

GNC ANTIMICROBIAL HAND WIPES- alcohol cloth
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

Use

- for sanitizing to decrease bacteria on skin when soap and water is not available

Warnings

For external use only.

Flammable, keep away from fire or flame

Do not use

- in the eyes, if contact occurs, rinse eyes thoroughly with water

Stop use and ask a doctor if

- irritation or redness develop and persists

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

Directions

- tear open packet and remove towelette
- wipe hand, finger and wrist areas thoroughly with towelette for 15 seconds
- dispose of used towelette properly
- children under 6 years of age should be supervised when using this product

Inactive Ingredients

Purified Water, Aloe Vera, Triethanolamine, Fragrance

Principal Display Panel - Carton Label

NDC 43655-3111-0

GNC
LIVE WELL

Antimicrobial
hand wipes

Ethyl Alcohol 66.5%

FRESH SCENT

25 PRE-MOISTENED TOWELETTES



Principal Display Panel - Pouch Label

NDC 43655-3111-0

GNC
LIVE WELL

Antimicrobial
hand wipes

Ethyl Alcohol 66.5%

FRESH SCENT

1 PRE-MOISTENED TOWELETTE



NDC 43655-3111-0
804610 BWG

**antimicrobial
hand wipes**

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FRESH SCENT

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Warnings

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Stop use and ask a doctor if irritation or redness develop and persist

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

Directions ■ tear open packet, remove towelette
■ wipe hand/wrist areas for 15 seconds and discard
■ supervise children under 6 years of age

See carton for full Drug Facts.

This unit is not intended for individual sale.

GNC ANTIMICROBIAL HAND WIPES

alcohol cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43655-3111
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)	
WATER (UNII: 059QF0KO0R)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43655-3111-0	25 in 1 BOX	05/01/2021	
1		1.7 mL in 1 POUCH; Type 0: Not a Combination Product		

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(43655-3111)

Revised: 1/2023

GNC Holdings LLC