

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

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**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**

<b>Drug Facts</b>	<b>Purpose:</b>
<b>Active Ingredients:</b> Avobenzone 2.7%, Homosalate 13.5%, Octisalate 4.5%, Octocrylene 9%..... Sunscreen	
<b>Uses:</b> • Helps prevent sunburn. • If used as directed with other sun protection measures (see <b>Directions</b> ), decreases the risk of skin cancer and early skin aging caused by the sun.	
<b>Warnings:</b> 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. <b>Stop use and ask a doctor</b> if rash occurs. <b>Keep out of reach of children.</b> If product is swallowed, get medical help.	
<b>Directions:</b> <b>Directions:</b> • 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	

<b>Drug Facts (Continued)</b> 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.	<b>Other Information:</b> • Protect this product from excessive heat and direct sun.
<b>Inactive Ingredients:</b> Isobutyl Denat., C12-15 Alkyl Brzozate Acrylates/Octylacrylamide Copolymer, Glycerin, Vitamin E	<b>Questions?</b> 1.800.611.4270

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

Under Wrap

Visible Label Area

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

**Route of Administration** TOPICAL

### Active Ingredient/Active Moiety

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

### Inactive Ingredients

Ingredient Name	Strength
<b>ALCOHOL</b> (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: 12/2025

SnugZ USA