ADVANCED- ethyl alcohol gel Marc Glassman, 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.

Advanced Hand Sanitizer 370

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

- 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 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 1050 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

Distributed by: Marc Glassman, Inc.

West 130th St, Cleveland, OH 44130

*Not manufactured or distributed by GOJO Industries, Inc. distributor of Purell Refreshing Gel Advanced Hand Sanitizer

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

DISTRIBUTED BY TOPCO ASSOC.LLC ELK GROVE VILLAGE, IL 60007 1-888-423-0139 topcare@topco.com

SES-MO-15036 SDA-WI-2486 DSP-MO-28 SDP-MO-34

principal display panel

Compare to Purell Advanced Hand Sanitizer

Refreshing Gel

Marcs Advanced Hand Sanitizer

WITH MOISTURIZERS

Kills More Than 99.99% Of Germs**

8 FL OZ (236 mL)



ADVANCED

ethyl alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:68998-370

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 (LINII: 059 OF 0KOOR)		

WATER (UNII: 059QF0KO0R)

GLYCERYL CAPRYLATE/CAPRATE (UNII: G7515SW10N)

GLYCERIN (UNII: PDC6A3C0OX)

ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)

.ALPHA.-TO COPHERO L ACETATE (UNII: 9E8 X80 D2L0) CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO) SULISOBENZONE (UNII: 1W6L629 B4K)

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:68998-370- 34	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/27/2012		
2	NDC:68998-370- 16	59 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/27/2012		
3	NDC:68998-370- 45	946 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/27/2012		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/27/2012	

Labeler - Marc Glassman, Inc (094487477)

Registrant - Vi-Jon (790752542)

Establishment			
Name	Address	ID/FEI	Business Operations
Vi-Jon		088520668	manufacture(68998-370)

Revised: 5/2020 Marc Glassman, Inc