

SNUGZ HAND SANITIZER GEL- hand sanitizer gel gel
SnugZ/USA, LLC

ZS - Hand Sanitizer Gel

Wet hands thoroughly with product and rub into skin until dry.

Warnings: For external use only. Flammable, keep away from fire. Do not use on damaged or broken skin. **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.

Inactive Ingredients: Aminomethyl Propanol, Carbomer, Water

Active Ingredients: Ethyl Alcohol 70% v/v

Purpose: Antiseptic

Keep out of reach of children.

Indications: Helps reduce bacteria on skin.

Hand Sanitizer Gel

Drug Facts	Purpose Antiseptic
Active Ingredient Ethyl Alcohol 70% v/v	
Uses: Helps reduce bacteria on skin.	
Warnings: For external use only. Flammable, keep away from fire. 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.	
Directions: Wet hands thoroughly with product and rub into skin until dry.	
Inactive Ingredients: Aminomethyl Propanol, Carbomer, Water	
Other: Store below 110°F (43°C)	
Questions? 1.800.611.4270	MADE IN THE USA.
DISTRIBUTED BY: (COMPANY NAME) (COMPANY CITY, STATE ZIP)	

1.0 fl. oz. (28 mL.) Unscented Hand Sanitizer

SNUGZ HAND SANITIZER GEL			
hand sanitizer gel gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76309-301
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
CARBOMER 940 (UNII: 4Q93RCW27E)	
AMINOMETHYL PROPANEDIOL (UNII: CZ7BU4QZJZ)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76309-301-01	28 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	
2	NDC:76309-301-02	56 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	
3	NDC:76309-301-04	112 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	
4	NDC:76309-301-08	224 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/01/2018	
5	NDC:76309-301-19	56 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	
6	NDC:76309-301-51	30 mL in 1 TUBE; Type 0: Not a Combination Product	01/01/2018	
7	NDC:76309-301-05	14 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	
8	NDC:76309-301-81	28 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	
9	NDC:76309-301-91	28 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	
10	NDC:76309-301-61	39.75 mL in 1 POUCH; Type 0: Not a Combination Product	01/01/2018	
11	NDC:76309-301-62	85.17 mL in 1 POUCH; Type 0: Not a Combination Product	01/01/2018	12/31/2020
12	NDC:76309-301-99	3785.41 mL in 1 JUG; Type 0: Not a Combination Product	04/20/2020	
13	NDC:76309-301-88	112 mL in 1 POUCH; Type 0: Not a Combination Product	04/20/2020	12/31/2020

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/01/2018	

Labeler - SnugZ/USA, LLC (615959228)

Registrant - SnugZ/USA, LLC (615959228)

Establishment

Name	Address	ID/FEI	Business Operations
SnugZ/USA, LLC		615959228	manufacture(76309-301)

Revised: 12/2023

SnugZ/USA, LLC