

SNUGZ SPF 50 SUNSCREEN SPRY- spf 50 sunscreen spray spray
SnugZ USA

ZSUNPL - SPF 50 Sunscreen Spray

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ZSUNPLS - SPF 50 Sunscreen Spray

Made in the USA from Domestic and Globally Sourced Materials. PABA FREE.

1.0 fl. oz. (28 mL.) BROAD SPECTRUM SPF 50 Unscented Sunscreen Spray

Peel here for Drug Facts

Drug Facts	Purpose:
Active Ingredients: Avobenzone 2.7%, Homosalate 13.5%, Octisalate 4.5%, Octocrylene 9%..... 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.	
Directions: Directions: • Apply liberally 15 minutes before sun exposure • Use a water-resistant sunscreen if swimming or sweating or immediately after towel drying • Reapply: • at least every 2 hours. • Sun Protection Measures. Spending time in the sun increases your risk of skin cancer and early skin aging. To decrease the risk, regularly use a sunscreen	

Drug Facts (Continued) with a broad spectrum SPF of 15 or higher and other sun protection measures including: • Limit time in sun, especially from 10 a.m. - 2 p.m. • Wear long-sleeve shirts, pants, hats, and sunglasses. • Children under six months of age. Ask a doctor.	Other Information: • Protect this product from excessive heat and direct sun.
Inactive Ingredients: Isobutyl Denat., C12-15 Alkyl Brzozate Acrylates, Octylacrylamide Copolymer, Glycerin, Vitamin E	
Questions? 1.800.611.4270	

DISTRIBUTED BY: (COMPANY NAME)
(COMPANY CITY, STATE ZIP)

Under Wrap

Visible Label Area

SNUGZ SPF 50 SUNSCREEN SPRY			
spf 50 sunscreen spray spray			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76309-525

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	4.2 g in 28 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	1.4 g in 28 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	2.8 g in 28 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	0.84 g in 28 mL

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76309-525-10	10 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/01/2019	
2	NDC:76309-525-71	28 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	06/01/2019	

Labeler - SnugZ USA (615959228)

Registrant - SnugZ/USA, LLC (615959228)

Establishment

Name	Address	ID/FEI	Business Operations
SnugZ USA		615959228	manufacture(76309-525)

Revised: 1/2024

SnugZ USA