ADVANCED HAND SANITIZER- ethyl alcohol gel
Retail Business Services

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.

Advanced Hand Sanitizer with Aloe
439.001/439AB-AC rev1

Active ingredient

Ethyl alcohol 70%

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 fire and 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 doctor if

- irritation or 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, aloe barbadensis leaf juice, glyceryl caprylate/caprate, glycerin, isopropyl myristate, tocopheryl acetate, carbomer or acrylates/C10-30 alkyl acrylate crosspolymer, fragrance, benzophenone-4, blue 1, yellow 5

DISTRIBUTED BY FOODHOLD U.S.A., LLC,

LANDOVER, MO 20785

1-877-846-9949, 2021 Ahold Delhaize Licensing Sarl

Quality guaranteed or your money back.

principal display panel

CAREONE

hand sanitizer

WITH ALOE

HELPS REDUCE BACTERIA ON THE SKIN

NET 8 fl oz (236 mL)

ADVANCED HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72476-439
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERYL CAPRYLATE/CAPRATE (UNII: G7515SW10N)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
SULISOBENZONE (UNII: 1W6L629B4K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:72476-439-34	236 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	01/04/2021	
2	NDC:72476-439-16	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/04/2021	
3	NDC:72476-439-45	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/04/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/04/2021	

Labeler - Retail Business Services (967989935)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(72476-439)

Establishment

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Vi-Jon, LLC		790752542	manufacture(72476-439)

Revised: 5/2023

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