

PAUL FRANK SUNNY DEFENSE SPF STICK- avobenzone, homosalate, octisalate, octocrylene stick
RBGROUP Co., Ltd

84326-624 Paul Frank Sunny Defense Sunscreen stick

Avobenzone 2.5%

Homosalate 9.5%

Octisalate 4.5%

Octocrylene 9.5%

Sunscreen

- 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.

For external use only

- on damaged or broken skin

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

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

and ask a doctor if rash occurs.

- Protect this product from excessive heat and direct sun.

Directions

- Apply liberally 15 minutes before sun exposure.
- Use a water resistant sunscreen if swimming or sweating.
- Reapply 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.

Ethylhexyl Palmitate, Synthetic Wax, Butyloctyl Salicylate, Dicaprylyl Carbonate, Ethylhexylglycerin, Water, Tocopherol, Allantoin, Butylene Glycol, 1,2-Hexanediol, Centella Asiatica Extract, Hydrolyzed Hyaluronic Acid, Hydrolyzed Sodium Hyaluronate, Dimethylsilanol Hyaluronate, Sodium Hyaluronate, Hyaluronic Acid, Potassium Hyaluronate, Sodium Hyaluronate Crosspolymer, Hydroxypropyltrimonium Hyaluronate, Sodium Hyaluronate Dimethylsilanol, Sodium Acetylated Hyaluronate



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avobenzone, homosalate, octisalate, octocrylene stick

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84326-624
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTOCRYLENE (UNII: 5A68WGF6VM) (OCTOCRYLENE - UNII:5A68WGF6VM)	OCTOCRYLENE	1.71 g in 18 g
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	0.45 g in 18 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	0.81 g in 18 g
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	1.71 g in 18 g

Inactive Ingredients

Ingredient Name	Strength
CENTELLA ASIATICA TRITERPENOIDS (UNII: 4YS74Q4G4J)	
DIMETHYLSILANOL HYALURONATE (UNII: Z853O1D4TE)	
SODIUM ACETYLATED HYALURONATE (UNII: WN66R7GL93)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
HYALURONIC ACID (UNII: S270N0TRQY)	
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)	
TOCOPHEROL (UNII: R0ZB2556P8)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
WATER (UNII: 059QF0K00R)	
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
ETHYLHEXYL PALMITATE (UNII: 2865993309)	
ALLANTOIN (UNII: 344S277G0Z)	
SYNTHETIC WAX (1900 MW) (UNII: 5M631N9V0S)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84326-624-02	1 in 1 BOX	11/05/2024	
1	NDC:84326-624-01	18 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	11/05/2024	

Labeler - RBGROUP Co., Ltd (987610097)

Establishment

Name	Address	ID/FEI	Business Operations
reBom Co., Ltd		695951708	manufacture(84326-624)

Revised: 11/2024

RBGROUP Co., Ltd