ADVANCED- ethyl alcohol gel
TOPCO Assoc. LLC

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

claims

TopCare

Hand Sanitizer Advanced

active ingredient

Ethyl alcohol 70%

Purpose

Antiseptic

Use

Uses

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

Warnings

Warnings

For external use only-hands

Flammable

Flammable. Keep away from heat and flame.

When using this product

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

Stop use and ask a doctor if skin irritation develops

Keep out of reach of children

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

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

Other information

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

Inactive ingredients

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

Disclaimer

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

claims

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

Adverse reaction

DISTRIBUTED BY TOPCO ASSOC.LLC

ELK GROVE VILLAGE, IL 60007

1-888-423-0139

topcare@topco.com

principal display panel

TopCare

Kills More Than

99.99% of Germs

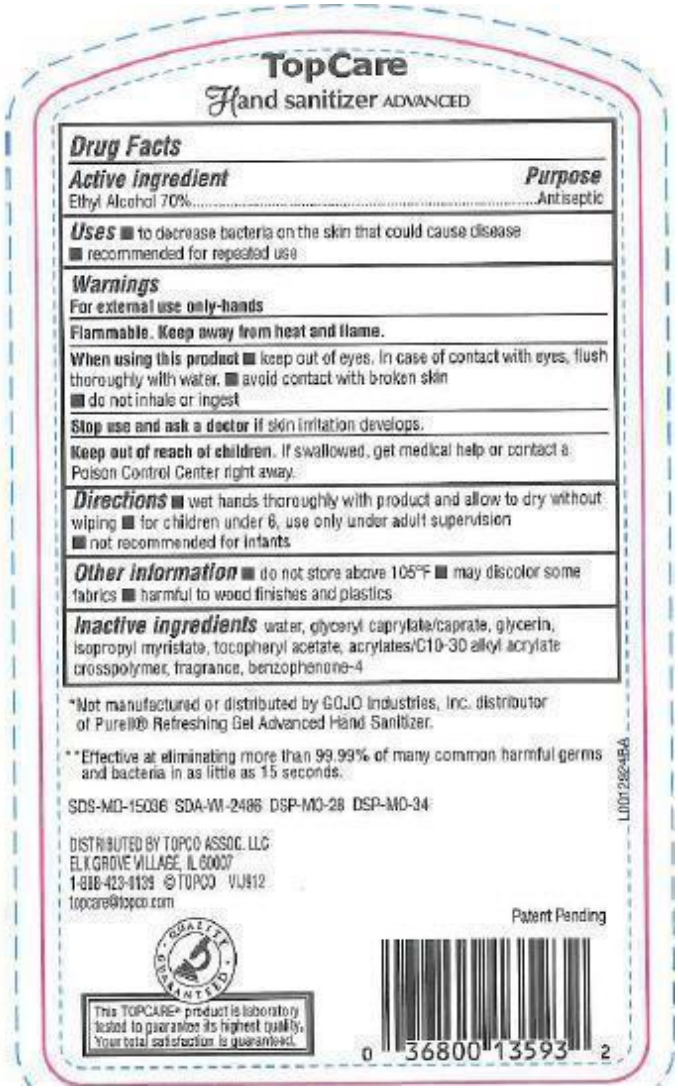
Hand

sanitizer

ADVANCED

Compare to Purell

32 FL OZ (946 mL)



ADVANCED

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:36800-370
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	616 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 COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	
SULISOBENZONE (UNII: 1W6L629B4K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:36800-370-34	236 mL in 1 BOTTLE, PUMP		
2	NDC:36800-370-16	59 mL in 1 BOTTLE, PUMP		
3	NDC:36800-370-45	946 mL in 1 BOTTLE, PUMP		

Marketing Information

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

Labeler - TOPCO Assoc. LLC (006935977)

Registrant - Vi-Jon (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		088520668	manufacture(36800-370)

Revised: 10/2012 TOPCO Assoc. LLC