BU SPF 30 BROAD SPECTRUM ALCOHOL-FREE PERFORMANCE SUNSCREEN - NATURAL CITRUS SCENT- octinoxate, octocrylene, octis alate, and avobenzone spray Bu Brands, 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.

bü SPF 30 Broad Spectrum Spray on Alcohol-Free Performance Sunscreen - Natural Citrus Scent

Drug Facts

Active Ingredients

Octinoxate 7.50% Octocrylene 7.50% Octisalate 5.00% Avobenzone 2.50%.

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.

Questions or Comments?

1.877.555.555 goodtobebu.com

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 occurs

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

Do not use near flame or while smoking

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
- Do not spray directly onto the face. Spray into hands, and apply to the face.
- children under 6 months: 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

Inactive Ingredients

Cyclopentasiloxane, Octyl Palmitate, Butyloctyl Salicylate, Diphenylsiloxy Phenyl Trimethicone, VP/Hexadecene Copolymer, Tocopherols, Bisabolol, Artemisia Pallens Flower Oil, Citrus Aurantium Dulcis (Orange) Peel Oil.

Other Information

• protect this product from excessive heat and direct sun

PRINCIPAL DISPLAY PANEL - 30 mL Bottle Label

SPF30

broad spectrum

spray-on alcohol-free performance sunscreen

Oil-Free Preservative-Free Natural Citrus Scent

bü

Water Resistant (80 Minutes) **1oz** (30mL)





spray-on alcohol-free performance

sunscreen

Oil-Free Preservative-Free Natural Citrus Scent



Water Resistant (80 Minutes) 10z (30mL)





Drug Facts

Active Ingredients Purpose
Octinovate 7.50%
Octocrylene 7.50%
Octisalate 5.00%
Avobenzone 2.50%.

Uses

helps prevent surdum
 it used as directed with
 other sun protection
 measures (see Directions),
 decreases the risk of skin
 cancer and early skin aging
 caused by the sun.

Questions or Comments? 1.877.555.555 goodtobebu.com

Manufactured for **bū brands, llc,** Azusa, CA 91702

PRODUCT OF USA Cruelty-free 7





BU SPF 30 BROAD SPECTRUM ALCOHOL-FREE PERFORMANCE SUNSCREEN - NATURAL CITRUS SCENT

octinoxate, octocrylene, octisalate, and avobenzone spray

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:70325-3003 Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7.5 g in 100 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	7.5 g in 100 mL
OCTISALATE (UNII: 4X49 Y0 59 6 W) (OCTIS ALATE - UNII:4X49 Y0 59 6 W)	OCTISALATE	5 g in 100 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	2.5 g in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
CYCLOMETHICONE 5 (UNII: 0THT5PCI0R)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8 D3)	
DIPHENYLSILO XY PHENYL TRIMETHICO NE (UNII: 1445L28B12)	
VINYLPYRROLIDONE/HEXADECENE COPOLYMER (UNII: KFR5QEN0N9)	
TOCOPHEROL (UNII: R0ZB2556P8)	
.ALPHABISABOLOL, (+/-)- (UNII: 36 HQN158 VC)	
DAVANA O IL (UNII: CL439 Y041G)	
ORANGE OIL (UNII: AKN3KSD11B)	

	Packaging					
:	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
	NDC:70325-3003-	98 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	12/0 1/20 16			
	NDC:70325-3003-	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/0 1/20 16			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC MONOGRAPH NOT FINAL	part352	12/0 1/20 16				

Labeler - Bu Brands, LLC (080075929)

Establishment				
Name	Address	ID/FEI	Business Operations	
Westwood Laboratories, Inc		832280635	MANUFACTURE(70325-3003)	

Revised: 1/2021 Bu Brands, LLC