

HAND SANITIZER- ethyl alcohol gel
Wal-Mart Stores, Inc.,

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.

Equate 119.000 119AA

Active ingredient

Ethyl Alcohol 62%

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

Inactive ingredients

water, carbomer, fragrance, glycerin, isopropyl myristate, red 40, yellow 5

claims

Effective at eliminating 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

Adverse reaction

DISTRIBUTED BY: Wal-Mart Stores, Inc.,
Bentonville, AR 72716

principal display panel

equate

Nectarine

Smoothie

Hand Sanitizer

KILLS 99.99% OF GERMS

- Moisturizers leave hands smooth

3 FL OZ (89 mL)



HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-720
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	620 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-720-21	88 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2018	

Labeler - Wal-Mart Stores, Inc., (051957769)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(49035-720)

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Revised: 5/2022

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