ADVANCED HAND SANITIZER- advanced hand sanitizer liquid
Chain Drug Marketing Association

Quality Choice 370.001/370AE Rev 2
Advanced Hand Sanitizer

Active ingredient

Ethyl alcohol 70% {v/v}

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 fire 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 if

- 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 105º F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

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

Disclaimer

*This product is not manufactured or distributed by GOJO Industries, Inc., distributor of Purell ®Advanced Hand Sanitizer.

Claims

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

Adverse Reaction

100% QC SATISFACTION GUARANTEED

Distributed by CDMA, Inc.

Novi, MI 48375

www.qualitychoice.com

Questions: 800-935-2362

Principal Display Panel

NDC 83324-065-02

QC ®

QUALITY CHOICE

Compare to Purell ®Advanced Hand Sanitizer*

Advanced Hand Sanitizer

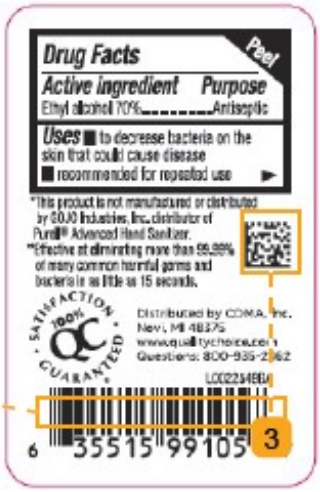
Original

Kills More Than 99.99% of Germs**

Leaves Hands Feeling Soft

With Moisturizers & Vitamin E

2 FL OZ (59 mL)



ADVANCED HAND SANITIZER

advanced hand sanitizer liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-065
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)	
GLYCERYL MONOCAPRYLOCAPRATE (UNII: G7515SW10N)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CARBOMER (UNII: 0A5MM307FC)	

ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
SULISOBENZONE (UNII: 1W6L629B4K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-065-08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/19/2026	
2	NDC:83324-065-02	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/19/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	02/19/2026	

Labeler - Chain Drug Marketing Association (011920774)

Registrant - Nice-Pak Products, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091520	manufacture(83324-065)

Establishment

Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091514	manufacture(83324-065)

Revised: 2/2026 Chain Drug Marketing Association