not a Combination Product	01/14/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	01/14/2019	

Labeler - Solskyn Personal Care LLC (080010329)

Establishment

Name	Address	ID/FEI	Business Operations
Accra Pac, Inc.		024213616	manufacture(70281-601)

Revised: 6/2019 Solskyn Personal Care LLC