

INSTANT HAND SANITIZER- alcohol gel

Safetec of America, Inc.

Reference Label Set Id: 3784c55f-928f-4766-902a-d09ba1ed42ae

NDC 61010-4111-1, Instant Hand Sanitizer-alcohol gel

Drug Facts

Active Ingredient

Ethyl Alcohol 70%

Purpose

Antiseptic

Uses

- for handwashing to decrease bacteria on skin whenever soap and water handwashing is not readily available
- helps prevent the risk of cross-contamination of bacteria that potentially can cause disease or infection
- recommended for repeated use

Warnings

For external use only. Flammable, keep away from fire or flame

When using this product

- do not get into eyes. If contact occurs, rinse eyes thoroughly with water

Stop use and ask doctor if

- irritation and redness develop and persists for more than 72 hours

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away

Directions

- place a small amount into one hand
- spread over both hands and wrist
- rub into the skin until dry
- do not wipe off; no rinsing required

Inactive ingredients

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

triethanolamine

Manufactured by **SAFETEC OF AMERICA, Inc.** Buffalo, NY 14215 800-456-7077
www.safetec.com

PRINCIPAL DISPLAY PANEL

PRINCIPAL DISPLAY PANEL - 34,688 fl. oz. Container Label

NDC 61010-4111-1

Safetec

instant

Hand

Sanitizer

Kills 99.9% of Germs

Enriched with

Aloe Vera

Fresh

34,688 fl.oz. (1,026 L) Reorder No. 17392



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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:61010-4111
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
D&C GREEN NO. 5 (UNII: 8J6RDU8L9X)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
WATER (UNII: 059QF0KO0R)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61010-4111-1	1026000 mL in 1 CONTAINER; Type 0: Not a Combination Product	12/16/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	12/16/2010	

Labeler - Safetec of America, Inc. (874965262)

Registrant - Safetec of America, Inc. (874965262)

Establishment

Name	Address	ID/FEI	Business Operations
Safetec of America, Inc.		874965262	manufacture(61010-4111)