

GRAPEFRUIT SPF 30 BROAD SPECTRUM- sunscreen aerosol
Bell International Laboratories, Inc.

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.

Hint SPF 30 Sunscreen Spray Grapefruit

Active Ingredients

Avobenzone 1.9%, Octinoxate 7.1%, Octisalate 3.9%, Octocrylene 6.6%

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

Flammable Contents under pressure. Do not puncture or incinerate. Do not store at temperatures above 120 °F.

For external use only

Do not use on damaged or broken skin.

Stop use and ask a doctor if rash occurs.

When using this product keep out of eyes. Rinse with water to remove.

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

Directions

Apply liberally 15 minutes before sun exposure. Spray on skin and rub complete and even coverage. Do not spray on face; spray on hands and rub onto face.

Reapply:

- after 80 minutes of swimming or sweating
- immediately after towel drying
- at least every 2 hours

Sun Protection Measure. 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 10am-2pm
- wear long-sleeved shirts, pants, hats and sun glasses.

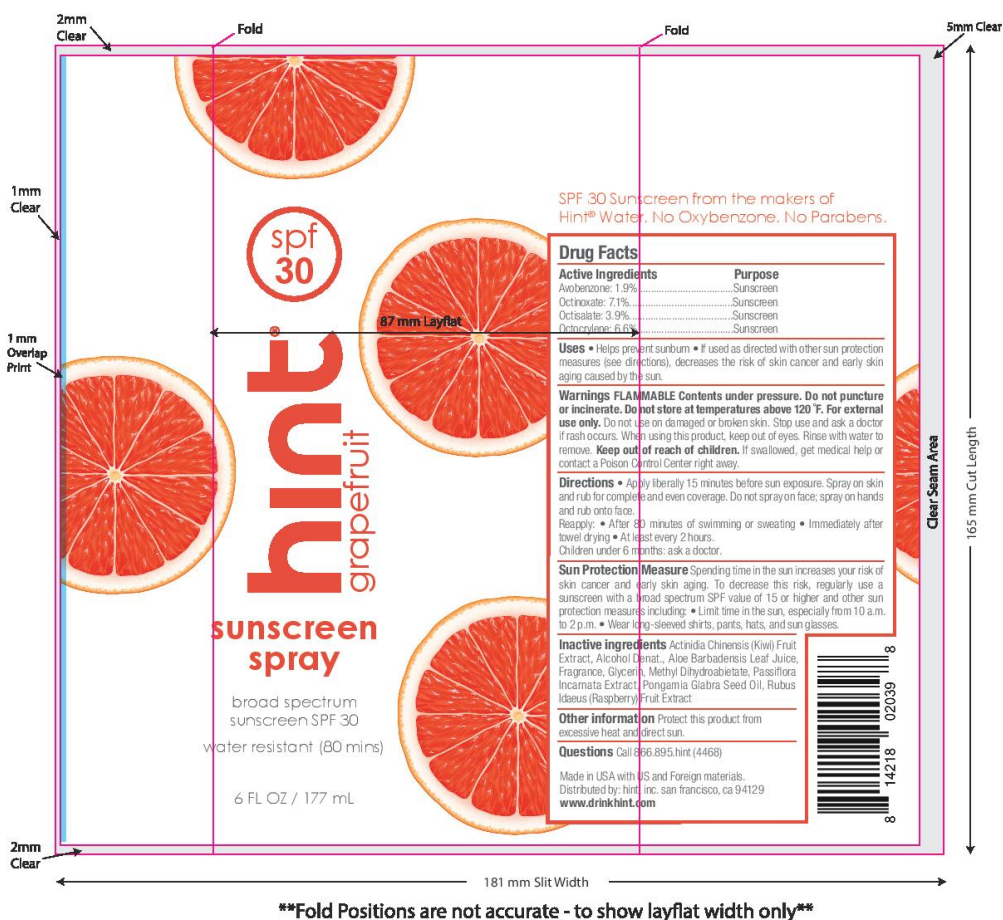
Inactive Ingredients

Actinidia Chinensis (Kiwi) Fruit Extract, Alcohol Denat., Aloe Barbadensis Leaf Huice, Fragrance, Glycerin, Methyl Dihydroabietate, Passiflora Incarnata Extract, Pongamia Glabra Seed Oil, Rubus Idaeus (Raspberry) Fruit Extract

Other Information

protect this product from excessive heat and direct sun.

PDP



GRAPEFRUIT SPF 30 BROAD SPECTRUM

sunscreens aerosol

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76150-204
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7.1 g in 100 mL
OCTISALATE (UNII: 4X49 Y0596 W) (OCTISALATE - UNII:4X49 Y0596 W)	OCTISALATE	3.9 g in 100 mL
OCTOCRYLENE (UNII: 5A68 WGF6 WM) (OCTOCRYLENE - UNII:5A68 WGF6 WM)	OCTOCRYLENE	6.6 g in 100 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	1.9 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958 V90 M)	
ALOE VERA LEAF (UNII: ZY8 1Z83 H0 X)	
PASSIFLORA INCARNATA FLOWER (UNII: K8 F3G29 S6 Z)	
METHYL DIHYDRO ABIETATE (UNII: 7666 FJ0 J9 F)	
RASPBERRY (UNII: 4N14V5R27W)	
KARUM SEED OIL (UNII: 62160 PU6 FJ)	
KIWI FRUIT (UNII: 71ES77 LGJC)	
GLYCERIN (UNII: PDC6 A3C0 OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76 150-204-13	178 mL in 1 CAN; Type 0: Not a Combination Product	11/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	11/01/2016	

Labeler - Bell International Laboratories, Inc. (967781555)

Revised: 1/2021

Bell International Laboratories, Inc.