

INSTANT HAND SANITIZER- alcohol gel
Safetec of America

61010-1112-2,-3,-8, Instant Hand Sanitizer

Active ingredient

Ethyl Alcohol 66.5%

Purpose

Antiseptic

Uses

- For handwashing to decrease bacteria on skin without soap and water

Warnings

For external use only

Flammable, keep away from fire or flame.

Keep out of reach of children.

Do not use in eyes, if this happens, rinse thoroughly with water.

Stop use, ask a doctor if irritation develops and persists for 72 hours. If ingested, get medical help or contact a Poison Control Center right away.

Directions

- wet hands & wrists thoroughly with product and allow to dry without wiping

Inactive ingredients

aloe vera, carbomer, D&C green #5, D&C yellow #10, fragrance, purified water, triethanolamine

PRINCIPAL DISPLAY PANEL - 0.9g Pouch Label

Safetec

A.B.H.C. TM MEETS CDC HANDWASHING RECOMMENDATIONS

**Instant
Hand
Sanitizer**

0.9 g (1/32 oz.)

Safetec of America, Inc.

Buffalo, NY 14215

800-456-7077

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Drug Facts

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PRINCIPAL DISPLAY PANEL - 144 Packets Box Label

NDC 61010-1112-3

Safetec

Infection Control

Instant Hand Sanitizer

Fresh Scent Instant
Hand Sanitizer with
Aloe Vera

A.B.H.C.™ MEETS CDC HANDWASHING RECOMMENDATIONS

144 Packets

Each Packet Net. Wt. 0.9g



INSTANT HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:61010-1112
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL	665 mL in 1 L
Inactive Ingredients				
Ingredient Name			Strength	
ALOE (UNII: V5VD430YW9)				
WATER (UNII: 059QF0KO0R)				
TROLAMINE (UNII: 9O3K93S3TK)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61010-1112-2	25 in 1 BOX	12/15/2010	
1	NDC:61010-1112-1	0.034 L in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:61010-1112-3	144 in 1 BOX	12/15/2010	
2	NDC:61010-1112-1	0.034 L in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:61010-1112-8	10 in 1 BOX	04/01/2020	
3	NDC:61010-1112-1	0.034 L in 1 POUCH; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		505G(a)(3)	12/15/2010	

Labeler - Safetec of America (874965262)

Establishment

Name	Address	ID/FEI	Business Operations
Safetec of America, Inc.		874965262	manufacture(61010-1112)

Revised: 2/2024

Safetec of America