

BU SPF 50 S BROAD SPECTRUM ALCOHOL-FREE PERFORMANCE SUNSCREEN - NATURAL WHITE SAGE SCENT- octinoxate, octocrylene, octisalate, avobenzone, and homosalate 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 50 S Broad Spectrum Spray on Alcohol-Free Performance Sunscreen - Natural White Sage Scent

Drug Facts

Active Ingredients

Octocrylene 10.00%
Octinoxate 7.50%
Homosalate 7.50%
Octisalate 5.00%
Avobenzone 3.00%

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

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, Butyloctyl Salicylate, Ethylhexyl Methoxycrylene, Diphenylsiloxy Phenyl Trimethicone, VP/Hexadecene Copolymer (and) Octyldodecanol, Polybutene, Tocopherols, Bisabolol, Lavendula Angustifolia Oil, Citrus Aurantium Bergamia (Bergamot) Fruit, Salvia Officinalis (Sage) Oil and Santalum Album (Sandlewood) Oil.

Other Information

- protect this product from excessive heat and direct sun

Questions or Comments?

Call: 310-456-8787 www.goodtobebu.com

PRINCIPAL DISPLAY PANEL - 98 mL Bottle Label

SPF50

broad
spectrum

spray-on
alcohol-free
performance
sunscreen

Oil-Free

Preservative-Free

with Antioxidants

Natural White Sage

Scent

bü

Water Resistant (80 Minutes)

3.3oz (98mL)

SPF **50**
broad spectrum

spray-on
alcohol-free
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Scent

bü

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Manufactured for bü brands, llc, Azusa, CA 91702

PRODUCT
OF USA
Cruelty-free

PRODUIT DE
ÉTATS-UNIS
Non testé sur
les animaux





BU SPF 50 S BROAD SPECTRUM ALCOHOL-FREE PERFORMANCE SUNSCREEN - NATURAL WHITE SAGE SCENT

octinoxate, octocrylene, octisalate, avobenzone, and homosalate spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70325-5003
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	10 g in 100 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	7.5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CYCLOMETHICONE 5 (UNII: 0THT5PC10R)	
ETHYLHEXYL METHOXYCRYLENE (UNII: S3KFG6Q5X8)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
DIPHENYLSILOXY PHENYL TRIMETHICONE (UNII: I445L28B12)	
VINYLPYRROLIDONE/HEXADECENE COPOLYMER (UNII: KFR5QEN0N9)	
OCTYLDODECANOL (UNII: 461N1O614Y)	
TOCOPHEROL (UNII: R0ZB2556P8)	
POLYBUTENE (1400 MW) (UNII: 1NA5AO9GH7)	
.ALPHA.-BISABOLOL, (+/-)- (UNII: 36HQN158VC)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
SAGE OIL (UNII: U27K0H1H2O)	
SANDALWOOD OIL (UNII: X7X01WMQ5F)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70325-5003-3	98 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2016	
2	NDC:70325-5003-1	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part352	12/01/2016	

Labeler - Bu Brands, LLC (080075929)

Establishment

Name	Address	ID/FEI	Business Operations
Westwood Laboratories, Inc		832280635	MANUFACTURE(70325-5003)

Revised: 1/2021

Bu Brands, LLC