SPF 30 SUNSCREEN STICK- avobenzone, homosalate, octisalate, octocrylene stick
OraLabs

Drug Facts

Active ingredient

Avobenzone 3.0%, Homosalate 10.0%, Octisalate 5.0%, Octocrylene 10%.....Sunscreen

Purpose

Sunscreen

Keep Out of Reach of Children

If swallowed get medical help or contact a Poison Control Center immediately.

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 or irritation develops and lasts.

Directions

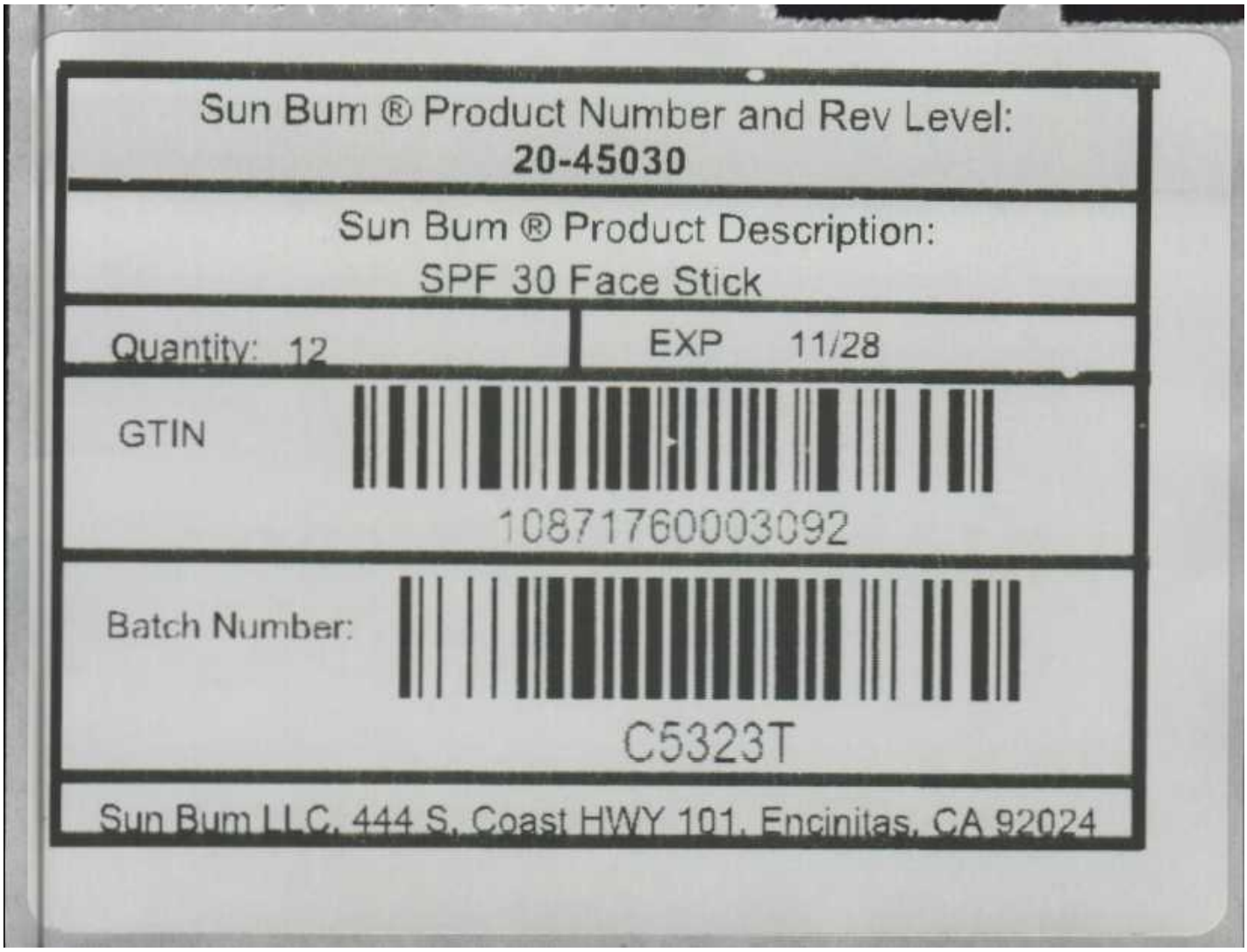
Apply liberally 15 minutes before sun exposure • reapply: after 80 minutes of swimming or sweating • immediately after towel drying • at least every 2 hours • **Sun Protection Measures.**Spending time in the sun increases your risk of skin cancer and early skin aging • To decrease the risk, regularly 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.

Inactive Ingredients

Paraffin wax, octyl palmitate, ozokerite, euphorbia cerifera (candelilla) wax, beeswax, butyloctyl salicylate, fragrance, polyethylene, polycrylene, cetyl alcohol, tocopheryl

acetate, ascorbyl palmitate

Package/Label Principal Display Panel



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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63645-329
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	30 mg in 1 g
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	100 mg in 1 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	50 mg in 1 g

OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	100 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
PARAFFIN (UNII: I9O0E3H2ZE)	240 mg in 1 g
ETHYLHEXYL PALMITATE (UNII: 2865993309)	200 mg in 1 g

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63645-329-01	13 g in 1 CONTAINER; Type 0: Not a Combination Product	12/08/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	352	12/08/2025	

Labeler - OraLabs (801824756)

Registrant - OraLabs (801824756)

Establishment

Name	Address	ID/FEI	Business Operations
OraLabs		801824756	manufacture(63645-329) , label(63645-329) , analysis(63645-329)

Revised: 12/2025 OraLabs