HAND SANITIZER- alcohol liquid
Medline Industries, Inc.

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.

Hand Sanitizer
495

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 doctor if

- irritation or redness develops
- condition persists for more than 72 hours

Keep out of reach 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 suervision
- not recommended for infants

Other information

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

inactive ingredients

water, Aloe barbadensis leaf juice, tocopheryl acetate, glycerin, isopropyl myristate, propylene glycol, carbomer, diisopropylamine

REF HH62G02UMC

www.medline.com

Made in USA with US and Foreign components

for Medline industries, Inc.,

Northfield, IL 60093 USA

1-800-MEDLINE RG17VJO

495.001/495AA Rev 1

principal display panel

UMC

UNIVERSITY MEDICAL CENTER

Hand Sanitizer 62% Ethyl ALCOHOL

Kills 99.99% of germs

Moisturizing & Vitamin E

Gentle enough for repeated use

2 FL OZ (59 mL)

HAND SANITIZER

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:53329-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
water (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
glycerin (UNII: PDC6A3C0OX)	
isopropyl myristate (UNII: 0RE8K4LNJS)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
DIISOPROPYLAMINE (UNII: BR9JLI40NO)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:53329-001-13	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/16/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/16/2017	

Labeler - Medline Industries, Inc. (025460908)

Registrant - Vi-Jon, Inc (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, Inc		088520668	manufacture(53329-001)

Revised: 5/2020 Medline Industries, Inc.