

INSTANT HAND SANITIZER CLASSIC- ethyl alcohol liquid
Universal Distribution Center LLC

INSTANT HAND SANITIZER Classic

Drug Facts

Active ingredient

Ethyl Alcohol 70% v/v

Purpose

Antimicrobial

Uses

hand sanitizer to help reduce bacteria on the skin.

Warnings

For external use only.

Flammable. Keep away from heat and flame.

When using this product avoid contact with face, eyes, and broken skin. In case of eye contact, flush with plenty of water and seek medical advice.

Stop use and ask a doctor if irritation or redness develops.

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

Directions

• wet hands thoroughly with product and rub into skin until dry. • Children under 6 years of age should be supervised by an adult when using.

Inactive ingredients

aqua (water), acrylates/c10-30 alkyl acrylate crosspolymer, aminomethyl propanol, aloe barbadensis leaf juice, denatonium benzoate, fragrance, glycerine, propylene glycol, tocopheryl acetate

With Vitamin E & Other Moisturizers

KILLS GERMS WITHOUT WATER

Mfd for and Distributed by:
Universal Distribution Center LLC
330 Applegarth Road,
Monroe Township, NJ 08831

www.universaldc.com

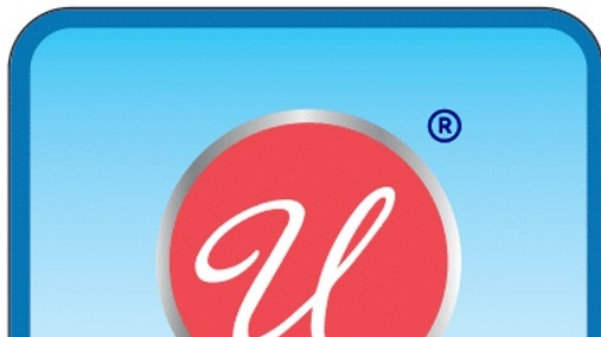
MADE IN CHINA

Packaging

Package: 236 mL



Package: 1 L (1000 mL)





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ITEM# 56032

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MADE IN CHINA

34 FL. OZ (1L)

INSTANT HAND SANITIZER CLASSIC

ethyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52000-434
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)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
ALOE VERA LEAF JUICE (UNII: RUE8E6T4NB)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-434-08	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	11/20/2025	
2	NDC:52000-434-01	1000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	11/20/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	11/20/2025	

Labeler - Universal Distribution Center LLC (019180459)

Revised: 11/2025

Universal Distribution Center LLC