

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

84326-424 Miffy Sunny Defense SPF 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 skin or broken skin
- and ask a doctor** if rash occurs.
- Keep out of eyes. Rinse with water to remove.

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

- 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 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

PDP_Miffy Sunny Defense SPF Stick



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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84326-424
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	0.55 g in 22 g
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	2.09 g in 22 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	2.09 g in 22 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	0.99 g in 22 g

Inactive Ingredients

Ingredient Name	Strength
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
WATER (UNII: 059QF0KO0R)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	

HYALURONATE SODIUM (UNII: YSE9PPT4TH)
ETHYLHEXYL PALMITATE (UNII: 2865993309)
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)
BUTYLOCTYL SALICYLATE (UNII: 2EH13UN8D3)
TOCOPHEROL (UNII: ROZB2556P8)
DIMETHYLSILANOL HYALURONATE (UNII: Z853O1D4TE)
ALLANTOIN (UNII: 344S277G0Z)
CENTELLA ASIATICA TRITERPENOIDS (UNII: 4YS74Q4G4J)
SODIUM ACETYLATED HYALURONATE (UNII: WN66R7GL93)

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84326-424-02	1 in 1 BOX	10/15/2024	
1	NDC:84326-424-01	22 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	10/15/2024	

Labeler - RBGROUP Co., Ltd (987610097)

Establishment

Name	Address	ID/FEI	Business Operations
reBom Co., Ltd		695951708	manufacture(84326-424)