HAND SANITIZER- ethyl alcohol gel
Uline

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.

Uline Hand Sanitizer
370/770

Active Ingredient

Ethyl alcohol 70%

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommend 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 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 thoroghly 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 105 F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

water, glyceryl caprylate/caprate, glycerin, isopropyl myristate, tocopheryl acetate, acrylates/C10-30 alkyl acrylate crosspolymer, fragrance, benzophenone-4

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

Distributed by: ULINE
12575 uline Drive, Pleasant Prairie, WI 53158
1-800-295-5510 - uline.com

Principal display panel

ENRICHED WITH MOISTURIZERS

ULINE

Hand Sanitizer

Kills more than 99.99% of germs*

S-21262

2 FL OZ (59 mL)

770.000/770AA/370AB

HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69790-770
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)	
GLYCERYL CAPRYLATE/CAPRATE (UNII: G7515SW10N)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 809Y72KV36)	
SULISOBENZONE (UNII: 1W6L629B4K)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69790-770-16	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/28/2016	
2	NDC:69790-770-96	221 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/28/2016	
3	NDC:69790-770-34	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/28/2016	
4	NDC:69790-770-88	1999 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/28/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/28/2016	

Labeler - Uline (039612668)

Registrant - Vi-Jon, Inc. (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, Inc.		088520668	manufacture(69790-770)

Revised: 1/2020 Uline