

ADVANCED HAND SANITIZER- alcohol gel
Walgreen CO

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.

Walgreens Advanced Hand Sanitizer
370.001/370E rev 1

Active Ingredient

Ethyl alcohol 70%

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 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

skin irritation develops.

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

Inactive ingredients

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

Questions?

Call 1-888-593-0593

Claims

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

*This product is 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 harmful germs and bacteria in as little as 15 seconds

Made in the U.S.A. with U.S. and foreign components

DISTRIBUTED BY: WALGREEN CO., 200 WILMOT., DEERFIELD, IL 60015

100% SATISFACTION GUARANTEED walgreens.com

principal display panel

Walgreens

Compare to Purell

ADVANCED Hand Sanitizer + Vitamin E

- Leaves hand soft
- Contains moisturizers
- Kills more than 99.99% of germs*

32 FL OZ (946 mL)



ADVANCED HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-0370
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: 059QF0K00R)	
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:0363-0370-16	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2013	
2	NDC:0363-0370-34	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2013	
3	NDC:0363-0370-42	828 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2013	
4	NDC:0363-0370-45	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/08/2013	

Labeler - Walgreen CO (008965063)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(0363-0370)

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Revised: 3/2022

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