

ANTISEPTIC- alcohol cloth
Safetec 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

Active ingredients

Ethyl Alcohol 66.5%

Purpose

Antiseptic

Uses

- Antiseptic cleansing of face, hands and body to decrease bacteria on skin without soap and water

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 or redness develop and persist for more than 72 hours

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
- unfold and use as a washcloth
- supervise children under 6 years of age

Inactive ingredients

aloe vera, fragrance, purified water, triethanolamine

Principal Display Panel - Safetec Antiseptic Towelette Carton Label

NDC 61010-2017-1

Safetec

**Antiseptic
Towelette**

For Professional and Hospital Use

Contents: 100 single-use, premoistened towelettes

Reorder No. 38400



Principal Display Panel - Safetec Antiseptic Towelette Packet Label

Safetec

**Antiseptic
Towelette**

For Professional and Hospital Use

Contents: 1 single-use, premoistened towelette

Manufactured by **SAFETEC OF AMERICA, Inc.**

Buffalo, NY 14215 800-456-7077 www.safetec.com



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ANTISEPTIC

alcohol cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:6 10 10-20 17
Route of Administration	TOPICAL		

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
aloe vera leaf (UNII: ZY8 1Z83H0 X)	
water (UNII: 059QF0KO0R)	
trolamine (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:6 10 10-20 17-0	.0019 L in 1 PACKET; Type 0: Not a Combination Product	05/01/2017	
2	NDC:6 10 10-20 17-1	100 in 1 BOX	05/01/2017	
2		.0019 L in 1 PACKET; 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	part333E	05/01/2017	

Labeler - Safetec of America, Inc. (874965262)

Establishment

Name	Address	ID/FEI	Business Operations
Safetec of America, Inc		874965262	MANUFACTURE(61010-2017)

Revised: 5/2017

Safetec of America, Inc.