

HAND SANITIZER- alcohol gel
Walmart Inc.

Equate 370.001-370AB-AE rev 1
Advanced Hand Sanitizer

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

Inactive ingredients

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

Claims

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

ADVERSE REACTION

Satisfaction guaranteed - For questions or comments please call 1-888-287-1915.

DISTRIBUTED BY: Walmart Inc.,

Bentonville, AR 72716

DSP-TN-21091

DSP-MO-20087

Pat. 9,161,982

PLASTIC BOTTLE

how2recycle.info

PRINCIPAL DISPLAY PANEL

NDC 79903-335-12

equate™

Advanced

Hand

Sanitizer

KILLS 99.99% OF GERMS*

with Vitamin E

- Moisturizers leave hands feeling smooth

Made in the USA with domestic and imported ingredients

Factory Certified

12 FL OZ (354mL)

NDC 79903-335-12

equate™

Advanced Hand Sanitizer

KILLS 99.99% OF GERMS*

with Vitamin E

- Moisturizers leave hands feeling smooth

12 FL OZ (354mL)

Pat. 8,161,082

DISTRIBUTED BY: Walmart Inc., Bentonville, AR 72716

DSP-TN-21091
DSP-MO-20087

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Drug Facts

Active ingredient	Purpose
Ethyl alcohol 70%.....	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 105°F ■ may discolor some fabrics ■ harmful to wool finishes and plastics

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

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

Satisfaction guaranteed - For questions or comments please call 1-888-287-1915.

Pat. 8,161,082

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HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79903-335
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 CAPRYLATE/CAPRATE (UNII: G7515SW10N)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
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:79903-335-12	355 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/10/2025	
2	NDC:79903-335-32	946 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/10/2025	
3	NDC:79903-335-60	1770 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/10/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	03/10/2025	

Labeler - Walmart Inc. (051957769)

Registrant - Nice-Pak Products, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091514	manufacture(79903-335)

Revised: 2/2026

Walmart Inc.