HAND SANITIZER- alcohol lotion
Walgreens

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 603

Active ingredient

Ethyl alcohol 62%

Purpose

Antiseptic

Use

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

Warnings

for external use only: hands

Flammable

Keep away from fire or flame

When using this product

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

Stop use and ask a docotor 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 105º F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

benzophenone-4, blue-1, carbomer,fragrance, glycerin, isopropyl alcohol, isopropyl myristate, tocopheryl acetate, water, yellow 5

claims

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

Adverse Reactions

DISTRIBUTED BY: WALGREEN CO.

200 WILMOT RD., DEERFIELD, IL 60015

100% SATISFACTION GUARANTEED

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

principal display panel

Well at

Walgreen

Hand

Sanitizer

Green Apple

Kills 99.99% of grms

2 FLL OZ (59 mL)

HAND SANITIZER

alcohol lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-0968
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	558 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SULISOBENZONE (UNII: 1W6L629B4K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0968-16	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/11/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/11/2019	

Labeler - Walgreens (008965063)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(0363-0968)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(0363-0968)

Revised: 12/2022 Walgreens