

719 WALNUT AVE HAND FRESH ALOE- benzalkonium chloride liquid

Personal Care Products

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.

PC-WM Foaming Hand Sanitizer 0555

Drug Facts

active ingredients

Benzalkonium Chloride

Purpose

Antiseptic

Use

hand sanitizer to help reduce bacteria on skin

Warnings

For external use only.

Flammable Keep away from heat and flame. Do not store 104°F/40°C

Stop use and ask a doctor if

irritation or redness develops.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away. (1-800-222-1222).

Directions

Wet hands thoroughly with the product and rub skin until dry. Children under 6 years of age should be supervised by an adult when using.

Inactive Ingredients

Cetrimonium Chloride, Citric Acid, Dimethicone, Disodium Cocoamphodiacetate, Aloe Barbadensis Leaf Juice, Fragrance, Glycerin, Tetrasodium EDTA Water (Aqua).

719 WALNUT AVE® FOAMING HAND SANITIZER fresh aloe product label

719 WALNUT AVE®

FOAMING HAND SANITIZER

fresh aloe

2 fl OZ (60 mL)

NDC 29500-0555-1

Distributed by:
Wal-Mart Stores, Inc.
Bentonville, AR 72716

Made in China



719 WALNUT AVE HAND FRESH ALOE

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:29500-0555
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	78 mg in 60 mL

Inactive Ingredients

Ingredient Name	Strength
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:29500-0555-1	60 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/19/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/19/2017	

Labeler - Personal Care Products (966155082)**Registrant** - Personal Care Products (966155082)

Revised: 5/2017

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