

SOLBAR THIRTY SPF 30- solbar thirty spf30 cream
Person and Covey

Solbar Thirty 30SPF

Purpose

Sun screen.

OTC - ACTIVE INGREDIENT SECTION

Homosalate

Octinoxate

Oxybenzone

Avobenzone

Octocrylene

INACTIVE INGREDIENT SECTION

Alcohol

Isobutyl Stearate

Cyclomethicone

PVP/Eicosene Copolymer

Dimethicone

Klucel Hydroxypropylcellulose

Acrylates/Octylacrylamide Copolymer

Indications and Use

Helps prevent sunburn. If used as directed with other sun protection measures, decreases the risk of skin cancer and early skin aging caused by the sun.

Dosage and Administration:

Apply liberally and evenly to all sun exposed areas of DRY skin 15 minutes before sun exposure. Reapply after 80 minutes of swimming or sweating and immediately after towel drying. Apply at least every 2 hours. For children under 6 months, ask a physician.

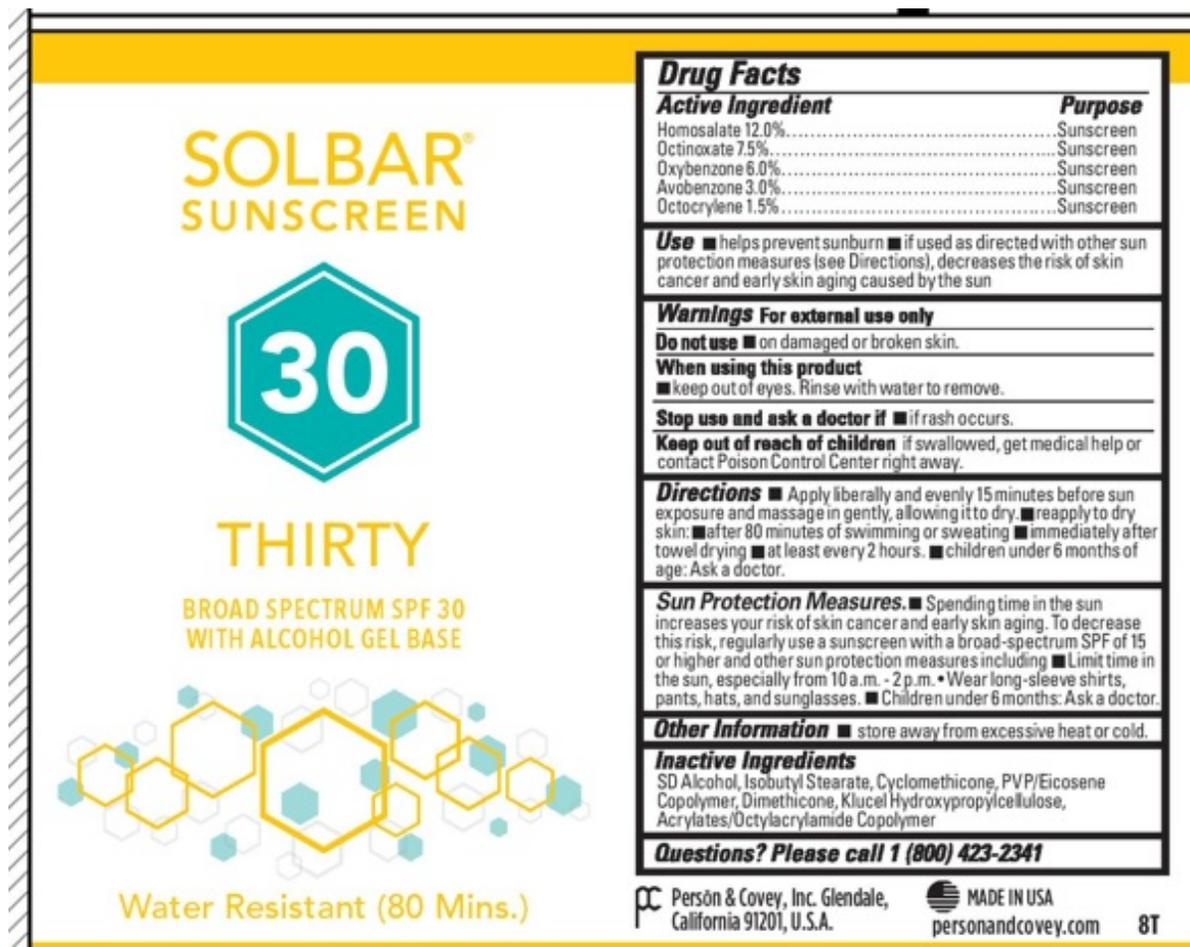
Keep out of Reach of Children:

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

Warnings

For external use only. Do not use on damaged or broken skin. Keep out of eyes. Rinse eyes thoroughly with water to remove. Stop use and ask a physician if rash or irritation develops and lasts. Store away from excessive heat and direct sun.

Package Label



SOLBAR[®]
SUNSCREEN

30

THIRTY

BROAD SPECTRUM SPF 30
WITH ALCOHOL GEL BASE

Water Resistant (80 Mins.)

Drug Facts

Active Ingredient	Purpose
Homosalate 12.0%	Sunscreen
Octinoxate 7.5%	Sunscreen
Oxybenzone 6.0%	Sunscreen
Avobenzone 3.0%	Sunscreen
Octocrylene 1.5%	Sunscreen

Use ■ 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 ■ if rash occurs.

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

Directions ■ Apply liberally and evenly 15 minutes before sun exposure and massage in gently, allowing it to dry. ■ reapply to dry skin: ■ after 80 minutes of swimming or sweating ■ immediately after towel drying ■ at least every 2 hours. ■ children under 6 months of age: Ask a doctor.

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 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-sleeve shirts, pants, hats, and sunglasses. ■ Children under 6 months: Ask a doctor.

Other Information ■ store away from excessive heat or cold.

Inactive Ingredients
SD Alcohol, Isobutyl Stearate, Cyclomethicone, PVP/Eicosene Copolymer, Dimethicone, Klucel Hydroxypropylcellulose, Acrylates/Octylacrylamide Copolymer

Questions? Please call 1 (800) 423-2341

pc Person & Covey, Inc. Glendale, California 91201, U.S.A.  MADE IN USA
personandcovey.com 8T

SOLBAR THIRTY SPF 30

solbar thirty spf30 cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0096-0685
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	0.12 g in 1 g

OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	0.075 g in 1 g
OXYBENZONE (UNII: 95OOS7VE0Y) (OXYBENZONE - UNII:95OOS7VE0Y)	OXYBENZONE	0.06 g in 1 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	0.03 g in 1 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	0.015 g in 1 g

Inactive Ingredients

Ingredient Name	Strength
CYCLOMETHICONE (UNII: NMQ347994Z)	
EICOSYL POVIDONE (2 EICOSYL BRANCHES/REPEAT) (UNII: XQQ9MKE2BJ)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
HYDROXYPROPYL CELLULOSE (UNII: RFW2ET671P)	
ALCOHOL (UNII: 3K9958V90M)	
ISOBUTYL STEARATE (UNII: V8DPR6HMX3)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0096-0685-04	105.6 g in 1 BOTTLE; Type 0: Not a Combination Product	06/01/1996	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	06/01/1996	

Labeler - Person and Covey (008482473)

Establishment

Name	Address	ID/FEI	Business Operations
Person and Covey		008482473	manufacture(0096-0685)

Revised: 12/2025

Person and Covey