FOREVER HAND SANITIZER HAND SANITIZER- alcohol gel
Aloe Vera of America, 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.

Drug Facts for an OTC drug listing SPL for Forever Hand Sanitizer

Active Ingredients

Active ingredient	Purpose
Ethyl Alcohol 62%	Antimicrobial

Inactive Ingredients

Water (Aqua)

Aloe Barbadensis Leaf Juice (Stabilized* Aloe Vera Gel)

Honey

Isopropyl Alcohol

Fragrance (Parfum)

Glycerin

Tetrahydroxypropyl

Ethylenediamine

Acrylates/C10-30 Alkyl Acrylate Crosspolymer

Tocopheryl Acetate
Isopropyl Myristate.

Purpose Section

Uses

- hand sanitizer to decrease bacteria on the skin
- recommended for repeated use

Warnings

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Fore external use only.

Flammable. Keep away from fire or flame.

Indications & Usage

- do not use in or near the eyes. If contact occurs, rinse throughly with water.
- avoid contact with broken skin.

Stop use and ask a doctor if

Stop use and ask a doctor if

- rash or irritation develops and lasts.

Keep out of reach of children

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

Dosage & Administration Directions

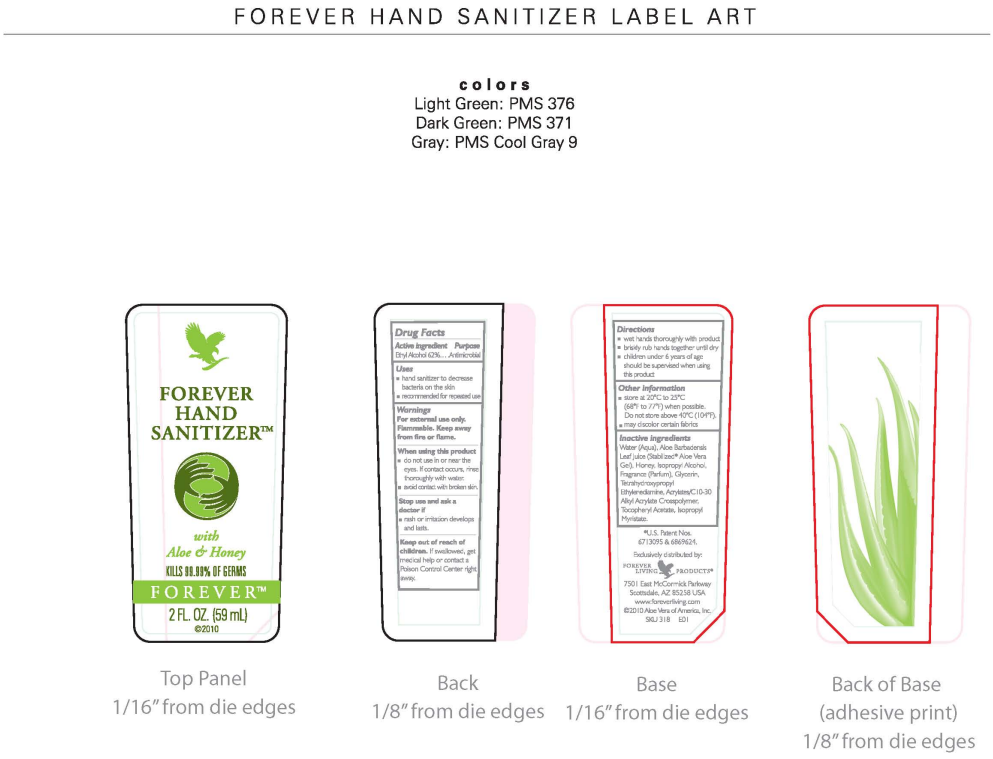
Directions

- wet hands thoroughly with product
- briskly rub hands together until dry
- children under 6 years of age should be supervised when using this product

Other information

- store at 20 °C to 25 °C (68°F to 77°F) when possible. Do not store above 40 °C (104° F).
- may discolor certain fabrics

Package Label Principal Display Panel



Label

FOREVER HAND SANITIZER HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11697-318
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	36.58 mL in 59 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
HONEY (UNII: Y9H1V576FH)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
LAVANDULA ANGUSTIFOLIA FLOWERING TOP (UNII: 9YT4B71U8P)	
CYMBOPOGON CITRATUS LEAF (UNII: 06JMS448M5)	
GLYCERIN (UNII: PDC6A3C0OX)	
EDETOL (UNII: Q4R969U9FR)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11697-318-05	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/16/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	08/16/2010	

Labeler - Aloe Vera of America, Inc. (049049463)

Establishment

Name	Address	ID/FEI	Business Operations

Dhaliwal Pharmaceuticals Laboratories,LLC		116933772	manufacture(11697-318)