GNC INSTANT HAND SANITIZER- alcohol gel
GNC Holdings LLC

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

Active ingredient

Ethyl Alcohol 66.5%

Purpose

Antiseptic

Uses

- hand sanitizer to help reduce bacteria on hands and skin
- recommended for repeated use

Warnings

For external use only

Flammable, keep away from fire or flames

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

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

Purified Water, Carbomer, Aloe Vera, Triethanolamine, Fragrance, D&C Green #5, D&C Yellow #10

Principal Display Panel - 118 mL Bottle Label

GNC
LIVE WELL

hand
sanitizer

Ethyl Alcohol 66.5%

FRESH SCENT

NDC 43655-1111-0

4.0 FL. OZ. (118 mL)

Principal Display Panel - 237 mL Bottle Label

GNC
LIVE WELL

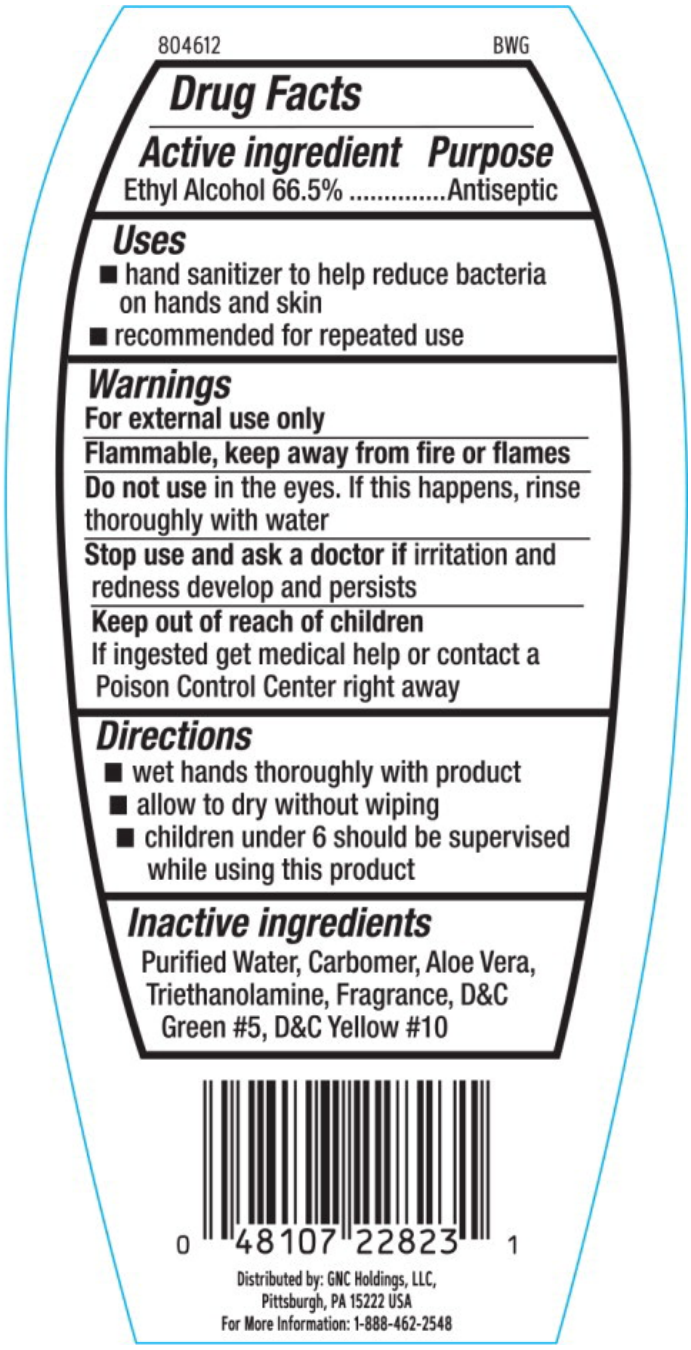
hand
sanitizer

Ethyl Alcohol 66.5%

FRESH SCENT

NDC 43655-1111-1

8.0 FL. OZ. (237 mL)



GNC INSTANT HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:43655-1111
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	66.5 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
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:43655-1111-0	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2021	
2	NDC:43655-1111-1	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2021	

Labeler - GNC Holdings LLC (117772252)

Registrant - Safetec of America, Inc. (874965262)

Establishment

Name	Address	ID/FEI	Business Operations
Safetec of America, Inc.		874965262	MANUFACTURE(43655-1111)