HAND SANITIZER- ethyl alcohol gel
Aldi 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.

Drug facts. Source 370.000-370AB

Active ingredients

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

Flammable, keep away from heat 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

- 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 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

other information

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

Adverse reaction

Distributed by ALDI, Inc.

Batavia, IL 60510

this Bottle Is Recyclable

370.000/370AB

principal display panel

this product is not manufactured or distributed by GOJO Industried, Inc.

distributor or Purell Refreshing Gel

Advanced Hand Sanitizer

compare to PURELL hand sanitizer

source

INVIGORATING GEL

ADVANCED

HAND SANITIZER

KILLS MORE THAN 99.99% OF GERMS

Advanced Formula

Less Drying Formula, Leaved Hand Soft

With Moisturizers to Leave Hands Smooth

8 fl oz

(236 mL)

HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:64024-370
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	700 mg in 1 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 INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
SULISOBENZONE (UNII: 1W6L629B4K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64024-370-34	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/02/2014	
2	NDC:64024-370-88	2000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/02/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/02/2014	

Labeler - Aldi Inc. (944259522)

Registrant - Vi-Jon (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		088520668	manufacture(64024-370)

Revised: 4/2020 Aldi Inc.