SANSPRAY-NA- non-alcohol antibacterial hand sanitizer spray Beacon Promotions, Inc.

Hand Sanitizer Spray

Purpose

Antiseptic

Warnings:

For external use only. Flammable. Keep away from heat or flame. Keep out of eyes, ears & mouth. In case of contact with eyes, rinse eyes thoroughly with water. Stop use and ask doctor if

redness or irritation develop and persist for more than 72 hours. Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away. Do not use on children less than 2 months of age. Do not use on open skin wounds.

Directions

Spray enough product on hands to cover all surfaces. Rub hands together until dry.

Inactive Ingredient:

Water, Aloe Vera, Glycerin, Propylene Glycol, Tocopherol, Lemon Fragrance.

Purpose

Anticeptic

Drug Facts

Active Ingredient Purpose Benzalkonium Chloride (0.1%)

Spray enough product on hands to cover all surfaces. Rub hands together until dry.

Keep out of reach of children

keep out of reach of children

Drug Facts		NON-ALCOHOLIC HAND SANITIZER		
Active Ingredient	Purpose	Directions: Spray enough product on hands to cover		
Benzalkonium Chloride (0.1%)	Antisentic	all surfaces. Rub hands together until dry		

Warnings: For external use only. Flammable. Keep away from heat or flame. Keep out of eyes, ears & mouth. In case of contact with eyes, rinse eyes thoroughly with water. Stop use and ask doctor if redness or irritation develop and persist for more than 72 hours. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Do not use on children less than 2 months of age. Do not use on open skin wounds.

Inactive Ingredients: Water, Aloe Vera, Glycerin, Propylene Glycol, Tocopherol, Lemon Fragrance.

Manufactured by Beacon Promotions, Inc., DBA Mixie Eagan, MN 55121 MADE in the USA with global materials LOT# 230216 EXP: 02-25 (0.33 oz.)

OVERLAP

SANSPRAY-NA

non-alcohol antibacterial hand sanitizer spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:70445-606

Route of Administration TOPICAL

Active Ingredient/Active Moiety

ı	,		
	Ingredient Name	Basis of Strength	Strength
	BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mg in 100 mL

Inactive Ingredients		
Ingredient Name	Strength	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
ALOE (UNII: V5VD430YW9)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
GLYCERIN (UNII: PDC6A3C0OX)		

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:70445-606-01	9.76 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/03/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/03/2024	

Labeler - Beacon Promotions, Inc. (119056382)

Establishment				
Name	Address	ID/FEI	Business Operations	
Beacon Promotions Inc.		119056382	manufacture(70445-606)	

Revised: 1/2024 Beacon Promotions, Inc.