HAND SANITIZER- ethyl alcohol gel
Walgreen Co

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.

Coastal Breeze Hand Sanitizer
604.000/604AA

Active ingredient

Ethyl alcohol 63%

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

For external use only: hands

Flammable

Keep away from heat and flame

When using this product

- keep out of eyes. Incase of contact with eyes, flush thoroughly with water
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor

- if irritation and redness develop
- condition persists 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

- wet hands thoroughly with product and allow to dry without wiping
- for children under 6, use only under adult supervision
- not recommended for infants

Other information

- do not store above 105ºF
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

water, glycerin, tocopheryl acetate, retinyl palmitate, acrylates/C10-30 alkyl acrylate crosspolymer, benzophenone-4, mannitol, cellulose, hydroxypropyl methylcellulose, fragrance, ext. violet 2, blue 1, red 40, ultramarines

Claims

*Effective at eliminating 99.99% of many common harmful germs and bacteria in as little as 15 seconds

Questions or Comments? 1-800-925-4733

Adverse reactions section

DISTRIBUTED BY: WALGREEN CO.

200 WILMOT RD., DEERFIELD, IL 60015

100% SATISFACTION GUARANTEED

walgreens.com

MADE IN U.S.A. WITH US AND FOREIGN COMPONENTS

Principal display panel

Well at Walgreens

NEW

Hand

Sanitizer

Coastal ocean scent

Kills 99.99% of germs*

2 FL OZ (59 mL)

HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-0596
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	63 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
glycerin (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
SULISOBENZONE (UNII: 1W6L629B4K)	
MANNITOL (UNII: 3OWL53L36A)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
EXT. D&C VIOLET NO. 2 (UNII: G5UX3K0728)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

FD&C RED NO. 40 (UNII: WZB9127XOA)	
ULTRAMARINE BLUE (UNII: I39WR998BI)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0596-16	59 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	01/26/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/26/2015	

Labeler - Walgreen Co (008965063)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(0363-0596)

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Revised: 2/2023 Walgreen Co