

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

ZSB- Beaded 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, Vitamin E, Water

Active Ingredient: Ethyl Alcohol 70% v/v

Purpose: Antiseptic

Keep out of reach of children.

Indications: Helps reduce bacteria on skin.

Beaded Hand Sanitizer Gel

Drug Facts

Active Ingredients: Ethyl Alcohol 70% v/v... Purpose: Antiseptic

Uses: Helps reduce bacteria on skin.

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Questions? 1.800.611.4270

Made in the USA From Domestic and Globally Sourced Materials

DISTRIBUTED BY: (COMPANY NAME)

(COMPANY CITY, STATE, ZIP)

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

SNUGZ BEADED HAND SANITIZER GEL

beaded hand sanitizer gel gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76309-351
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	
AMINOMETHYL PROPANEDIOL (UNII: CZ7BU4QZJZ)				
ETHYLHEXYL PALMITATE (UNII: 2865993309)				
WATER (UNII: 059QF0KO0R)				
CARBOMER 940 (UNII: 4Q93RCW27E)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76309-351-01	28 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	
2	NDC:76309-351-02	56 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	
3	NDC:76309-351-04	112 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	
4	NDC:76309-351-08	224 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/01/2018	
5	NDC:76309-351-19	56 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	
6	NDC:76309-351-51	30 mL in 1 TUBE; Type 0: Not a Combination Product	01/01/2018	
7	NDC:76309-351-61	39.75 mL in 1 POUCH; Type 0: Not a Combination Product	01/01/2018	
8	NDC:76309-351-62	85.17 mL in 1 POUCH; Type 0: Not a Combination Product	01/01/2018	12/31/2020
9	NDC:76309-351-81	28 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	
10	NDC:76309-351-91	28 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	
11	NDC:76309-351-05	14 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2018	
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(76309-351)

Revised: 12/2023

SnugZ/USA, LLC