HAND SANITIZER- alcohol gel
CDMA

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.

Quality Choice Advanced Hand Sanitizer with Aloe
439.001/439AC rev 1

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 heat 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

- skin irritation develops
- 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 witout 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

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

Questions?

1-888-593-0593

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

Distributed by CDMA, Inc.

43157 W Nine Mile

Novi, MI 48376

www.qualitychoice.com

Questions 800-935-2362

principal display panel

QC QUALITY CHOICE

Compare to Germ-X Advanced Hand Sanitizer with Aloe

Advanced

Hand Sanitizer

With Aloe

Kills More Than 99.99% of Germs*

Leaves hands feeling soft

With Moisturizer & Vitamin E

8 FL OZ (236 mL)

HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-439
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.70 mL in 1 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 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:63868-439-34	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/05/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/05/2021	

Labeler - CDMA (011920774)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(63868-439)

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Revised: 3/2022 CDMA