INSTANT HAND SANITIZER- instant hand sanitizer gel
Global Protection USA, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Instant Hand Sanitizer

Drug Facts

Active Ingredient

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 a 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 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

Distributed by
Global Protection USA, Inc.
West Berlin, NJ 08091
800-957-8955

PACKAGE LABEL - PRINCIPAL DISPLAY PANEL – BOTTLE

Global Protection, Inc. “Provider of Homeland Security Equipment”

Instant Hand Sanitizer

Kills 99.9% of Germs

Enriched with Aloe Vera

2 Fl. oz. (59ML)

INSTANT HAND SANITIZER

instant hand sanitizer gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50225-355
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	Alcohol	60 g in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Aloe (UNII: V5VD430YW9)	
D&C Green no. 5 (UNII: 8J6RDU8L9X)	
D&C Yellow no. 10 (UNII: 35SW5USQ3G)	
water (UNII: 059QF0KO0R)	

Product Characteristics			
Color	green (green)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:50225-355-01	59 mL in 1 BOTTLE		
2	NDC:50225-355-02	236 mL in 1 BOTTLE		
3	NDC:50225-355-03	473 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333	01/06/2010	

Labeler - Global Protection USA, Inc. (030123488)

Registrant - Safetec of America, Inc. (874965262)

Establishment

Name	Address	ID/FEI	Business Operations
Safetec of America, Inc.		874965262	MANUFACTURE

Revised: 1/2010 Global Protection USA, Inc.