ALCOHOL- alcohol liquid
CVS

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.

Ocean Breeze Hand Sanitizer
471.000/471AA

Active ingredient

Ethyl Alcohol 62%

Purpose

Antiseptic

Use

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

Warnings

For exterenal 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 thoroghly 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, carbomer, fragrance, glycerin, isopropyl myristate, blue 1

Questions?

Satisfaction quaranteed - For questions or comments please call 1-888-287-1915

adverse reaction

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

Distributed by: CVS Pharmacy, Inc

One CVS Drive, Woonsocket, RI

CVS.com

Made in the USA with U.S and foreign components

Principal display panel

ocean breeze scent

antibacterial

hand sanitizer

with moisturizers

kills 99.99% of germs

fast & effective

moisturizers leave hands feeling smooth

paraben & phthalate free

CVS Pharmacy

8 FL OZ (236 mL)

ALCOHOL

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69842-471
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
water (UNII: 059QF0KO0R)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-471-16	89 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/03/2020	
2	NDC:69842-471-34	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/03/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/10/2019	

Labeler - CVS (062312574)

Registrant - Vi Jon, Inc. (088520668)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, Inc.		088520668	manufacture(69842-471)

Establishment

Name	Address	ID/FEI	Business Operations
Vi Jon, Inc.		790752542	manufacture(69842-471)

Revised: 2/2023 CVS