

**INSTANT HAND SANITIZER- alcohol gel**  
**Safetec of America, Inc.**

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**61010-1112, Instant Hand Sanitizer**

**Drug Facts**

**Active Ingredients**

Ethyl Alcohol 66.5%

**Purpose**

Antiseptic

**Uses**

- for handwashing to decrease bacteria on skin without soap and water
- recommended for repeated use

**Warnings**

**For external use only**

**Flammable, keep away from fire or flame**

**Do not use** in the eyes. If this happens, rinse 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 ingested get medical help or contact a Poison Control Center right away

**Directions**

- wet hands & wrists thoroughly with product
- allow to dry without wiping
- children under 6 should be supervised while using this product

**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 - 8 fl. oz. Bottle Label**

**Safetec**

**instant**

**Hand**

**Sanitizer**

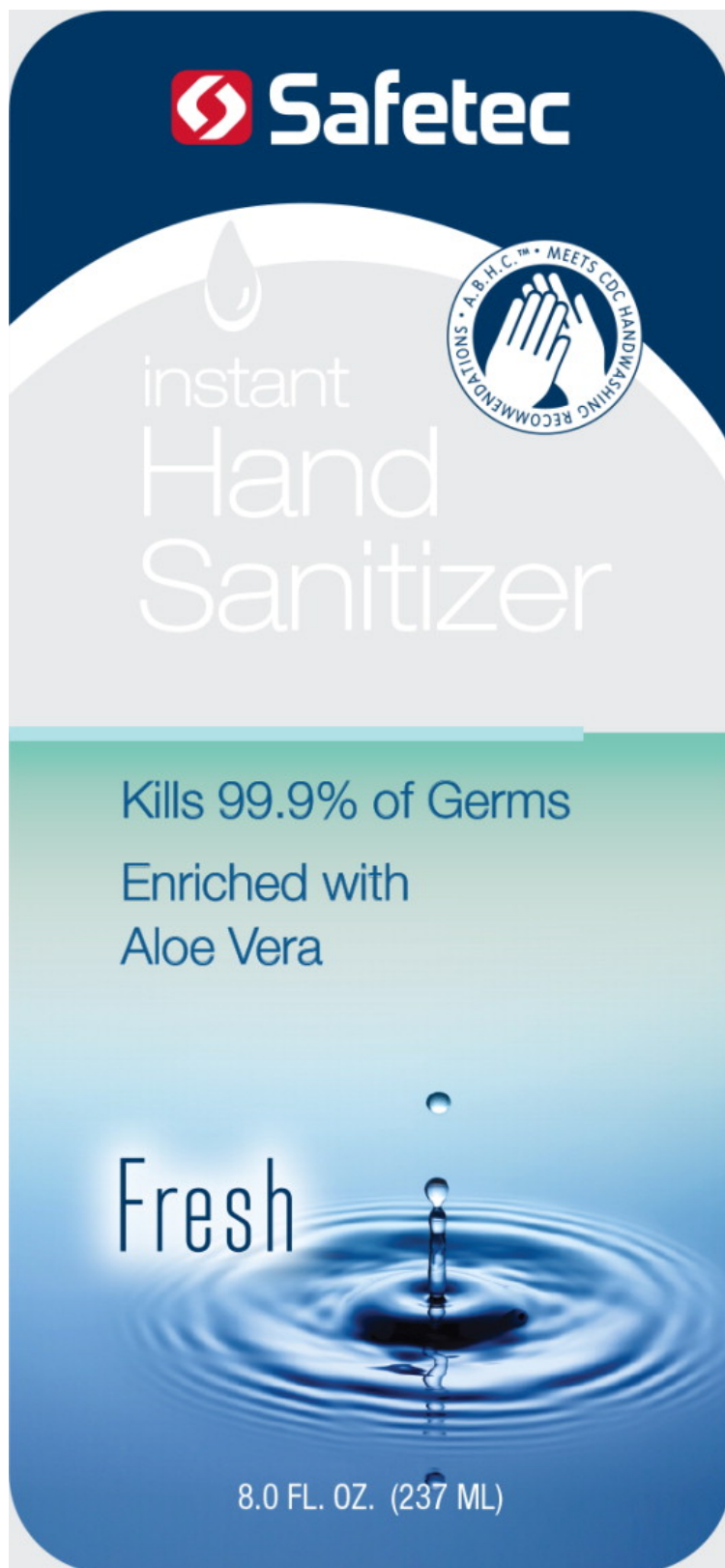
Kills 99.9% of Germs

Enriched with

Aloe Vera

**Fresh**

8.0 FL. OZ. (237 ML)



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Reorder No. 17370 (Cap)

Reorder No. 17371 (Pump)

## INSTANT HAND SANITIZER

alcohol gel

### Product Information

**Product Type**

HUMAN OTC DRUG

**Item Code (Source)**

NDC:61010-1112

**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
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
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:61010-1112-5	0.118 L in 1 BOTTLE; Type 0: Not a Combination Product	03/22/2012	
2	NDC:61010-1112-6	0.237 L in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/22/2012	
3	NDC:61010-1112-0	0.046 L in 1 BOTTLE; Type 0: Not a Combination Product	03/22/2012	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	03/22/2012	

**Labeler** - Safetec of America, Inc. (874965262)**Establishment**

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